

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/05/97ISR Number: 10000158Report Type:Expedited (15-DaCompany Report #001-0945-970644
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Apnoea Clonic Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
2700 MG PER DAY(900 MG TID), ORAL		Overdose					
100 MG, PRN				Tylenol (Paracetamol)	SS		
10 MG, QHS, UNKNOWN				Elavil (Amitriptyline Hydrochloride)	SS		

Date:11/10/97ISR Number: 3000018-XReport Type:Expedited (15-DaCompany Report #001-0945-970659
 Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest Respiratory Distress	Health Professional	Neurontin 400 Mg (Gabapentin)	PS		ORAL
400MG BID,PER ORAL				Wellbutrin (Amfebutamone Hydrochloride)	C		

Date:11/10/97ISR Number: 3000198-6Report Type:Expedited (15-DaCompany Report #001-0945-970563
 Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG, Initial or Prolonged BID,PER ORAL	2 WK	Blood Creatine Phosphokinase Increased Cardiomegaly Chest Pain Haemoglobin Decreased Lung Infiltration Pericarditis Pleural Effusion Red Blood Cell Count Decreased White Blood Cell Count Decreased	Health Professional	Neurontin Estradiol Levothyroxine Mvi Vitamins Calcium	PS C C C C C		ORAL

Date:11/12/97ISR Number: 3000388-2Report Type:Expedited (15-DaCompany Report #001-0945-970660
Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG, TID, Other PER ORAL		Bradycardia Cardiac Arrest Convulsion Extrasystoles Syncope	Health Professional	Neurontin 300 Mg (Gabapentin)	PS		ORAL

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Date:11/12/97ISR Number: 3000531-5Report Type:Expedited (15-DaCompany Report #001-0945-970469

Age:39 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1200 MG/900	Complex Partial Seizures	Health	Neurontin	PS		ORAL
Initial or Prolonged MG/1200MG	Feeling Drunk	Professional				
DAILY, ORAL	Liver Function Test					
	Abnormal		Daypro	SS		
	Vomiting		Dilantin Kapseals	C		
			Dilantin Kapseals	C		
			Folic Acid	C		
			Premarin	C		
			Sudafed	C		
			Synthroid	C		
			Vitamins	C		

Date:11/12/97ISR Number: 3097789-3Report Type:Expedited (15-DaCompany Report #001-0945-970414

Age: Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Congenital Anomaly 400 MG,TID, /	Blood Creatinine	Health	Neurontin	PS		
EXPOSURE	Increased	Professional				
INUTERO	Complications Of Maternal					
400 MG,TID,	Exposure To Therapeutic		Tegretol	SS		ORAL
EXPOSURE IN	Drugs					
UTERO	Hydronephrosis					
STOPPED 1	Multiple Congenital		Depo-Provera	SS		
MONTH BEFORE	Abnormalities					
CONCEPTION	Vesicoureteric Reflux					
			Prenatal Vitamins	SS		

Date:11/13/97ISR Number: 3005992-3Report Type:Direct
Age:75 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 400MG PO 6 MON	Asthenia		Neurontin	PS		ORAL
Initial or Prolonged	Brain Neoplasm Cerebellar Infarction Condition Aggravated Depressed Level Of Consciousness Dizziness Hypotension Mental Impairment Myocardial Infarction					

Date:11/20/97ISR Number: 3002019-4Report Type:Expedited (15-DaCompany Report #001-0945-970373
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Breath Sounds Decreased Haemoptysis Hypoxia Left Ventricular Failure Lung Infiltration Lymphocyte Count Decreased Muscle Spasms Neutrophil Count

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Dose	Duration	Increased Pneumonia Pulmonary Haemorrhage Pyrexia	Report Source	Product	Role	Manufacturer	Route
		Rales Sinus Tachycardia White Blood Cell Count Increased	Health Professional	Neurontin (Gabapentin) Trazodon Ultram (Tramadol Hydrochloride)	PS C C		

Date:11/20/97ISR Number: 3002190-4Report Type:Expedited (15-DaCompany Report #001-0945-970481
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNK		Retinitis Pigmentosa	Health Professional	Neurotin	PS		

Date:11/20/97ISR Number: 3005890-5Report Type:Direct Company Report #
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 TID		Nightmare		Gabapentin	PS		

Date:11/21/97ISR Number: 3002209-0Report Type:Expedited (15-DaCompany Report #001-0945-970680
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300 MG TID, Initial or Prolonged PER ORAL		Arthralgia Gastroenteritis Helicobacter Joint Stiffness Sedation	Health Professional	Neurontin Risperidone (Risperidone) Prevacid (Lansoprazole) Propulsid	PS SS C		ORAL

(Cisapride) C
Xanax (Alprazolam) C

Date:12/01/97ISR Number: 3003154-7Report Type:Expedited (15-DaCompany Report #001-0945-970696
Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 600MG TID, PER ORAL		Balance Disorder Fall Fatigue	Consumer	Neurontin Vasotec Levothyroxine Multiple Vitamins Potassium Indomethacin	PS C C C C C		ORAL

Date:12/01/97ISR Number: 3003565-XReport Type:Expedited (15-DaCompany Report #001-0945-970562
Age:31 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Coordination Abnormal Dysarthria Palpitations Sedation

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		Speech Disorder Tongue Oedema				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Health Professional	Neurontin 300mg /Gabapentin	PS	
300MG, TID, PER ORAL				Klonopin	C	
				Prozac	C	
				Ritalin	C	
				Tylenol	C	

Date:12/01/97ISR Number: 3006217-5Report Type:Direct
Age:55 YR Gender:Female I/FU:I

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Gabapentin	PS		ORAL
300 MG PO QID 2 MON				Atenolol	C		
				Chlorthalidone	C		

Date:12/03/97ISR Number: 3007268-7Report Type:Direct
Age:39 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 100MG P.O.		Pruritus		Gabapentin	PS		ORAL
Initial or Prolonged Q8HOUR		Urticaria					
Other							

Date:12/05/97ISR Number: 3004252-4Report Type:Expedited (15-DaCompany Report #97USA12050
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Congenital Anomaly
 TRANSPLACENTAL
 Complications Of Maternal
 Exposure To Therapeutic
 TRANSPLACENTAL
 Health
 Professional
 Tegretol
 Neurontin
 PS
 SS
 Drugs
 Congenital Genitourinary
 Abnormality
 Hydronephrosis
 Urethral Valves

Date:12/09/97ISR Number: 3004776-XReport Type:Expedited (15-DaCompany Report #001-0945-970644
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2700 MG (900, Initial or Prolonged MG TID), PER		Apnoea Clonic Convulsion	Health Professional	Neurontin	PS		ORAL
ORAL		Encephalopathy	Company				
100 MG PRN		Overdose	Representative	Tylenol	SS		
10 MG QHS		Speech Disorder		Elavil	SS		

Date:12/10/97ISR Number: 3006332-6Report Type:Expedited (15-DaCompany Report #044-0945-970079
 Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Chest Pain Coordination Abnormal Epilepsy

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pleuritic Pain Vision Blurred	Foreign Health Professional	Neurontin Tegretol Warfarin Digoxin Frumil Sotalol Clindamycin Cephradine	PS C C C C C C C		

Date:12/11/97ISR Number: 3007729-0Report Type:Direct
Age:25 YR Gender:Female I/FU:I

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Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	250 QID		Attention		Depakote	PS		ORAL
Other	300 MG QID		Deficit/Hyperactivity Disorder Thrombocytopenia		Neurontin	SS		

Date:12/29/97ISR Number: 3015056-0Report Type:Expedited (15-DaCompany Report #001-0945-970680
Age:43 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 900 MG (300 Initial or Prolonged MG, TID) PER			Arthralgia	Health Professional	Neurontin	PS		ORAL
ORAL	0.5 MG (,HS),	1 MON	Hormone Increased Joint Stiffness Reiter'S Syndrome		Risperidone	SS		ORAL
PER ORAL			Sedation Thyroxine Decreased		Xanax Prevacid Propulsid	C C C		

Date:12/31/97ISR Number: 3012904-5Report Type:Direct
Age:40 YR Gender:Male I/FU:I

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Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 300 MG III PO	Dermatitis Exfoliative		Gabapentin	PS		ORAL
Initial or Prolonged TID	Face Oedema Oedema Peripheral					

Date:12/31/97ISR Number: 3012906-9Report Type:Direct
Age:48 YR Gender:Female I/FU:I

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Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 300 MG PO QD	Confusional State		Gabapentin	PS	Parke Davis	ORAL
Initial or Prolonged	Coordination Abnormal Fall Hallucination		Trental Diltiazem Atenolol Elavil	C C C C		

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Date:12/31/97ISR Number: 3013663-2Report Type:Expedited (15-DaCompany Report #001-0945-973040

Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 1200 MG (300 Initial or Prolonged MG, QID), PER ORAL	Convulsion Infection Vision Blurred	Consumer	Neurontin	PS		ORAL
			Lasix	C		
			Klonopin	C		
			Coumadin	C		
			Aldactone	C		
			Lactulose	C		
			Neomycin	C		

Date:12/31/97ISR Number: 3078536-8Report Type:Periodic Company Report #9720452

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2.00 GRAM TOTAL: DAILY: ORAL	Convulsion Gastric Disorder Nausea	Health Professional	Zithromax	PS		ORAL
1200.00 MG TOTAL: DAILY: ORAL	Vomiting		Tegretol	SS		ORAL
ORAL			Neurontin	SS		ORAL
ORAL			Paxil	C		
			Birth Control	C		

Date:01/02/98ISR Number: 3014010-2Report Type:Expedited (15-DaCompany Report #001-0945-973023

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 600-1500 MG Initial or Prolonged PER DAY PER			Medication Error	Literature	Neutotin	PS	ORAL
ORAL	3	MON	Professional				

Date:01/05/98ISR Number: 3013768-6Report Type:Expedited (15-DaCompany Report #001-0945-973040
 Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1200 MG (300 Initial or Prolonged MG, QID), PER			Convulsion	Consumer	Neurontin	PS	ORAL
ORAL			Infection				
			Vision Blurred				
			Visual Disturbance	Lasix	C		
				Klonopin	C		
				Coumadin	C		
				Aldactone	C		
				Lactulose	C		
				Neomycin	C		

Date:01/07/98ISR Number: 3017460-3Report Type:Direct Company Report #
 Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG 1T QID			Balance Disorder	Neurontin	PS		
			Headache				

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Date:01/13/98ISR Number: 3016621-7Report Type:Expedited (15-DaCompany Report #001-0945-970680

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 900 MG (300 Initial or Prolonged MG, TID), PER	Arthralgia	Health	Neurontin	PS		ORAL
ORAL	Blood Thyroid Stimulating Hormone Increased	Professional				
0.5 MG (,HS), PER ORAL	Joint Stiffness Sedation		Risperidone	SS		ORAL
	Thyroxine Decreased		Depakote Xanax Prevacid Propulsid Mellaril	C C C C C		

Date:01/15/98ISR Number: 3016250-5Report Type:Expedited (15-DaCompany Report #001-0945-970579

Age:32 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2700 MG (900 MG, TID) PER	Coma Condition Aggravated	Health Professional	Neurontin 300mg (Gabapentin)	PS		ORAL
ORAL	Haemorrhagic Stroke Hyponatraemia		Tegretol (Carbamazepine) (Risperidone) (Lorazepam) (Levothyroxine) (Fluoxetine)	C C C C C		

Date:01/29/98ISR Number: 3020666-0Report Type:Expedited (15-DaCompany Report #001-0945-980045

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Deep Vein Thrombosis	Health	Neurontin	PS		
Initial or Prolonged			Professional	Estrogens	C		

Date:01/29/98ISR Number: 3086385-XReport Type:Periodic Company Report #8-97303-006K
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Redux	PS		ORAL
15 MG TWICE		Oedema					
DAILY, ORAL		Palpitations		Nadolol	SS		
20 MG ONCE		Pruritus					
DAILY				Neurontin	SS		
300 MG THREE							
TIMES DAILY				Zoloft	SS		
100 MG FOUR							
TIMES DAILY				Aspirin	C		
				Chroimium P	C		
				Darvocet	C		
				Daypro	C		
				K Dur	C		
				Lasix	C		
				Multivitamin	C		

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Date:01/30/98ISR Number: 3021463-2Report Type:Expedited (15-DaCompany Report #001-0945-980037

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Disorder	Health	Neurontin	PS		
UNK			Professional	Doxepin	C		
				Zoloft	C		

Date:02/02/98ISR Number: 3116164-6Report Type:Periodic Company Report #A0057361

Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dermatitis	Health	Lamictal	PS		ORAL
100 MG /TWICE			Professional				
PER DAY/ ORAL			Company	Gabapentin	SS		ORAL
100 MG /TWICE			Representative				
PER DAY/ORAL							

Date:02/03/98ISR Number: 3022196-9Report Type:Expedited (15-DaCompany Report #044-0073-980004

Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abortion Spontaneous	Foreign	Phenytoin Sodium	PS		ORAL
400 MG DAILY			Health				
Initial or Prolonged			Professional	Neurontin	SS		
PER ORAL							
400 MG DAILY							

Date:02/04/98ISR Number: 3024241-3Report Type:Direct Company Report #

Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Headache Gabapentin PS
900 MG/DAY

Date:02/04/98ISR Number: 3024243-7Report Type:Direct Company Report #
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nausea		Gabapentin	PS		
300 MG TID				Apap	C		
				Ecasa	C		
				Colestipol	C		
				Felodipine	C		
				Hctz	C		
				Lisinopril	C		
				Metroprolol	C		
				Simvastatin	C		

Date:02/04/98ISR Number: 3024245-0Report Type:Direct Company Report #
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Mouth		Gabapentin	PS		ORAL
300 MG PO TID				Baclofen	C		
				Propoxyphene	C		
				Lorazepam	C		
				Maalox	C		
				Docusate Sodium	C		
				...	C		
				Ofloxacin	C		

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 C

Date:02/04/98ISR Number: 3024246-2Report Type:Direct
 Age:74 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue		Gabapentin	PS		
300 MG /DAY		Fluid Retention		Allopurinol	C		
		Oedema		Bethanechol	C		
				Biscodyl	C		
				Bumetanide	C		
				Captopril	C		
				Fentanyl	C		
				Finasteride	C		
				Guaifenisin	C		
				Ipratropium	C		
				Isosorbide	C		
				Metolazone	C		

Date:02/04/98ISR Number: 3024250-4Report Type:Direct
 Age:55 YR Gender:Male I/FU:I

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysarthria		Gabapentin	PS		
300 MG				Amitriptyline	C		
				Aspirin	C		
				Atenolol	C		
				Baclofen	C		
				Docusate	C		
				Pepcid	C		
				Motrin	C		
				Prazosin	C		
				Zoloft	C		
				Trazodone	C		

Date:02/04/98ISR Number: 3024251-6Report Type:Direct
Age:52 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Lethargy		Gabapentin	PS		ORAL
300 MG PO TID							

Date:02/04/98ISR Number: 3024252-8Report Type:Direct
Age:69 YR Gender:Male I/FU:I

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Amnesia		Gabapentin	PS		ORAL
300 MG PO BID							

Date:02/04/98ISR Number: 3024257-7Report Type:Direct
Age:71 YR Gender:Male I/FU:I

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oedema		Gabapentin	PS		
900 MG/DAY							
				Apap	C		
				Allopurinol	C		
				Atenolol	C		

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Diltiazem C
 Fluvastatin C
 Gemfibrosil C
 Isosorbide C
 Naftifine C
 Potassium Chloride C
 Prazosin C

Date:02/05/98ISR Number: 3023942-0Report Type:Expedited (15-DaCompany Report #001-0945-980053
 Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 3600 MG	Arthralgia	Consumer	Neurontin	PS		ORAL
Initial or Prolonged (DAILY), PER	Difficulty In Walking					
ORAL	Myalgia					
			Levaquin	C		
			Diflucan	C		
			Herbs	C		

Date:02/05/98ISR Number: 3024370-4Report Type:Direct Company Report #
 Age:76 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG ONE	Fatigue		Gabapentin	PS		
TID						
			Insulin Nph	C		
			Vit E	C		
			Doxepin	C		
			Famotidine	C		

Date:02/05/98ISR Number: 3024454-0Report Type:Direct Company Report #
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Health	Gabapentin	PS		
300 MG Q HS			Professional				

Date:02/05/98ISR Number: 3024466-7Report Type:Direct Company Report #
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bone Pain	Health	Gabapentin	PS		
300 MG BID			Professional	Asa	C		
		Headache		Prazosin	C		
		Neck Pain		Capsaicin	C		
				Simvastatin	C		
				Methocarbamol	C		

Date:02/06/98ISR Number: 3029535-3Report Type:Direct Company Report #
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Psychotic Disorder		Gabapentin	PS		
500MG TO							
Initial or Prolonged							
2100MG/DAY				Allegra	C		
				Lasix	C		

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Synthroid	C
Lithium	C
Xyprexa	C
Premarin	C
Temazepam	C
Stelazine	C

Date:02/17/98ISR Number: 3030420-1Report Type:Expedited (15-DaCompany Report #044-0945-973003
 Age:48 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 500 MG DAILY, Initial or Prolonged PER ORAL	Hepatic Function Abnormal	Foreign Health	Neurontin	PS		ORAL
		Professional	Ferrous Sulfate Sodium Docusate Phosphate Enema Diazepam	C C C C		

Date:02/17/98ISR Number: 3031315-XReport Type:Expedited (15-DaCompany Report #9802119
 Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required 100.00 MG Intervention to TOTAL DAILY Prevent Permanent ORAL Impairment/Damage ORAL ORAL	Alopecia Back Pain Condition Aggravated Intervertebral Disc Protrusion Migraine Pain In Extremity Stress Thyroid Disorder	Consumer	Zoloft Prilosec Neurontin	PS SS SS		ORAL ORAL ORAL

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 800 MG		Antinuclear Antibody	Consumer	Neurontin	PS		ORAL
Initial or Prolonged (DAILY) PER		Positive	Health				
ORAL		Dna Antibody Positive	Professional				
		Supraventricular Tachycardia Systemic Lupus Erythematosis					

Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - INTRA VENOUS	135MG/M2 IV	Anaemia		Taxol	PS		
Initial or Prolonged Q21 DAYS		Asthenia					
300MG/PO		Ecchymosis		Neurontin	SS		
INTRA VENOUS	75 MG/M2 IV	Haematochezia		Cisplatin	SS		
Q21 DAYS		Neuropathy Peripheral					
		Purpura Thrombocytopenia					

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Date:02/25/98ISR Number: 3036490-9Report Type:Direct
Age:73 YR Gender:Male I/FU:I

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300 MG PO BID		Coordination Abnormal		Gabapentin	PS		ORAL
AND 600 MG HS		Dizziness					
		Muscle Twitching					
		Sedation					
		Vision Blurred					

Date:02/25/98ISR Number: 3036674-XReport Type:Expedited (15-DaCompany Report #971217-107013517
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cerebrovascular Accident	Study	Ultram	PS		ORAL
50 MG, PRN,			Health				
Initial or Prolonged			Professional	Neurontin	SS		
ORAL				Prozac	C		

Date:02/27/98ISR Number: 3041985-8Report Type:Direct
Age:50 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Platelet Count Decreased	Health	Neurontin	PS		
900MG QD			Professional				

Date:02/27/98ISR Number: 3044063-7Report Type:Direct
Age:57 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Decreased Activity	Health	Gabapentin	PS		
500 MG TO							

2100 MG/DAY

Psychotic Disorder

Professional

Allegra	C
Lasix	C
Synthroid	C
Lithium	C
Zyprexa	C
Stelazine	C
Premarin	C
Temazepam	C

Date:03/02/98ISR Number: 3129834-0Report Type:Periodic Company Report #9712638
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Choreoathetosis	Health	Zoloft	PS		ORAL
100.00 MG		Coordination Abnormal	Professional				
TOTAL: DAILY:		Drug Interaction					
ORAL		Headache		Neurontin	SS		ORAL
900.00 TOTAL:		Nervous System Disorder					
TID: ORAL		Speech Disorder		Buspar	C		
		Thinking Abnormal					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3141207-3Report Type:Periodic
Age:56 YR Gender:Male I/FU:I

Company Report #001-0945-973038

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300 MG (100 Initial or Prolonged MG, TID), PER ORAL	Skin Disorder	Health Professional	Neurontin	PS		ORAL
			Haldol (Haloperidol)	C		
			Cogentin	C		
			Debrox	C		
			Prilosec	C		
			Phazyme	C		
			Depakote	C		
			Asa	C		

Date:03/02/98ISR Number: 3148217-0Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-970029

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 800 MG (400 MG, BID), PER ORAL	Chest Pain Headache Leukopenia Myalgia Nausea	Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:03/02/98ISR Number: 3148218-2Report Type:Periodic
Age:47 YR Gender:Female I/FU:I

Company Report #001-0945-970047

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability 300 MG (QHS), PER ORAL	Diplopia Medication Error Vith Nerve Paralysis	Health Professional	Neurontin 300 Mg (Gabapentin)	PS		ORAL

Insulin	C
Glucophage	C
Heemalog	C
Senokot	C

Date:03/02/98ISR Number: 3148221-2Report Type:Periodic Company Report #001-0945-970051
 Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 Disability MG, TID) PER ORAL		Cardiac Failure Deafness Delirium Hallucination, Auditory Lethargy Medication Error Renal Failure	Health Professional	Neurontin 300 Mg (Gabapentin)	PS		ORAL
				Pinolol	C		
				Amitriptyline	C		
				Mestinon	C		
				Prozac	C		
				Klonopin	C		
				Phentermine	C		
				Fenfluramine	C		
				Percocet	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3148223-6Report Type:Periodic
Age:41 YR Gender:Female I/FU:I

Company Report #001-0945-970059

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (600 MG, BID) PER ORAL	Blood Creatinine Increased Blood Urea Increased Encephalopathy	Health Professional	Neurontin 300 Mg (Gabapentin)	PS		ORAL
			Acyclovir	C		
			Mycophenolate	C		
			Tacrolimus	C		
			Lactulose	C		
			Insulin	C		
			Ursodid	C		
			Lasix	C		
			Prednisone	C		
			Procardia	C		
			Prilosec	C		

Date:03/02/98ISR Number: 3148226-1Report Type:Periodic
Age:46 YR Gender:Male I/FU:I

Company Report #001-0945-970109

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged PER ORAL	Deep Vein Thrombosis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Elavil	C		
			Motrin	C		
			Ultram	C		
			Axid	C		

Date:03/02/98ISR Number: 3148229-7Report Type:Periodic
Age:10 YR Gender:Male I/FU:I

Company Report #001-0945-970141

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization -	Convulsion	Consumer	Neurontin 300 Mg			

Initial or Prolonged SEE TEXT, PER	Dermatitis		(Gabapentin)	PS	ORAL
ORAL			Topimax	SS	ORAL
PER ORAL			Sabril	C	

Date:03/02/98ISR Number: 3148231-5Report Type:Periodic Company Report #001-0945-970175
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER		Phlebitis	Health Professional	Neurontin 300 Mg (Gabapentin)	PS		ORAL
ORAL				Dilantin Decadron	C C		

Date:03/02/98ISR Number: 3148234-0Report Type:Periodic Company Report #001-0945-970195
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1500 MG (500 MG, TID), PER		Mania	Health Professional	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

(Folic Acid) C
 Ativan C
 (Valproic Acid) C
 Multivitamins C
 (Lithium) C
 Tylenol C

Date:03/02/98ISR Number: 3148237-6Report Type:Periodic Company Report #001-0945-970203
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG, TID), PER		Medication Error - Renal Failure Acute	Health Professional	Neurontin 300 Mg (Gabapentin)	PS		ORAL

ORAL

Coumadin C
 Lasix C
 (Enalapril) C

Date:03/02/98ISR Number: 3148240-6Report Type:Periodic Company Report #001-0945-970214
 Age:82 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Difficulty In Walking Disorientation Medication Error Mental Disorder	Consumer	Neurontin (Gabapentin)	PS		

Date:03/02/98ISR Number: 3148242-XReport Type:Periodic Company Report #001-0945-970300
 Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Cardiac Failure	Consumer	Neurontin 300 Mg			

Initial or Prolonged Congestive (Gabapentin) PS ORAL
 600 MG (300
 MG, TID), PER
 ORAL
 (Quinapril) C
 Synthroid C
 Meclizine C
 (Famoridine) C
 (Cimetidine) C
 (Calcium) C

Date:03/02/98ISR Number: 3148244-3Report Type:Periodic Company Report #001-0945-970302
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Deafness Ear Discomfort	Health Professional	Neurontin 300 Mg (Gabapentin)	PS		ORAL

300 MG (ONE
 DOSE) PER
 ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3148246-7Report Type:Periodic
Age:76 YR Gender:Female I/FU:I

Company Report #001-0945-970367

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 MG, TID), PER ORAL		Thermal Burn Vomiting	Consumer	Neurontin 300 Mg (Gabapentin)	PS		ORAL
1 TAB BID/1 1/2 TABS AT BEDTIME (200 MG), PER ORAL				Tegretol 200mg (Carbamazepine)	SS		ORAL
				Synthroid (Quinapril) (Melozine) (Famotidine)	C C C C		

Date:03/02/98ISR Number: 3148248-0Report Type:Periodic
Age:41 YR Gender:Female I/FU:I

Company Report #001-0945-970383

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 1200 MG (600 MG, BID), PER ORAL		Medication Error Psychotic Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				(Clonazepam) Metamucil Tylenol (Olanzapine) Multivitamins Electroconvulsive	C C C C C C		

Date:03/02/98ISR Number: 3148251-0Report Type:Periodic Company Report #001-0945-970388
Age:7 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Anaemia	Health	Neurontin			
Initial or Prolonged	Lymphocytosis	Professional	(Gabapentin)	PS		ORAL
2 GM, PER						
Other						
ORAL						
			(Zinc)	C		
			Tranxene	C		

Date:03/02/98ISR Number: 3148253-4Report Type:Periodic Company Report #001-0945-970433
Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Abnormal Behaviour	Health	Neurontin 300 Mg			
Initial or Prolonged	Coordination Abnormal	Professional	(Gabapentin)	PS		ORAL
600 MG (300						
MG, TID) PER	Drug Level Above					
ORAL	Therapeutic					
	Mania		Tegretol	C		
	Medication Error		(Lithium)	C		
	Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3148254-6Report Type:Periodic
Age:36 YR Gender:Female I/FU:I

Company Report #001-0945-970449

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Asthenia	Health	Neurontin			
800 MG		Bradyphrenia	Professional	(Gabapentin)	PS		ORAL
(DAILY) PER		Diarrhoea					
ORAL		Disturbance In Attention					
		Dysphagia		Ultram	C		
		Memory Impairment		Trazodone	C		
		Nausea					
		Sedation					
		Tongue Discolouration					
		Tongue Disorder					
		Vomiting					

Date:03/02/98ISR Number: 3148255-8Report Type:Periodic
Age:7 YR Gender:Female I/FU:I

Company Report #001-0945-970454

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Gingivitis	Consumer	Neurontin 300 Mg			
300 MG		Rectal Haemorrhage		(Gabapentin)	PS		ORAL
(DAILY), PER		Weight Increased					
ORAL							

Date:03/02/98ISR Number: 3148258-3Report Type:Periodic
Age:39 YR Gender:Female I/FU:I

Company Report #001-0945-970486

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Deep Vein Thrombosis	Health	Neurontin			
PER ORAL			Professional	(Gabapentin)	PS		ORAL
				Depakote	C		
				Dilantin	C		

Date:03/02/98ISR Number: 3148260-1Report Type:Periodic Company Report #001-0945-970500
Age:8 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3 DOSES, PER		Abnormal Behaviour	Health Professional	Neurontin (Gabapentin)	PS		ORAL

ORAL

Date:03/02/98ISR Number: 3148518-6Report Type:Periodic Company Report #001-0945-970514
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged "MAXIMUM DOSAGE" PER		Medication Error Pneumonia	Health Professional	Neurontin (Gabapentin)	PS		ORAL

ORAL

Date:03/02/98ISR Number: 3148519-8Report Type:Periodic Company Report #001-0945-970538
Age:36 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

Disability

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600MG (DAILY)		Blood Potassium Increased Medication Error	Health Professional	Neurontin (Gabapentin)	PS		ORAL
PER ORAL		Oedema					
		White Blood Cell Count Increased		Klonopin	C		
				Zyprexa	C		
				Percocet	C		
				Trilafon	C		

Date:03/02/98ISR Number: 3148520-4Report Type:Periodic Company Report #001-0945-970549
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Atrioventricular Block First Degree	Health Professional	Neurontin 300mg (Gabapentin)	PS		ORAL
1200MG (600MG		Bradycardia					
, BID) PER		Cerebrovascular Accident					
ORAL		Supraventricular Extrasystoles		Aspirin	C		

Date:03/02/98ISR Number: 3148523-XReport Type:Periodic Company Report #001-0945-970574
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bradycardia	Consumer	Neurontin 300mg (Gabapentin)	PS		ORAL
Hospitalization - Initial or Prolonged 2700MG		Condition Aggravated					
(900MG, TID)		Convulsion					
PER ORAL		Dizziness					
		Mental Impairment		Dilantin Kapseals			
		Visual Disturbance		100mg (Phenytoin			

SEE TEXT, PER

Sodium)

SS

ORAL

ORAL

Aspirin	C
Sertraline	C
Oxybutynin	C
Fellodipine	C
Docusate	C

Date:03/02/98ISR Number: 3148525-3Report Type:Periodic
Age:47 YR Gender:Male I/FU:I

Company Report #001-0945-970608

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900MG (300MG		Apnoea Lethargy	Health Professional	Neurontin 300mg (Gabapentin)	PS		ORAL

TID) PER ORAL

Procardia	C
Paxil	C
Cipro	C
Fosamax	C
Lactulose`	C
Insulin	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3148527-7Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-970613

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900MG (300MG TID) PER ORAL		Deep Vein Thrombosis	Health Professional	Neurontin 300mg (Gabapentin)	PS		ORAL
				Elavil	C		

Date:03/02/98ISR Number: 3148528-9Report Type:Periodic
 Age:35 YR Gender:Female I/FU:I

Company Report #001-09145-970636

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 600MG (DAILY) PER ORAL		Hypoaesthesia	Consumer	Neurontin 300mg (Gabapentin)	PS		ORAL

Date:03/02/98ISR Number: 3148529-0Report Type:Periodic
 Age:59 YR Gender:Male I/FU:I

Company Report #001-0945-970673

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900MG (300MG TID) PER ORAL		Abdominal Pain Upper Dizziness Headache	Consumer	Neurontin 300mg (Gabapentin)	PS		ORAL
				Methadone	C		

Date:03/02/98ISR Number: 3148530-7Report Type:Periodic
 Age:79 YR Gender:Male I/FU:I

Company Report #001-0945-970692

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Pyrexia	Health	Neurontin 300mg			

Initial or Prolonged 1800MG (600MG	Spinal Fracture	Professional	(Gabapentin)	PS	ORAL
TID) PER ORAL					
				Cefizox	C
				Zithromax	C
				Coumadin	C
				Digoxin	C
				Ativan	C
				Norvasc	C
				Inderal La	C
				Zantac	C

Date:03/02/98ISR Number: 3148531-9Report Type:Periodic Company Report #001-0945-973011
Age:73 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900MG (300MG	Aphonia Decreased Appetite Dysphagia	Consumer	Neurontin Capsules 300mg (Gabapentin)	PS		ORAL
TID) PER ORAL						
	Medication Error Pneumonia Aspiration Sedation Weight Decreased		Morphine Percocet Dilaudid	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/98ISR Number: 3148532-0Report Type:Periodic
Age:79 YR Gender:Female I/FU:I

Company Report #001-0945-973033

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900MG (300MG TID) PER ORAL	Activated Partial Thromboplastin Time Prolonged	Health Professional	Neurontin (Gabapentin)	PS		ORAL
	International Normalised Ratio Increased Phlebitis Prothrombin Time Prolonged		Coumadin Synthroid Bumex Allopurinol Maxzide	C C C C C		

Date:03/02/98ISR Number: 3148535-6Report Type:Periodic
Age:63 YR Gender:Female I/FU:I

Company Report #001-0945-973034

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration UNKNOWN 1200MG (400MG TID) UNK	Medication Error Oedema Peripheral		Neurontin (Gabapentin)	PS		
			Digoxin Verapamil Calcitonin Soma Meprobamate	C C C C C		

Date:03/02/98ISR Number: 3148536-8Report Type:Periodic
Age:35 YR Gender:Male I/FU:I

Company Report #001-0945-973042

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1600MG (DAILY) PER	Haematemesis Medication Error	Consumer	Neurontin (Gabapentin)	PS		ORAL

ORAL

Dilantin	C
Percocet	C
Propulsid	C
Elavil	C

Date:03/04/98ISR Number: 3040401-XReport Type:Expedited (15-DaCompany Report #001-0945-980134
Age:43 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 300 MG DAILY Intervention to PER ORAL Prevent Permanent Impairment/Damage	Retinal Tear	Health Professional	Neurontin	PS		ORAL

Date:03/04/98ISR Number: 3040402-1Report Type:Expedited (15-DaCompany Report #001-0945-980139
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Blood Creatine Phosphokinase Increased Rhabdomyolysis	Health Professional	Neurontin Insulin Digoxin Ms Contin Zantac Ecotrin	PS C C C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/04/98ISR Number: 3040403-3Report Type:Expedited (15-DaCompany Report #001-0945-980150

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 180 MG DAILY	Blood Potassium Decreased	Health	Neurontin	PS		
Initial or Prolonged	Heart Rate Decreased	Professional	Atenolol Verapamil	C C		

Date:03/04/98ISR Number: 3050449-7Report Type:Direct

Company Report #

Age:28 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Required 1200 MG PO	Cutaneous Lupus	Health	Neurontin	PS		ORAL
Intervention to TID	Erythematosis	Professional				
Prevent Permanent Impairment/Damage	Rash Erythematous					

Date:03/10/98ISR Number: 3049193-1Report Type:Direct

Company Report #

Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
300 MG TID	Diplopia	Health	Gabapentin	PS		ORAL
	Vision Blurred	Professional	Enalapril Fentanyl Nortriptyline Hydromorphone Naproxen	C C C C C		

Date:03/10/98ISR Number: 3088051-3Report Type:Direct

Company Report #

Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

300 MG BID	Nausea	Gabapentin	PS
	Vomiting	Xanax	C

Date:03/11/98ISR Number: 3054874-XReport Type:Expedited (15-DaCompany Report #001-0945-980140
 Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 900 MG (300 Initial or Prolonged MG , TID), PER ORAL;	Coma Convulsion Pneumonia Aspiration	Health Professional	Neurontin	PS		ORAL

SEVERAL YEARS

THERAPY

Mysoline	C
Multiple Other Medications	C

Date:03/11/98ISR Number: 3054887-8Report Type:Expedited (15-DaCompany Report #001-0945-973038
 Age:56 YR Gender:Male I/FU:F

Outcome Hospitalization - Initial or Prolonged	PT Electroencephalogram Abnormal Haemorrhagic Stroke Obsessive-Compulsive 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
300 MG (100 MG, TID), PER ORAL		Skin Disorder Transient Ischaemic Attack	Health Professional	Neurontin	PS		ORAL
				Haldol	C		
				Debrox	C		
				Prilosec	C		
				Phazyme	C		
				Depakote	C		
				Cogentin	C		
				Atarax	C		
				Asa	C		

Date:03/12/98ISR Number: 3053249-7Report Type:Expedited (15-DaCompany Report #001-0945-970227
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG BID, Initial or Prolonged PER ORAL		Fatigue	Health Professional	Neurontin	PS		ORAL
				Entacyl	C		
				Elavil	C		
				Flonase	C		
				Hydrocodone	C		
				Naproxen	C		
				Rocephin	C		
				Prednisone	C		
				Morphine	C		

Date:03/13/98ISR Number: 3055506-7Report Type:Expedited (15-DaCompany Report #001-0945-980174
Age:1 DY Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization - SEE TEXT, Initial or Prolonged OTHER	Benign Congenital Hypotonia Blood Glucose Increased Complications Of Maternal Exposure To Therapeutic Drugs	Health Professional	Neurontin Klonopin Synthroid Zoloft Prenatal Vitamins	PS SS C C C
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Date:03/13/98ISR Number: 3055508-0Report Type:Expedited (15-DaCompany Report #001-0945-980163
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hodgkin'S Disease Mass	Consumer	Neurontin Dilantin Fosamax	PS C C		

Date:03/16/98ISR Number: 3056553-1Report Type:Expedited (15-DaCompany Report #044-0945-980005
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1200 MG (400 Initial or Prolonged MG, TID), PER ORAL		Arthralgia Arthritis Synovitis	Foreign Health Professional	Neurontin	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lamotrigine C

Date:03/17/98ISR Number: 3056019-9Report Type:Expedited (15-DaCompany Report #001-0945-980165
 Age:83 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hypoxia	Health	Neurontin	PS		
400 MG (,100 Hospitalization - MG QAM, 100 Initial or Prolonged MG		Pneumonia	Professional				
		Pneumonitis					
		Pulmonary Fibrosis		Miacalcin	C		
		Rib Fracture		Testosterone	C		
		White Blood Cell Count Increased					

Date:03/18/98ISR Number: 3056375-1Report Type:Direct Company Report #
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other I PO Q HS, I PO BID		Dermatitis		Neurontin	PS	Parke-Davis	ORAL
		Haemorrhage					
		Skin Fissures					

Date:03/19/98ISR Number: 3057780-XReport Type:Direct Company Report #
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG PO TID 4 WK		Oedema Peripheral		Neurontin	PS		ORAL
				Baclofen	C		
				Oxycontin	C		
				Flexeril	C		

Date:03/23/98ISR Number: 3058470-XReport Type:Expedited (15-DaCompany Report #001-0945-980172
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Astrocytoma	Consumer	Neurontin	PS		ORAL
3600 MG		Condition Aggravated					
(DAILY) PER							
ORAL							
				Dilantin With Phenobarbital Tylenol	C C		

Date:03/23/98ISR Number: 3058472-3Report Type:Expedited (15-DaCompany Report #001-0945-970427
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - PER ORAL		Asthenia	Health	Neurontin	PS		ORAL
Initial or Prolonged		Chromaturia Fatigue Feeling Jittery Haematocrit Decreased Haemolysis Haemolytic Anaemia Palpitations Restlessness	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/23/98ISR Number: 3058475-9Report Type:Expedited (15-DaCompany Report #001-0945-980106
 Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Antinuclear Antibody	Health	Neurontin	PS		ORAL
800 MG		Positive	Professional				
(DAILY) PER		Dna Antibody Positive					
ORAL		Leukopenia					
		Rheumatoid Factor					
		Positive					
		Supraventricular					
		Tachycardia					
		Systemic Lupus					
		Erythematosis					

Date:03/24/98ISR Number: 3058769-7Report Type:Expedited (15-DaCompany Report #001-0945-980178
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Benign Intracranial	Health	Neurontin	PS		ORAL
Hospitalization -		Hypertension	Professional				
900 MG		Headache					
Initial or Prolonged		Vision Blurred					
(300MG, TID)							
PER ORAL							

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		Chest Pain	Health	Redux	PS		ORAL
Other		Drug Withdrawal Syndrome	Professional				
15 MG DAILY		Dyspnoea		Desyrel	SS		ORAL
ORAL	1 YR						
50 MG AT							

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

Date:03/31/98ISR Number: 3059424-XReport Type:Expedited (15-DaCompany Report #001-0945-980037
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Chemotherapy Cardiotoxicity Attenuation	Health Professional	Neurontin (Doxepin) Zoloft (Sertraline Hydrochloride)	PS C C		

Date:03/31/98ISR Number: 3067072-0Report Type:Direct Company Report #
Age:28 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

Required Intervention to Prevent Permanent Dose Impairment/Damage PO 300MG TID	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pancreatitis		Gabapentin	PS		ORAL
				Saquinavir	C		
				Fluconazole	C		
				Dapsone	C		
				Megace	C		
				Zovirax	C		
				Ms Contin	C		
				Zantac	C		
				Epivir	C		
				Bactrim	C		
				Trazadone	C		

Date:04/01/98ISR Number: 3058750-8Report Type:Expedited (15-DaCompany Report #001-0945-980203
 Age:60 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Ammonia Increased Autoimmune Hepatitis Confusional State Hepatic Function Abnormal	Health Professional Company Representative	Neurontin	PS		

Date:04/01/98ISR Number: 3064583-9Report Type:Direct Company Report #
 Age:42 YR Gender:Female I/FU:I

Outcome Dose Required 2 PO TID	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alanine Aminotransferase Increased Blood Creatine Phosphokinase Increased Myositis	Health Professional	Neurontin	PS		ORAL

Date:04/02/98ISR Number: 3058131-7Report Type:Expedited (15-DaCompany Report #001-0945-980204
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatinine Increased Renal Tubular Necrosis	Health Professional	Neurontin Ticlid Hydralazine Axid	PS C C C		

Date:04/05/98ISR Number: 3153527-7Report Type:Periodic Company Report #97USA10341
 Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Stevens-Johnson Syndrome	Health	Tegretol	PS		ORAL
DAILY, ORAL			Professional	Neurontin	SS		ORAL
900 MG DAILY,							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/07/98ISR Number: 3071810-0Report Type:Direct
Age:72 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - NS	Acute Prerenal Failure		Gabapentin	PS		
Initial or Prolonged	Mental Impairment					

Date:04/13/98ISR Number: 3072359-1Report Type:Direct
Age:97 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death 200MG Q HS	Choreoathetosis		Neurontin	PS		
Hospitalization - Initial or Prolonged	Dysarthria Dyskinesia Gait Disturbance Mental Impairment Urinary Incontinence					

Date:04/13/98ISR Number: 3073159-9Report Type:Direct
Age:29 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 600MG 9AM; Initial or Prolonged 600MG PO Q AFTERNOON;900 MG PO Q PM,	Amnesia Diplopia Dysarthria Gait Disturbance Sedation		Gabapentin	PS	Parke Davis	ORAL
			Carbamazepine	C		
			Combivir	C		
			Diazepam	C		
			Zoloft	C		
			Oramorph	C		

Date:04/14/98ISR Number: 3064329-4Report Type:Expedited (15-DaCompany Report #001-0945-980174
Age:1 DY Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Complications Of Maternal	Health	Neurontin	PS		
Initial or Prolonged	Exposure To Therapeutic	Professional	Klonopin	SS		
	Drugs		Synthroid	C		
	Hypotonia		Zoloft	C		
			Prenatal Vitamins	C		

Date:04/14/98ISR Number: 3064330-0Report Type:Expedited (15-DaCompany Report #001-0945-980140
Age:53 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Coma	Health	Neurontin	PS		ORAL
300 MG TID						
Initial or Prolonged	Convulsion	Professional				
ORAL						
	Drug Ineffective		Mysoline	C		
	Pneumonia Aspiration					

Date:04/20/98ISR Number: 3072440-7Report Type:Direct Company Report #
Age:60 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Asthenia	Health	Gabapentin	PS		
300 MG Q DAY						
Initial or Prolonged	Neuropathy Peripheral	Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/22/98ISR Number: 3070077-7Report Type:Periodic
Age:79 YR Gender:Female I/FU:F

Company Report #001-0945-973033

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 900 MG (300 Initial or Prolonged MG TID), PER ORAL	Activated Partial Thromboplastin Time Prolonged	Health Professional	Neurontin	PS		ORAL
	International Normalised Ratio Increased Phlebitis Prothrombin Time Prolonged		Coumadin Synthroid Bumex Allopurinol Digxin Maxzide	C C C C C C		

Date:04/24/98ISR Number: 3072536-XReport Type:Direct
Age: Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly QAM QNOON QPM QHS	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Heart Disease Congenital Premature Baby Premature Labour		Neurontin	PS	Parke-Davis	

Date:04/28/98ISR Number: 3072215-9Report Type:Expedited (15-DaCompany Report #001-0945-980172
Age:26 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 3600 MG (DAILY) PER	Astrocytoma Condition Aggravated	Consumer	Neurontin	PS		ORAL

Glioma

ORAL

Tylenol	C
Felbatol	C
Phenobarbital	C
Dilantin	C

Date:05/01/98ISR Number: 3073037-5Report Type:Direct
 Age:54 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 600 MG BID Initial or Prolonged ORAL	Coordination Abnormal		Gabapentin	PS	Parke-Davis	ORAL

Ibuprofen	C
Multivitamins	C
Famotidine	C
Benzotropine	C
Lorazepam	C
Lithium	C
Levothyroxine	C
Thioridazine	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/01/98ISR Number: 3080313-9Report Type:Direct
Age:44 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300 MG TID		Confusional State		Gabapentin	PS		
		Dizziness					
		Irritability					

Date:05/01/98ISR Number: 3080639-9Report Type:Expedited (15-DaCompany Report #
Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Chromatopsia		Gabapentin	PS		ORAL
300 MG QAM PO							
Intervention to		Vision Blurred					
AND 400 MG							
Prevent Permanent							
TID PO							
Impairment/Damage				Prednisone	C		ORAL
4 MG PO QD							
				Amlodipine	C		ORAL
10 MG PO QD							
				Clonidine	C		ORAL
0.1 MG PO BID							
				Bicitra	C		ORAL
2 TABS PO TID							
				Digoxin	C		ORAL
0.125 MG PO							
				Penuk	C		ORAL
250 MG PO BID							

Date:05/04/98ISR Number: 3074314-4Report Type:Expedited (15-DaCompany Report #001-0945-980165
Age:83 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cryptogenic Organizing	Health	Neurontin	PS		ORAL
400 MG (,100							
Hospitalization -		Pneumonia	Professional				
MG QAM, 100							

Initial or Prolonged
MG Q NOON,
200 MG QHS)PO

Hypoxia
Pneumonitis
Pulmonary Fibrosis
Rib Fracture

Miacalcin C
Testosterone C

Date:05/05/98ISR Number: 3073884-XReport Type:Expedited (15-DaCompany Report #001-0945-980283
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG Initial or Prolonged (,DAILY), PER ORAL		Sepsis	Health Professional	Neurontin	PS		ORAL

Insulin C
Warfarin C
Darvocet C
Phoslo C

Date:05/05/98ISR Number: 3073885-1Report Type:Expedited (15-DaCompany Report #001-0945-980111
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 3600 MG (1200 Other MG, TID)		Fatigue	Health	Neurontin	PS		
		Hepatic Function Abnormal	Professional				
		Hepatitis C Insomnia Liver Function Test Abnormal		Accupril	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/06/98ISR Number: 3072734-5Report Type:Direct
 Age:43 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation		Gabapentin	PS	Parke-Davis	ORAL
600MG TID							
ORAL							
12.5 MG BID				Clozapine	SS	Novartic	ORAL
ORAL							
				Divalproex	C		
				Fluphenazine	C		
				Levothyroxine	C		
				Lithium Carbonate	C		
				Benzotropine	C		
				Lorazepam	C		

Date:05/08/98ISR Number: 3074736-1Report Type:Expedited (15-DaCompany Report #001-0945-980286
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Osteonecrosis	Health Professional Company Representative	Neurontin	PS		
Other							

Date:05/08/98ISR Number: 3074737-3Report Type:Expedited (15-DaCompany Report #001-0945-980292
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bronchioloalveolar	Consumer	Neurontin	PS		ORAL
Hospitalization - 1200 MG (300 MG,QID)		Carcinoma					
Initial or Prolonged							
Other		Drug Withdrawal Syndrome		Soma (Carisoprodol)	C		
Required		Dysgeusia		(Hydrocodone)	C		
Intervention to		Dyspnoea		Amitriptyline	C		
Prevent Permanent		Headache		Prozac (Fluoxetine)			

Impairment/Damage

Malaise
Pancreatitis
Pneumonia Mycoplasmal
Pyrexia
Tongue Oedema

Hydrochloride)

C

Date:05/11/98ISR Number: 3076680-2Report Type:Expedited (15-DaCompany Report #001-0945-980285
Age:58 YR Gender:Male I/FU:I

Outcome PT
Hospitalization - Bladder Disorder
Initial or Prolonged Burning Sensation
Feeling Of Body
Temperature Change
Hypoaesthesia
Nerve Conduction Studies
Abnormal
Paraesthesia
Polyneuropathy
Rash Erythematous
Rash Macular
Rash Pruritic
Red Blood Cell
Sedimentation Rate
Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Syncope

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
600 - 2400 MG		Literature	Neurontin	PS		ORAL
PER DAY PER		Health				
ORAL		Professional				
			Amitriptyline	C		
			Bupivacaine	C		
			Clonidine	C		
			Clonidine	C		
			Dilantin	C		
			Carbamazepine	C		
			Mexilatine	C		
			Desimpramine	C		

Date:05/12/98ISR Number: 3076586-9Report Type:Expedited (15-DaCompany Report #001-0073-980205
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Sodium Decreased Convulsion Inappropriate Antidiuretic Hormone Secretion Mental Disorder	Health Professional	Dilantin Neurontin	PS SS		

Date:05/13/98ISR Number: 3079264-5Report Type:Expedited (15-DaCompany Report #001-0945-970219
 Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 900-1200 MG Other (DAILY) PER ORAL		Blindness Condition Aggravated Eye Haemorrhage Macular Degeneration	Consumer	Neurontin Tegretol	PS C		ORAL

Trandate C
Hydrochlorothiaz C

Date:05/13/98ISR Number: 3079266-9Report Type:Expedited (15-DaCompany Report #001-0945-980311
Age:37 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Abdominal Discomfort
Initial or Prolonged	Dermatitis
Other	Diarrhoea
	Dizziness
	Ecchymosis
	Facial Palsy
	Fatigue
	Hypoaesthesia
	Loss Of Consciousness
	Mental Disorder
	Migraine
	Movement Disorder
	Nausea
	Palpitations
	Pyrexia
	Speech Disorder
	Syncope

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Tremor Weight Decreased				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Consumer	Neurontin	PS	
SEE TEXT, PER						ORAL
ORAL						

Date:05/19/98ISR Number: 3079442-5Report Type:Direct
Age:35 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dystonia		Gabapentin	PS		
				Phenytoin	SS		
				Vancomycin	C		
				Ranitidine	C		
				Docusate	C		
				Fes04	C		
				Ibuprofen	C		
				Codeinie	C		
				Ofloxacin	C		

Date:05/19/98ISR Number: 3081180-XReport Type:Expedited (15-DaCompany Report #001-0073-980221
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Dilantin	PS		ORAL
300 MG (100							
MG, TID), PER		Blood Potassium Decreased					
ORAL		Diarrhoea					
		Dysarthria		Neurontin	SS		ORAL
900 MG (300							
MG,TID), PER		Face Oedema					
ORAL		Fluid Retention					
		Oedema Peripheral		Glucotrol	C		
		Tongue Oedema		K-Dur	C		

Weight Increased

Date:05/21/98ISR Number: 3082334-9Report Type:Expedited (15-DaCompany Report #001-0945-980326
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2400 MG		Chest Pain	Health	Neurontin	PS		
Initial or Prolonged		Medication Error	Professional	Glucotrol	C		
				Lodine	C		
				Klonopin	C		
				Imitrex	C		
				Ultram	C		

Date:05/26/98ISR Number: 3082534-8Report Type:Expedited (15-DaCompany Report #001-0945-980320
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death UNKNOWN	1800 MG (600	Drowning	Health	Neurontin	PS		
MG,TID)		Overdose	Professional				
UNKNOWN		Sedation		Doxepin	SS		
		Syncope		Methadone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/26/98ISR Number: 3167664-4Report Type:Periodic
 Age:7 YR Gender:Male I/FU:I

Company Report #001-0916-970004

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 700 MG (100 Initial or Prolonged MG, 7 TIMES DAILY)	Condition Aggravated Convulsion Otitis Media	Health Professional	Carbamazepine	PS		
400 MG (100 MG, QID)			Neurontin (Gabapentin)	SS		ORAL
5 DAY			Ritalin (Methylphenidate Hydrochloride)	SS		ORAL

Date:05/26/98ISR Number: 3167667-XReport Type:Periodic
 Age:7 YR Gender:Male I/FU:I

Company Report #001-0916-970004

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 700 MG (100 Initial or Prolonged MG, 7 TIMES DAILY)	Convulsion Otitis Media	Health Professional	Carbamazepine	PS		
400 MG (100 MG QID)			Neurontin (Gabapentin)	SS		ORAL
5 DAY			Ritalin (Methylphenidate Hydrochloride)	SS		ORAL

Date:05/28/98ISR Number: 3084794-6Report Type:Expedited (15-DaCompany Report #001-0945-980346
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Neoplasm Malignant	Health	Neurontin	PS		ORAL
300 MG DAILY, Other PER ORAL			Professional				

Date:05/28/98ISR Number: 3084796-XReport Type:Expedited (15-DaCompany Report #001-0945-940034
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG DAILY, Initial or Prolonged PER ORAL		Blister	Health	Neurontin	PS		ORAL
Disability		Infection	Professional				
		Joint Stiffness Oedema Peripheral Skin Exfoliation Toxic Epidermal Necrolysis		Mysoline	C		

Date:06/02/98ISR Number: 3087573-9Report Type:Expedited (15-DaCompany Report #001-0945-980357
Age:42 YR Gender:Female I/FU:I

Outcome	PT
Other	Arthralgia Dermatitis Drug Hypersensitivity Dyspnoea Face Oedema Increased Appetite Myalgia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Weight Increased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
3600 MG (900 MG, QID), PER ORAL		Health Professional	Neurontin	PS		ORAL
			Stelazine	C		
			Xanax	C		

Date:06/02/98ISR Number: 3087579-XReport Type:Expedited (15-DaCompany Report #001-0073-980253
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 400 MG (,150 Initial or Prolonged MG IN AM AND HS,100 MG AT NOON), PER ORAL MONTHS 900 MG (300 MG.TID), PER ORAL. WEEKS		Arthralgia	Health Professional	Dilantin	PS		ORAL
		Dermatitis					
		Dermatomyositis					
		Drug Interaction					
		Hypotension					
		Muscle Atrophy		Neurontin	SS		ORAL

Date:06/03/98ISR Number: 3089637-2Report Type:Direct Company Report #
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG PO TID Initial or Prolonged		Anorexia		Gabapentin	PS		ORAL
		Dehydration					

Pyrexia

Date:06/04/98ISR Number: 3090234-3Report Type:Direct
Age:35 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema		Neurontin	PS	Parke Davis	
900MG TID(WITH DOSAGE GRADUALLY INCREASED FROM 300MG		Rash Maculo-Papular					
				Hydrocortisone	C		
				Multivitamin With Minerals	C		
				Sertraline	C		
				Ramitidine	C		
				Synthroid	C		
				Divalprox Na	C		
				Folic Acid	C		
				Thiamine	C		

Date:06/08/98ISR Number: 3092316-9Report Type:Expedited (15-DaCompany Report #001-0945-980159
Age: Gender:Female I/FU:I

Outcome	PT
Disability	Asthenia Blood Sodium Decreased Body Temperature Decreased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route		
500 MG		Dysphagia	Consumer	Neurontin	PS				
		Dyspnoea							
		Fatigue							
		Hypotension							
		Pain							
		Neurontin							
		SS							
		Paraesthesia						Tegretol	C
		Parosmia							
		Pneumonia							
Tinnitus									
Vertigo									
Vision Blurred									
Weight Decreased									

Date:06/11/98ISR Number: 3092515-6Report Type:Expedited (15-DaCompany Report #001-0073-980205

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Sodium Decreased	Health	Dilantin	PS		
		Condition Aggravated	Professional	Neurontin	SS		
		Convulsion					
		Inappropriate					
		Antidiuretic Hormone					
		Secretion					
		Mental Disorder					

Date:06/11/98ISR Number: 3092519-3Report Type:Expedited (15-DaCompany Report #001-0945-980203

Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Ammonia Increased	Health	Neurontin	PS		
		Confusional State	Professional				
		Liver Function Test	Company				
		Abnormal	Representative				

Date:06/18/98ISR Number: 3096884-2Report Type:Expedited (15-DaCompany Report #001-0945-980286

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Osteonecrosis	Health	Neurontin	PS		
UNKNOWN	UNKNOWN		Professional	Steroid	SS		
UNKNOWN	UNKNOWN		Company				
SEVERAL YEARS			Representative				
AGO							

Date:06/19/98ISR Number: 3096905-7Report Type:Expedited (15-DaCompany Report #034-0945-980003
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blister	Foreign	Neurontin	PS		ORAL
UNK, PER ORAL							
Initial or Prolonged		Dermatitis	Health				
			Professional				
			Company				
			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/19/98ISR Number: 3096906-9Report Type:Expedited (15-DaCompany Report #034-0945-980004
Age:70 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - UNK, PER ORAL	Blister	Foreign	Neurontin	PS		ORAL
Initial or Prolonged	Dermatitis	Health Professional Company Representative				

Date:06/23/98ISR Number: 3097741-8Report Type:Expedited (15-DaCompany Report #001-0945-980392
Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 1500 MG	Coma	Consumer	Neurontin	PS		ORAL
Initial or Prolonged (EVERY NIGHT), PER ORAL	Confusional State					
	Convulsion					
	Overdose					
	Pneumonia		Lasix	C		
	Tremor		Tylenol W/Codeine	C		
	Ulcer Haemorrhage		Corag	C		
			Lanoxin	C		
			Humalog	C		
			Humalin	C		
			Colchicine	C		

Date:06/24/98ISR Number: 3097902-8Report Type:Expedited (15-DaCompany Report #001-0945-980346
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death 300 MG	Gallbladder Cancer	Health Professional	Neurontin	PS		ORAL
Other (,DAILY), PER						

ORAL

Date:06/30/98ISR Number: 3100267-6Report Type:Expedited (15-DaCompany Report #001-0945-980417
Age:42 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion	Health	Neurontin			
Initial or Prolonged	Cough	Professional	(Gabapentin)	PS		
UNK, UNKNOWN						
	Lung Disorder		Tegretol			
	Rash Maculo-Papular		(Carbamazepine)	SS		
UNK, UNKNOWN						
			Sdz Gli 328	C		
			(Ganciclovir)	C		
			Phenobarb			
			(Phenobarbital			
			Sodium)	C		
			(Glipizide)	C		

Date:06/30/98ISR Number: 3100291-3Report Type:Expedited (15-DaCompany Report #044-0945-980036
Age:37 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Abdominal Pain Upper	Foreign	Neurontin			
Initial or Prolonged	Back Pain	Health	(Gabapentin)	PS		
2400 MG (800						
MG, TID) ,	Dysuria	Professional				
UNKOWN						
	Muscle Spasms					
	Urinary Incontinence		(Carbamazepine)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Labetalol) C

Date:07/01/98ISR Number: 3100340-2Report Type:Expedited (15-DaCompany Report #001-0945-980348
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2400 MG Initial or Prolonged (,DAILY), PER		Diarrhoea	Consumer	Neurontin	PS		ORAL
ORAL		Gastrointestinal Disorder Liver Function Test Abnormal Tinnitus					

Date:07/01/98ISR Number: 3100344-XReport Type:Expedited (15-DaCompany Report #002-0945-980011
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 900 MG (,DAILY), PER		Death	Foreign	Neurontin	PS		ORAL
ORAL		Health Professional					
				Lorazepam	C		
				Risperidone	C		

Date:07/01/98ISR Number: 3104400-1Report Type:Direct
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG PO TID		Headache		Gabapentin	PS		
		Sedation		Cisapride	C		
				Salsalate	C		
				Lansoprazol	C		
				Prazosin	C		

Date:07/01/98ISR Number: 3104422-0Report Type:Direct
 Age:43 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Condition Aggravated		Ativan	PS		
		Medication Error		Buspar	SS		
		Psychotic Disorder		Cardura	SS		
				Depakote	SS		
				Desyrel	SS		
				Haldol	SS		
				Mellaril	SS		
				Neurontin	SS		
				Propilsid	SS		
				Risperdal	SS		

Date:07/06/98ISR Number: 3102982-7Report Type:Expedited (15-DaCompany Report #001-0945-980423
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Interaction	Health	Neurontin	PS		
		Oesophageal Ulcer	Professional	Unspecified Antacid	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/06/98ISR Number: 3102984-0Report Type:Expedited (15-DaCompany Report #001-0945-980412

Age:7 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haematocrit Decreased	Health	Neurontin	PS		ORAL
300 MG		Haemoglobin Decreased	Professional				
(,EVERY 8							
HOURS), PER							
ORAL							

Tetrabenazine	C
Lorazepam	C
Chloral Hydrate	C
Acetaminophen	C
Ibuprofen	C
Vancomycin	C
Ceftazidime	C
Cefazolin	C

Date:07/06/98ISR Number: 3184947-2Report Type:Periodic Company Report #91692

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression	Other	Klonopin Tablets			
OCCLUSIVE		Headache		(Clonazepam)	PS		
DRESSING	2.0000	Mood Altered					
X PER DAY		MG 1.0					
		Sedation					
				Neurontin			
				(Gabapentin)	SS		ORAL

Date:07/07/98ISR Number: 3102593-3Report Type:Expedited (15-DaCompany Report #001-0945-980283

Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 300 MG	Condition Aggravated	Health	Neurontin	PS	ORAL
Initial or Prolonged (DAILY), PER	Neuropathy Peripheral	Professional			
ORAL	Pain In Extremity				
	Sepsis		Insulin	C	
			Warfarin	C	
			Darvocet	C	
			Phoslo	C	

Date:07/07/98ISR Number: 3102594-5Report Type:Expedited (15-DaCompany Report #001-0945-980422
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Health	Neurontin	PS		ORAL
1500 MG (300		Petit Mal Epilepsy	Professional				
MG, 5X DAILY)							
PER ORAL							
ONCE WEEKLY				DiFlucan	SS		
				Diphenhydramine	C		
				Bactrim	C		
				Amitriptyline	C		
				Acetaminophen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/07/98ISR Number: 3102596-9Report Type:Expedited (15-DaCompany Report #001-0945-980421
Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300 MG Initial or Prolonged (DAILY), PER ORAL	Disorientation Monoparesis	Consumer	Neurontin	PS		ORAL

Date:07/07/98ISR Number: 3103871-4Report Type:Direct Company Report #
Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300 MG TID PO 4 DAY Initial or Prolonged	Lethargy Mental Impairment Tremor		Neurontin	PS		ORAL

Date:07/10/98ISR Number: 3104309-3Report Type:Expedited (15-DaCompany Report #033-0945-980019
Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 1200 MG DAILY Hospitalization - Initial or Prolonged	Cardiac Arrest Convulsion Pulmonary Embolism	Foreign Health Professional	Neurontin Tegritol	PS SS		

Date:07/13/98ISR Number: 3108784-XReport Type:Direct Company Report #
Age:80 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG PO QD 1 WK Initial or Prolonged	Choreoathetosis Tardive Dyskinesia Vomiting		Neurontin Premarin Levothyroxine	PS C C		ORAL

Digoxin	C
Duragesic	C
Oxycodan	C
Compazine	C
Ultram	C
Zoloft	C
Dss	C
Senna	C
Scopolamine Patch	C

Date:07/14/98ISR Number: 3104962-4Report Type:Expedited (15-DaCompany Report #001-0945-980449
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Gastrinoma	Consumer	Neurontin	PS		ORAL
1600 MG (800		Migraine					
MG, BID), PER		Retinal Vein Thrombosis					
ORAL		Vision Blurred		Darvon			
		Visual Acuity Reduced		(Dextropropoxyphene Hydrochloride)	C		
				Asa (Acetylsalicylic Acid)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/14/98ISR Number: 3104964-8Report Type:Expedited (15-DaCompany Report #034-0945-980003

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300 MG	Duration Dermatitis	Foreign	Gabapentin	PS		ORAL
Initial or Prolonged (DAILY) PER	Eosinophilia	Health				
ORAL	Phantom Pain	Professional				
	Pruritus	Company	(Ipratropium			
	Rash Maculo-Papular	Representative	Bromide)	C		
	Rash Vesicular		(Lorazepam)	C		
	Thrombocythaemia		(Salbutamol)	C		
			(Triflusal)	C		
			"Budenoside"	C		
			"Oxpentilfylline:"	C		

Date:07/14/98ISR Number: 3104967-3Report Type:Expedited (15-DaCompany Report #034-0945-980004

Age:73 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300 MG	Duration Blister	Foreign	Neurontin	PS		ORAL
Initial or Prolonged (DAILY) PER	Dermatitis	Health				
ORAL	Pruritus	Professional				
	Pyrexia	Company	Carbamazepine	C		
	Rash Maculo-Papular	Representative	Amitriptyline	C		

Date:07/16/98ISR Number: 3105982-6Report Type:Expedited (15-DaCompany Report #001-0945-980447

Age:65 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 3200 MG (800	Duration Inappropriate	Health	Neurontin	PS		ORAL
Initial or Prolonged MG,QID), PER	Antidiuretic Hormone	Professional				

Secretion

ORAL

Multiple Other Medications

C

Date:07/16/98ISR Number: 3105983-8Report Type:Expedited (15-DaCompany Report #001-0945-980392
Age:47 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1500 MG Initial or Prolonged (,EVERY NIGHT), PER	Coma Confusional State Convulsion Medication Error	Consumer	Neurontin	PS		ORAL
ORAL	Pneumonia Tremor Ulcer Haemorrhage		Lasix Tylenol W/Codeine Corag Lanoxin Humalog Humulin (Colchine)	C C C C C C C		

Date:07/16/98ISR Number: 3105987-5Report Type:Expedited (15-DaCompany Report #001-0945-980357
Age:42 YR Gender:Female I/FU:F

Outcome	PT
Other	Arthralgia Dermatitis Drug Hypersensitivity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Dyspnoea Face Oedema Increased Appetite	Report Source	Product	Role	Manufacturer	Route
3600 MG (900 MG, QID), PER ORAL		Myalgia Weight Increased	Health Professional	Neurontin	PS		ORAL
				Stelazine Xanax	C C		

Date:07/17/98ISR Number: 3108232-XReport Type:Direct
Age:60 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 300 MG ONE Intervention to CAP, BY MOUTH Prevent Permanent Impairment/Damage		Bradycardia Hypotension		Gabapentin	PS		
				Lovastatin Docusate Hydrocortisone	C C C		

Date:07/20/98ISR Number: 3107255-4Report Type:Expedited (15-DaCompany Report #001-0945-980451
Age:8 DY Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 4400 MG Initial or Prolonged (,DAILY) SEE Other TEXT FOR THERAPY DATES 600 MG (,DAILY) SEE		Complications Of Maternal Exposure To Therapeutic Drugs Dehydration Haemorrhage Intracranial	Health Professional	Neurontin	PS		
				Tegretol	SS		

TEXT FOR

THERAPY DATES

(Folic Acid) C

Date:07/21/98ISR Number: 3107303-1Report Type:Expedited (15-DaCompany Report #001-0073-980221
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (100 MG, TID), PER ORAL		Blood Potassium Decreased Diarrhoea Dysarthria	Consumer	Dilantin	PS		ORAL
900MG (100 MG, TID) PER ORAL		Eyelid Oedema Face Oedema Fluid Retention Memory Impairment		Neurontin Capsules 100 Mg (Gabapentin)	SS		ORAL
		Oedema Peripheral Tongue Oedema Weight Increased		Glucotrol (Glipizide) K-Dur (Potassium Chloride)	C C		

Date:07/28/98ISR Number: 3110720-7Report Type:Expedited (15-DaCompany Report #001-0945-980292
Age:51 YR Gender:Male I/FU:F

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
1200 MG (300 MG QID) PER ORAL		Adenoma Benign	Health	Neurontin	PS		ORAL
		Chills	Professional				
		Dysgeusia					
		Dyspnoea		Soma	C		
		Headache		Hydrocodone	C		
		Lung Infiltration		Amitriptyline	C		
		Pancreatitis		Prozac	C		
		Pneumonia Mycoplasmal					
		Pyrexia					
		Tongue Oedema					

Date:07/28/98ISR Number: 3110729-3Report Type:Expedited (15-DaCompany Report #001-0073-980221

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Potassium Decreased	Health	Dilantin	PS		ORAL
300 MG (100 MG TID) PER ORAL		Diarrhoea	Professional				
		Dysarthria					
900 MG (300 MG TID) PER ORAL		Eyelid Oedema		Neurontin	SS		ORAL
		Face Oedema					
		Fluid Retention					
		Memory Impairment		Glucotrol	C		
		Oedema Peripheral		K-Dur	C		
		Tongue Oedema					
		Weight Increased					

Date:07/31/98ISR Number: 3111333-3Report Type:Expedited (15-DaCompany Report #044-0945-980059

Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Pulmonary Embolism	Foreign	Neurontin	PS		ORAL
3200MG			Health				
(1600MG,BID),			Professional				
PER ORAL				Zyprexa	SS		ORAL
100MG (,							
DAILY), PER							
ORAL							
				Camcolit	C		
				Epilim	C		
				Dolmatil	C		
				Tertroxin	C		
				Zimovane	C		

Date:07/31/98ISR Number: 3252994-8Report Type:Periodic Company Report #001-0991-980850
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Consumer	Rezulin Tablets 400			
Other		Weight Increased		Mg (Troglitazone)	PS		ORAL
400 MG							
(DAILY) PER							
ORAL							
				Neurontin Capsules			
300 MG, PER				300 Mg (Gabapentin)	SS		ORAL
ORAL							
				Prilosec	C		
				Mevacor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Synthroid	C
Humulin	C
Lasix	C
Quinine	C
Percocet	C

Date:08/04/98ISR Number: 3116637-6Report Type:Expedited (15-DaCompany Report #001-0945-980543
 Age:26 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - (,TID), PER Initial or Prolonged ORAL	Abdominal Pain Anxiety Condition Aggravated Lower Limb Fracture Pancreatitis Staphylococcal Infection	Consumer	Neurontin Lamictal Paxil Zyprexa Ancef	PS C C C C		ORAL

Date:08/05/98ISR Number: 3113424-XReport Type:Expedited (15-DaCompany Report #033-0945-980020
 Age:20 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1200 MG Initial or Prolonged (DAILY), PER ORAL	Pyrexia Rash Maculo-Papular Toxic Skin Eruption	Foreign Health Professional	Neurontin Alepsal Gardenal Lansoyl Di-Antalvic	PS C C C C		ORAL

Date:08/05/98ISR Number: 3113742-5Report Type:Direct Company Report #
 Age:79 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Hospitalization - Creatinine Renal
Initial or Prolonged Clearance Decreased
Oedema Peripheral

Gabapentin PS
Losartan SS
Troglitrazone SS

Date:08/07/98ISR Number: 3114479-9Report Type:Expedited (15-DaCompany Report #001-0945-980480
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	900MG	Amnesia	Consumer	Neurontin	PS		ORAL
(300MG,TID),		Aphasia					
PER ORAL		Arthritis					
		Catatonia		Dilantin (Phenytoin			
		Confusional State		Sodium)	C		
				Unspecified			
				Antihypertensive			
				Meds	C		
				Depakote	C		
				Clonidine	C		
				Prednisone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/07/98ISR Number: 3114481-7Report Type:Expedited (15-DaCompany Report #001-0945-980533

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1500MG, Initial or Prolonged DAILY, PER ORAL	Pulmonary Thrombosis	Health Professional	Neurontin	PS		ORAL
			Phenytoin Coumadin (Warfarin Sodium)	C C		

Date:08/19/98ISR Number: 3118959-1Report Type:Expedited (15-DaCompany Report #002-0945-980014

Age:60 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 300 MG DAILY PER ORAL	Gait Disturbance Lethargy	Foreign Health	Neurontin	PS		ORAL
	Liver Function Test Abnormal Neuroleptic Malignant Syndrome Parkinson'S Disease Pyrexia	Professional	Sinemet (Levodopa, Carbidopa)	C		

Date:08/19/98ISR Number: 3118964-5Report Type:Expedited (15-DaCompany Report #001-0945-980451

Age:8 DY Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - UNKNOWN 4400 MG DAILY Initial or Prolonged UNKNOWN	Cerebral Haemorrhage Neonatal	Health Professional	Neurontin	PS		
Other UNKNOWN 600 MG DAILY	Complications Of Maternal		Tegretol	SS		

UNKNOWN Exposure To Therapeutic
 Drugs (Folic Acid) C
 Dehydration

Date:08/19/98ISR Number: 3119094-9Report Type:Direct Company Report #
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Abnormal Dreams		Gabapentin	PS		ORAL
300 MG TID PO							
Intervention to		Increased Appetite		Lorazepam	C		
Prevent Permanent		Platelet Count Decreased		Psyllium	C		
Impairment/Damage		Red Blood Cell Count		Premarin	C		
		Decreased		Paroxetine	C		
		Thirst		Furosemide	C		
		White Blood Cell Count		Hydroxyzine	C		
		Decreased		Thiamine Hcl	C		
				Multi-Vitamins	C		

Date:08/19/98ISR Number: 3119095-0Report Type:Direct Company Report #
 Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Aggression		Gabapentin	PS		ORAL
100 MG TID PO							
Intervention to		Agitation		Multi-Vitamins	C		
Prevent Permanent		Hallucination					
Impairment/Damage							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/19/98ISR Number: 3119097-4Report Type:Direct
Age:76 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Required	Fall		Gabapentin	PS		ORAL
300 MG/QD/PO						
Intervention to	Gait Disturbance		Thiothixene	C		
Prevent Permanent			Multi-Vitamins	C		
Impairment/Damage			Benztropine Mesylate	C		
			Lactulose	C		

Date:08/20/98ISR Number: 3119479-0Report Type:Expedited (15-DaCompany Report #033-0945-980026
Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Fall	Foreign	Neurontin	PS		ORAL
900 MG (300						
Initial or Prolonged	Femoral Neck Fracture	Health				
MG, TID)						
	Muscular Weakness	Professional	Lioresal (Baclofen)	C		

Date:08/26/98ISR Number: 3121928-9Report Type:Expedited (15-DaCompany Report #001-0945-980578
Age:72 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Asthenia	Consumer	Neurontin	PS		ORAL
200 MG						
Initial or Prolonged	Confusional State					
(DAILY), PER						
ORAL	Coordination Abnormal					
	Dehydration		Sinemet	C		
	Fatigue		Empirin	C		
	Haemoglobin Decreased		Tylenol	C		
	Nightmare		Klonopin	C		
	Platelet Count Decreased		Prozac	C		
			Amantadine	C		
			Mirapex	C		

Date:08/27/98ISR Number: 3122068-5Report Type:Direct
Age:86 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG TID Initial or Prolonged	Confusional State Coordination Abnormal Dizziness Dysarthria		Neurontin	PS		

Date:09/01/98ISR Number: 3124222-5Report Type:Expedited (15-DaCompany Report #049-0945-980016
Age:1 DY Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other IN UTERO	Anaemia	Foreign	Neurontin	PS		
EXPOSURE	Apgar Score Low	Health				
3 MG DAILY- IN UTERO	Complications Of Maternal Exposure To Therapeutic Drugs	Professional	Valproate	SS		
EXPOSURE	Neonatal Apnoeic Attack					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/98ISR Number: 3124225-0Report Type:Expedited (15-DaCompany Report #049-0945-980017

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - PER ORAL Initial or Prolonged 3 GM DAILY PER ORAL	Convulsion	Foreign Health Professional	Neurontin Valproate	PS SS		ORAL ORAL

Date:09/01/98ISR Number: 3124785-XReport Type:Expedited (15-DaCompany Report #001-0945-980608

Age:22 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 2400 MG (900 MG QAM, 600 MG QNOON AND QPM, 300 MG QHS) IN UTERO 400 MG DAILY, IN UTERO EXPOSURE	Arthralgia Breast Hyperplasia Complications Of Maternal Exposure To Therapeutic Drugs Hirsutism Migraine Precocious Puberty	Consumer	Neurontin Felbatol	PS SS		

Date:09/03/98ISR Number: 3125675-9Report Type:Direct

Company Report #

Age:34 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300MG TID PO Initial or Prolonged	Dyspnoea Tongue Coated Tongue Disorder		Neurantin	PS		ORAL

Urticaria

Date:09/04/98ISR Number: 3126807-9Report Type:Expedited (15-DaCompany Report #001-0945-980618
 Age:52 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1500 MG (500	Coronary Artery Occlusion	Health	Neurontin	PS		ORAL
Initial or Prolonged MG, TID), PER	Inappropriate	Professional				
Other ORAL	Antidiuretic Hormone					
Required Intervention to Prevent Permanent Impairment/Damage	Secretion		Coumadin	C		
			Trazodone	C		
			Lasix	C		
			Serzone	C		
			Rezulin	C		

Date:09/08/98ISR Number: 3127826-9Report Type:Expedited (15-DaCompany Report #033-0945-980020
 Age:20 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1200 MG	Acne	Foreign	Neurontin	PS		ORAL
Initial or Prolonged (DAILY), PER	Pyrexia	Health				
ORAL	Rash Maculo-Papular	Professional				
	Toxic Skin Eruption		Alepsal	C		
			Gardenal	C		
			Lansoyl	C		
			Di-Antalvic	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/10/98ISR Number: 3127350-3Report Type:Expedited (15-DaCompany Report #001-0945-980578
 Age:72 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 200 MG DAILY Initial or Prolonged PER ORAL	Asthenia	Consumer	Neurontin	PS		ORAL
	Confusional State					
	Coordination Abnormal		Sinemet	C		
	Dehydration		Tylenol	C		
	Dyskinesia		Klonopin	C		
	Fatigue		Prozac	C		
	Feeding Disorder		Amantadine	C		
	Haemoglobin Decreased		Mirapex	C		
	Nightmare					
	Platelet Count Decreased					
	Vitamin B12 Decreased					

Date:09/11/98ISR Number: 3128706-5Report Type:Direct Company Report #
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 200 MG TID Initial or Prolonged ORAL	Bleeding Time Prolonged		Gabapentin	PS		ORAL
	Blood Fibrinogen					
	Increased		Ciprofloxacin	C		
	Haematocrit Decreased		Ranitidine	C		
	Haemoptysis		Sulfameth/Trimeth	C		
	Thrombocytopenia					

Date:09/13/98ISR Number: 3134279-3Report Type:Expedited (15-DaCompany Report #001-0945-980447
 Age:65 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 3200 MG (800 MG QID) PER	Inappropriate Antidiuretic Hormone Secretion	Health Professional	Neurontin (Gabapentin)	PS		ORAL

ORAL

Multiple Other Medications (Unspecified) C

Date:09/15/98ISR Number: 3237561-4Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #8-98175-071A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaemia	Consumer	Duract	PS		ORAL
ORAL		Cough		Neurontin	SS		
				Ergomar	C		
				Imipiramine	C		
				Talacen	C		
				Vicodin	C		
				Zoloft	C		

Date:09/16/98ISR Number: 3131206-XReport Type:Direct
Age:38 YR Gender:Female I/FU:I

Company Report #

Outcome
Required
Intervention to
Prevent Permanent

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Camcolit (Lithium
Carbonate) C
Epilim (Valproate
Sodium) C
Dolmatil (Sulpiride) C
Tertroxin
(Liothyronine
Sodium) C
Zimovane (Zopiclone) C

Date:09/23/98ISR Number: 3134277-XReport Type:Expedited (15-DaCompany Report #001-0945-980543
Age:26 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - UNKNOWN TID	Abdominal Pain	Consumer	Neurontin	PS		ORAL
Initial or Prolonged PER ORAL	Lower Limb Fracture					
	Pancreatitis		Lamictal	C		
	Staphylococcal Infection		Paxil	C		
			Zyprexa (Olanzapine)	C		
			Ancef (Cefazolin Sodium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/29/98ISR Number: 3136219-XReport Type:Expedited (15-DaCompany Report #001-0945-98-649
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Pneumonia	Health	Neurontin	PS		ORAL
Hospitalization - 2400 MG (800 MG TID), PER Initial or Prolonged ORAL		Respiratory Failure	Professional				

Date:10/01/98ISR Number: 3137324-4Report Type:Expedited (15-DaCompany Report #033-0945-980037
Age:1 DY Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly 3 TABLET		Complications Of Maternal	Foreign	Neurontin	PS		
Other (DAILY)		Exposure To Therapeutic	Health				
PLACETAL		Drugs	Professional				
		Foetal Distress Syndrome		Depamide	C		
		Foetal Growth Retardation		(Valpromide)			
		Neonatal Respiratory		Tranxene			
		Distress Syndrome		(Clorazepate Dipotassium)	C		

Date:10/01/98ISR Number: 3137327-XReport Type:Expedited (15-DaCompany Report #049-0945-980016
Age:1 DY Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - IN UTERO		Apgar Score Low	Foreign	Neurontin	PS		
Initial or Prolonged EXPOSURE		Blood Incompatibility	Health				
Congenital Anomaly 3 GM DAILY-		Haemolytic Anaemia Of	Professional	Valproate	SS		
Other IN UTERO		Newborn					

Complications Of Maternal

EXPOSURE

Exposure To Therapeutic
Drugs
Congenital Central
Nervous System Anomaly
Convulsion
Premature Labour

Date:10/01/98ISR Number: 3137329-3Report Type:Expedited (15-DaCompany Report #049-0945-980017

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - PER ORAL		Complications Of Maternal	Foreign	Neurontin	PS		ORAL
Initial or Prolonged 3 GM DAILY		Exposure To Therapeutic Drugs	Health Professional	Valproate	SS		ORAL
PER ORAL		Epilepsy		Diazepam	SS		
INTRAVENOUS	INTRAVENOUS	Premature Labour		Felbamate	SS		
INTRAVENOUS	INTRAVENOUS						

Date:10/01/98ISR Number: 3260849-8Report Type:Periodic Company Report #001-0073-970439

Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300/400 MG (QHS), PER ORAL		Convulsion Drug Level Below Therapeutic Pneumonia	Health Professional	Dilantin Kalseals 100 Mg (Phenytoin Sodium)	PS		ORAL
3600 MG DAILY				Neurontin 400 Mg (Gabapentin)	SS		ORAL

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Freedom Of Information (FOI) Report

(400 MG) PER

ORAL

Cardizem C
Isordil C

Date:10/02/98ISR Number: 3137732-1Report Type:Expedited (15-DaCompany Report #001-0073-980398
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 500 MG DAILY Initial or Prolonged PER ORAL		Convulsion	Consumer	Dilantin Kapseals	PS		ORAL
UNKNOWN 2400 MG (400 MG DAILY UNKNOWN		Erectile Dysfunction Medication Error Myocardial Infarction Pulmonary Oedema		Neurontin	SS		
				Folic Acid Norvasc (Amlodipine Besilate) Lipitor (Atorvastatin)	C C C C		

Date:10/05/98ISR Number: 3138779-1Report Type:Expedited (15-DaCompany Report #001-0945-980412
Age:7 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG (EVERY Initial or Prolonged 8 HOURS), PER Other ORAL	8 HR	Anaemia Condition Aggravated Haematocrit Decreased Haemoglobin Decreased Haemorrhage	Health Professional	Neurontin	PS		ORAL
				Tetrabenazine Lorazepam Chloral Hydrate Acetaminophen Ibuprofen	SS C C C C		

Vancomycin C
 Ceftazidime C
 Cefazolin C

Date:10/06/98ISR Number: 3138768-7Report Type:Expedited (15-DaCompany Report #001-0945-980665
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	3600 MG DAILY	Abortion Spontaneous	Health Professional	Neurontin	PS		

Date:10/06/98ISR Number: 3138771-7Report Type:Expedited (15-DaCompany Report #001-0945-980691
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1200 MG (400 MG, TID) PER	Cardiac Arrest Dysphonia	Consumer	Neurontin	PS		ORAL

Prilosec
 (Omeprazole) C
 Klonopin
 (Clonazepam) C
 Welbutrin
 (Amfebutamone
 Hydrochloride) C
 Prolupsid

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Cisapride) C
 Buspar (Buspirone
 Hydrochloride) C

Date:10/06/98ISR Number: 3138774-2Report Type:Expedited (15-DaCompany Report #049-0945-980019
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agranulocytosis	Foreign	Neutrontin	PS		ORAL
PER ORAL			Health Professional				

Date:10/07/98ISR Number: 3139352-1Report Type:Expedited (15-DaCompany Report #9905902
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Aphasia	Foreign	Gabitril	PS		ORAL
25.000 MG PO							
Initial or Prolonged		Convulsion					
QD							
Other		Neurosis		Gabapentin	SS		
		Obsessive Thoughts		Carbamazepine	C		
		Status Epilepticus					

Date:10/13/98ISR Number: 3141708-8Report Type:Expedited (15-DaCompany Report #001-0945-980692
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Benign Hepatic Neoplasm	Health	Neurontin	PS		ORAL
1500 MG							
Initial or Prolonged		Deafness	Professional				
(DAILY) PER							
Disability		Haemangioma					
ORAL							
		Ototoxicity		Lithium	C		
				Buspirone	C		
				Carbamazepine	C		

Date:10/13/98ISR Number: 3142422-5Report Type:Expedited (15-DaCompany Report #001-0945-980693
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1200 MG DAILY Initial or Prolonged PER ORAL	Anxiety Myocardial Infarction Paraesthesia Rash Erythematous	Consumer	Neurontin Darvocet-N Humulin-R Humulin-N	PS C C C		ORAL

Date:10/13/98ISR Number: 3142473-0Report Type:Expedited (15-DaCompany Report #033-0945-980038
Age: Gender:Not SpecifiI/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death PLACENTAL	Complications Of Maternal Exposure To Therapeutic Drugs Intra-Uterine Death Neoplasm	Foreign Health Professional	Neurontin	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/13/98ISR Number: 3142512-7Report Type:Expedited (15-DaCompany Report #033-0945-980034

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Foreign	Neurontin	PS		ORAL
ORAL		Complications Of Maternal Exposure To Therapeutic Drugs Mass	Health Professional				

Date:10/15/98ISR Number: 3143029-6Report Type:Expedited (15-DaCompany Report #001-0945-980578

Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 200 MG		Confusional State	Consumer	Neurontin	PS		ORAL
Initial or Prolonged (DAILY), PER		Coordination Abnormal					
ORAL		Dehydration					
		Fatigue		Sinemet	C		
		Haemoglobin Decreased		Empirin	C		
		Muscular Weakness		Tylenol	C		
		Nightmare		Klonopin	C		
		Platelet Count Decreased		Prozac	C		
		Vitamin B12 Decreased		Amantadine	C		
				Mirapex	C		

Date:10/16/98ISR Number: 3143128-9Report Type:Expedited (15-DaCompany Report #033-0945-980034

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Foreign	Neurontin			
2400MG			Health	(Gabapentin)	PS		ORAL
(DAILY) PER			Professional				

ORAL

Tegretol

C

Date:10/16/98ISR Number: 3143129-0Report Type:Expedited (15-DaCompany Report #033-0945-980038

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Complications Of Maternal Exposure To Therapeutic	Foreign Health	Neurontin (Gabapentin)	PS		OTHER
2400MG (DAILY)		Drugs	Professional				
PLACENTTAL		Foetal Disorder					
		Neoplasm		Tegretol	C		

Date:10/19/98ISR Number: 3144271-0Report Type:Direct

Company Report #

Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ALPHA		Cognitive Disorder		Tegretol	PS		ORAL
CAPSULES PO, BETA TABLETS		Condition Aggravated					
PO, DOUBLE		Depressed Level Of Consciousness					
BLINDED ALPHA				Gabapentin	SS		ORAL
CAPSULES PO, BETA TABLETS							
PO, DOUBLE							

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BLINDED

ALPHA

CAPSULES PO,

BETA TABLETS

PO, DOUBLE

BLINDED

Lamotrigine

SS

ORAL

Date:10/28/98ISR Number: 3149020-8Report Type:Expedited (15-DaCompany Report #001-0945-980759

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Drug Interaction	Health	Neurontin	PS		
Initial or Prolonged		Drug Toxicity	Professional	Tylenol	SS		
		Gastrointestinal Motility Disorder					
		Intentional Misuse					

Date:10/29/98ISR Number: 3149662-XReport Type:Expedited (15-DaCompany Report #001-0945-980748

Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Depression	Health	Neurontin	PS		ORAL
1200 MG (600		Euphoric Mood	Professional				

MG BID) PER

ORAL

Prozac

C

Ultram

C

Lodine

C

Date:10/29/98ISR Number: 3149665-5Report Type:Expedited (15-DaCompany Report #001-0945-980760

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin	PS		ORAL
900 MG (300		Drug Interaction					
MG TID) PER		Meningitis					
ORAL		Oedema Peripheral		Dilantin	SS		
200 MG IN AM;							
300 MG IN PM							
				Estrace	C		
				Prilosec	C		
				Ms Contin	C		

Date:10/30/98ISR Number: 3150210-9Report Type:Expedited (15-DaCompany Report #9834536
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Arthralgia	Consumer	Vistaril	PS		ORAL
50.00 MG		Hypersensitivity					
Intervention to		Paralysis					
TOTAL:PRN:ORA							
Prevent Permanent							
L							
Impairment/Damage				Zoloft	SS		ORAL
50.00 MG							
TOTAL:DAILY:O							
RAL							
600.00 MG				Neurontin	SS		ORAL
TOTAL:TID:ORA							
L							
				Synthroid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/30/98ISR Number: 3150242-0Report Type:Expedited (15-DaCompany Report #001-0945-980749

Age:5 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Benign Intracranial		Neurontin	PS		ORAL
300 MG		Hypertension					
(DAILY), PER							
ORAL				Diamox	C		

Date:11/02/98ISR Number: 3150275-4Report Type:Expedited (15-DaCompany Report #1998-002643

Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anaemia	Foreign	Viramune	PS		ORAL
200 MG/PO		Ascites	Study	Nelfinavir	SS		ORAL
Initial or Prolonged		Asthenia	Health	Indinavir	SS		ORAL
2000 MG/PO		Balance Disorder	Professional	Nevrontin	SS		ORAL
2400 MG/PO		Blood Alkaline		Acyclovir	C		
1200 MG/PO		Phosphatase Increased		Septra Ds	C		
		Blood Creatinine		Megace	C		
		Increased		Losec	C		
		Blood Lactate		Stemetil	C		
		Dehydrogenase Increased		Ms Contin	C		
		Blood Sodium Decreased		Clarithromycin	C		
		Depressed Level Of		Ethambutol	C		
		Consciousness		Fluconazole	C		
		Diarrhoea					
		Dyspnoea					
		Fall					
		Haemoglobin Decreased					
		Heart Rate Increased					
		Hypotension					
		Malaise					
		Metabolic Acidosis					
		Nausea					
		Oedema Peripheral					

Oral Intake Reduced
Ph Urine Decreased
Proteinuria
Renal Failure Acute
Renal Tubular Disorder
Sedation
Vomiting

Date:11/02/98ISR Number: 3151612-7Report Type:Expedited (15-DaCompany Report #001-0945-980716
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Neurontin	PS		

Date:11/02/98ISR Number: 3152104-1Report Type:Expedited (15-DaCompany Report #9834323
Age:72 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Balance Disorder Drug Interaction Nervous System Disorder	Foreign 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
50.00 MG		Diflucan	PS		ORAL
TOTAL;					
DAILY;	ORAL				
400.00 MG		Gabapentin	SS		ORAL
TOTAL;	ORAL				
ORAL		Prednisolone	SS		ORAL

Date:11/02/98ISR Number: 3268071-6Report Type:Periodic
Age:71 YR Gender:Female I/FU:F

Company Report #001-0991-980850

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oedema Peripheral Weight Increased	Consumer	Rezulin Tablets 400 Mg (Troglitazone)	PS		ORAL
400 MG							
(,DAILY), PER							
ORAL							
				Neurontin Capsules 300 Mg (Gabapentin)	SS		ORAL
300 MG, PER							
ORAL							
				Prilosec (Omeprazole)	C		
				Mevacor (Lovastatin)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Humulin (Insulin Human Injection, Isophane)	C		
				Lasix (Furosemide)	C		

(Quinine) C
 Percocet
 (Paracetamol,
 Oxycodone
 Hydrochloride,
 Oxycodone C

Date:11/02/98ISR Number: 3271014-2Report Type:Periodic Company Report #001-0991-981543
 Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG (, DAILY), PER ORAL		Oedema Peripheral Weight Increased	Consumer	Rezulin Tablets 200 Mg (Troglitazone)	PS		ORAL
600 MG (,TID), PER ORAL				Neurontin (Gabapentin)	SS		ORAL
				Humulin	C		
				Dyazide	C		
				Hyzaar	C		
				Catapres	C		
				Indocin	C		
				Lipitor	C		
				Zoloft	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/05/98ISR Number: 3152772-4Report Type:Expedited (15-DaCompany Report #001-0945-980783

Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 400 MG (100 Initial or Prolonged MG, QID), PER ORAL	Diarrhoea Dizziness Orthostatic Hypotension	Consumer	Neurontin	PS		ORAL
			Cozaar	C		
			Insulin	C		
			Nph Insulin	C		
			Proamatine	C		
			Lasix	C		

Date:11/06/98ISR Number: 3153238-8Report Type:Direct

Company Report #

Age:46 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - UNKNOWN UNKNOWN 3 DAY Initial or Prolonged	Dermatitis Face Oedema Pruritus	Health Professional	Neurontin Dilantin	PS C		

Date:11/09/98ISR Number: 3154270-0Report Type:Expedited (15-DaCompany Report #001-0073-980398

Age:63 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 500 MG (, Initial or Prolonged DAILY), PER ORAL 2400 MG (2400 MG, DAILY)	Convulsion Drug Interaction Erectile Dysfunction Myocardial Infarction Pulmonary Oedema	Consumer	Dilantin Neurontin (Folic Acid)	PS SS C		ORAL

Norvasc C
Lipitor C
Neurontin C
.. C

Date:11/09/98ISR Number: 3154759-4Report Type:Expedited (15-DaCompany Report #001-0945-980665
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health	Neurontin			
		Complications Of Maternal	Professional	(Gabapentin)	PS		
UNKNOWN	3600 MG DAILY						
		Exposure To Therapeutic					
UNKNOWN		Drugs					

Date:11/11/98ISR Number: 3155171-4Report Type:Direct Company Report #
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Thrombocytopenia		Neurontin	PS	Parke-Davis	ORAL
300MG QID							

Date:11/13/98ISR Number: 3157205-XReport Type:Expedited (15-DaCompany Report #001-0945-980692
Age:44 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

Disability

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1500 MG DAILY		Benign Hepatic Neoplasm Condition Aggravated	Health Professional	Neurontin (Gabapentin)	PS		ORAL
PER ORAL		Deafness Neurosensory					
		Haemangioma		Lithium	C		
		Ototoxicity		Buspirone	C		
				Carbamazepine	C		

Date:11/16/98ISR Number: 3158141-5Report Type:Expedited (15-DaCompany Report #049-0945-980019
Age:71 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agranulocytosis	Foreign	Neurontin	PS		ORAL
1200 MG Hospitalization - (DAILY), PER Initial or Prolonged ORAL		Cardiac Arrest	Health Professional				
Other 40 MG (DAILY), PER ORAL		Clonic Convulsion		Antra	SS		ORAL
		Condition Aggravated					
		Convulsion					
6 MG (DAILY), PER ORAL		Csf Protein Increased		Arelix	SS		ORAL
		Delirium					
		Depressed Level Of Consciousness		Lopirin	C		
		Depression		Phenhydan	C		
		Diarrhoea		Xanef	C		
		Encephalopathy		Lopedium	C		
		Gastrointestinal Haemorrhage		Lasix	C		
		Infection					
		Multi-Organ Failure					
		Pneumonia					
		Proteinuria					

Psychotic Disorder
 Renal Failure
 Respiratory Failure
 Sepsis
 Staphylococcal Sepsis
 Whipple'S Disease

Date:11/19/98ISR Number: 3160681-XReport Type:Direct
 Age: Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1 TID Initial or Prolonged	Cystitis		Neurontin	PS	Parke-Davis	ORAL
	Fibroadenoma Of Breast		Darvocet	C		
	Head Injury		Percocet	C		
	Lung Disorder		Mebaral	C		
	Pneumonia		Tenormin	C		
	Tricuspid Valve Incompetence					

Date:11/20/98ISR Number: 3160861-3Report Type:Expedited (15-DaCompany Report #001-0945-980830
 Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1800 MG (600 Initial or Prolonged MG, TID), PER	Delirium	Health	Neurontin	PS		ORAL
	Intentional Misuse	Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Neurontin SS

Date:11/23/98ISR Number: 3161609-9Report Type:Expedited (15-DaCompany Report #001-0945-980829
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anaphylactic Reaction	Consumer	Neurontin	PS		ORAL
900 MG (300 Required MG TID) PER Intervention to ORAL Prevent Permanent Impairment/Damage		Anuria Pruritus		Bactrim Atarax Prozac Reglan Xanax Prevacid	SS C C C C C		

Date:11/27/98ISR Number: 3163650-9Report Type:Expedited (15-DaCompany Report #9839160
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 50.00 MG Initial or Prolonged TOTAL:DAILY Required Intervention to ORAL Prevent Permanent ORAL Impairment/Damage		Alcohol Problem Intentional Misuse	Consumer	Zoloft Navane Alcohol Neurontin Buspar Cozaar	PS SS SS SS C C		ORAL ORAL ORAL ORAL

Date:11/30/98ISR Number: 3163941-1Report Type:Expedited (15-DaCompany Report #9838796
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 5.00 MG		Arterial Occlusive Disease	Consumer	Glucotrol	PS		ORAL
TOTAL: DAILY: Required ORAL		Cardiac Failure					
Intervention to ORAL		Congestive		Neurontin	SS		ORAL
Prevent Permanent Impairment/Damage		Diabetic Neuropathy		Glucophage	C		
		Emotional Disorder		Propulsid	C		
		Fluid Retention		Prilosec	C		
		Glycosylated Haemoglobin Increased		Zocor	C		
		Haematoma		Vitamins	C		
		Oedema Peripheral					
		Peripheral Coldness					
		Pseudomonas Infection					
		Skin Discolouration					
		Skin Ulcer					
		Staphylococcal Infection					
		Wound Infection					

Date:11/30/98ISR Number: 3164189-7Report Type:Expedited (15-DaCompany Report #001-0945-980839
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other X 1 DOSE	1 DAY	Feeling Abnormal	Consumer	Neurontin	PS		
		Suicidal Ideation	Other				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/98ISR Number: 3165284-9Report Type:Expedited (15-DaCompany Report #033-0945-980038

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abortion Missed	Foreign Health	Neurontin (Gabapentin)	PS		
2400 MG		Complications Of Maternal					
(,DAILY),		Exposure To Therapeutic	Professional				
PLACENTAL		Drugs					
		Congenital Anomaly		Tegretol	C		
		Hydrops Foetalis					
		Mediastinum Neoplasm					
		Neuroblastoma					
		Oedema					
		Teratoma					

Date:12/01/98ISR Number: 3165338-7Report Type:Expedited (15-DaCompany Report #001-0945-980840

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Psychiatric Symptom	Consumer	Neurontin	PS		

Date:12/02/98ISR Number: 3165631-8Report Type:Direct

Company Report #

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Complications Of Maternal		Phenobarbital	PS		ORAL
150 MG PO QD;		Exposure To Therapeutic					
SEE IMAGE FOR		Drugs					
THERAPY		Sedation		Klonopin	SS		
1 MG; SEE		Small For Dates Baby					
IMAGE FOR							
THERAPY							

Neurontin

SS

1300 MG; SEE

IMAGE FOR

THERAPY

Date:12/04/98ISR Number: 3166592-8Report Type:Expedited (15-DaCompany Report #001-0945-980859

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Deafness	Health	Neurontin	PS		
900 MG (,			Professional				
DAILY)							

Date:12/04/98ISR Number: 3167133-1Report Type:Expedited (15-DaCompany Report #001-0945-980749

Age:5 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Benign Intracranial Hypertension	Health	Neurontin (Gabapentin)	PS		ORAL
300 MG			Professional				
(DAILY), PER							
ORAL				Diamox	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/07/98ISR Number: 3168027-8Report Type:Expedited (15-DaCompany Report #001-0945-980649

Age:74 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acute Respiratory	Health	Neurontin	PS		ORAL
2400 MG (800 Hospitalization - MG, TID) PER Initial or Prolonged ORAL	1 YR	Distress Syndrome Lung Disorder Necrosis Pneumonia Respiratory Failure	Professional				

Date:12/07/98ISR Number: 3168258-7Report Type:Direct

Age:67 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour		Gabapentin	PS		ORAL
300 MG / QHS / PO		Coordination Abnormal Hypomania		Amlodipine Besylate Multi-Vitamins Lactulose Glipizide Metformin Carbamazepine Lorazepam	C C C C C C C		

Date:12/07/98ISR Number: 3168267-8Report Type:Direct

Age:56 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ecchymosis		Neurontin	PS		ORAL
Other 300 MG AM PO				Paxil Coumadin	C C		

Date:12/09/98ISR Number: 3168761-XReport Type:Direct
Age: Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 400 MG PO BID Initial or Prolonged Other	Diplopia		Neurontin	PS		ORAL

Date:12/09/98ISR Number: 3169039-0Report Type:Expedited (15-DaCompany Report #001-0945-980860
Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required 1200 MG (600 Intervention to MG, BID) PER Prevent Permanent ORAL Impairment/Damage	Anaphylactic Shock Bronchospasm Cough Dyspnoea Face Oedema	Health Professional	Neurontin Paxil	PS C		ORAL

Date:12/09/98ISR Number: 3169210-8Report Type:Expedited (15-DaCompany Report #001-0073-980589
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Drug Effect Decreased Liver Function Test

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Abnormal
Platelet Count Decreased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
4800 MG	(DAILY), PER	Health Professional	Dilantin (Phenytoin Sodium) Neurontin (Gabapentin)	PS SS		ORAL
ORAL			Unspecified Chemotherapy Phenobarb Neurontin	SS C C		

Date:12/10/98ISR Number: 3169886-5Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Flushing Swelling		Neutontin Aspirin Some Inhalers	PS C C		

Date:12/14/98ISR Number: 3170533-7Report Type:Expedited (15-DaCompany Report #981207-016014873
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 75 MG, BID, Required ORAL Intervention to 500 MG, QD, Prevent Permanent ORAL		Alanine Aminotransferase Increased Aspartate Aminotransferase Increased	Foreign Health Professional	Topamax (Topiramate) Tablets Paracetamol	PS SS		ORAL ORAL

Impairment/Damage 600 MG, QD, ORAL	Blood Bilirubin Increased Blood Creatinine Increased	Gabapentin Vigabatrin	SS SS	ORAL ORAL
2 UNKNOWN, BID, ORAL	Drug Toxicity Encephalopathy Gamma-Glutamyltransferase Increased Hepatitis Renal Failure Acute Sepsis			

Date:12/14/98ISR Number: 3170599-4Report Type:Expedited (15-DaCompany Report #046-0945-980003
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - SEE TEXT, PER Initial or Prolonged ORAL		Epilepsy Grand Mal Convulsion	Foreign Health Professional	Neurontin Renitec Seloken Trombyl Cipramil Lamictal	PS C C C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/15/98ISR Number: 3171087-1Report Type:Expedited (15-DaCompany Report #001-0945-980874
Age:55 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 600 MG (200 Initial or Prolonged MG TID) PER ORAL	Condition Aggravated Schizophrenia Stress	Consumer	Neurontin	PS		ORAL
			Vitamin Nos	C		
			Zantac	C		
			Multivitamins(Ergoca lciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Vitamin B-12	C C		

Date:12/16/98ISR Number: 3170868-8Report Type:Expedited (15-DaCompany Report #001-0945-980870
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Idiopathic Thrombocytopenic Purpura	Health Professional	Neurontin (Gabapentin)	PS		

Date:12/16/98ISR Number: 3231547-1Report Type:Periodic Company Report #8-98175-071A
Age:49 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 25MG DAILY AS NEEDED ORAL	Anaemia Cough	Health Professional	Duract	PS		ORAL
300MG DAILY ORAL			Neurontin (Gabapentin)	SS		ORAL
			Ergomar	C		

Imipramine	C
Klonopin	C
Lipitor	C
Neurontin	C
Premarin	C
Reglan	C
Vicodin	C
Vitamin E	C
Zoloft	C

Date:12/17/98ISR Number: 3171707-1Report Type:Expedited (15-DaCompany Report #001-0945-980608
 Age:22 MON Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Arthralgia	Consumer	Neurontin	PS		
2400 MG (,900		Complications Of Maternal					
MG QAM, 60 MG		Exposure To Therapeutic					
QNOON AND		Drugs					
QPM, 300 MG		Hair Growth Abnormal					
QHS), OTHER)		Hypertrophy Breast		Felbatol	SS		
400 MG		Migraine					
(,DAILY),		Precocious Puberty					
OTHER							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/17/98ISR Number: 3171710-1Report Type:Expedited (15-DaCompany Report #001-0945-980869
 Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Medication Error Sepsis	Health Professional	Neurontin	PS		

Date:12/17/98ISR Number: 3171727-7Report Type:Expedited (15-DaCompany Report #001-0073-980597
 Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG AM/200MG PM, PER ORAL 1200MG (600MG BID) PER ORAL		Idiopathic Thrombocytopenic Purpura	Health Professional	Dilantin Neurontin	PS SS		ORAL ORAL

Date:12/18/98ISR Number: 3171758-7Report Type:Expedited (15-DaCompany Report #049-0945-980025
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 2400 MG (,DAILY), PER ORAL		Renal Impairment	Foreign Health Professional Company Representative	Neurontin	PS		ORAL

Date:12/22/98ISR Number: 3173529-4Report Type:Direct Company Report #
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Acute Tonsillitis		Phenytoin	PS		
Initial or Prolonged		Angioneurotic Oedema		Gabapentin	SS		
		Dermatitis					
		Face Oedema					
		Liver Function Test					
		Abnormal					
		Rash Generalised					
		Rash Maculo-Papular					
		Stomatitis					
		Throat Irritation					
		Urinary Tract Infection					

Date:12/29/98ISR Number: 3174159-0Report Type:Direct Company Report #
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Tardive Dyskinesia		Gabapentin	PS		ORAL
300 MG PO TID							
Initial or Prolonged							
+100 MG HS 2							
Disability							
DAYS							
				Clozapine	C		
				Baclofen	C		
				I-Thyroxine	C		
				Ipratropium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/29/98ISR Number: 3176418-4Report Type:Expedited (15-DaCompany Report #001-0945-980839
 Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Abnormal	Health	Neurontin	PS		
X 1 DOSE		Mania Suicidal Ideation	Professional				

Date:12/30/98ISR Number: 3176821-2Report Type:Expedited (15-DaCompany Report #001-0945-980859
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Deafness	Health	Neurontin	PS		
900 MG			Professional				
(DAILY)							

Date:12/30/98ISR Number: 3176825-XReport Type:Expedited (15-DaCompany Report #049-0945-980025
 Age:73 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Arrhythmia	Foreign	Neurontin	PS		ORAL
3200 MG (800							
Hospitalization -		Blood Urea Decreased	Health				
MG, QID), PER							
Initial or Prolonged		Cardio-Respiratory Arrest	Professional				
ORAL							
		Creatinine Renal Clearance Decreased Dehydration Drug Toxicity Electrolyte Imbalance Pulmonary Embolism Renal Impairment Vomiting					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other Required 900 MG (300 Intervention to MG, TID) PER Prevent Permanent ORAL Impairment/Damage		Anaphylactoid Reaction Pruritus	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
UNKNOWN	UNKNOWN			Bactrim (Sulfamethoxazole, Trimethoprim)	SS		
				Atarax (Hydroxyzine Hydrochloride)	C		
				Prozac (Fluoxetine Hydrochloride)	C		
				Reglan (Metoclopramide)	C		
				Xanax (Alprazolam)	C		
				Prevacid (Lansoprazole)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1800 MG (600 Initial or Prolonged MG TID) PER		Delirium Intentional Misuse	Health Professional	Neurontin	PS		ORAL

Freedom Of Information (FOI) Report

ORAL

Date:01/06/99ISR Number: 3177775-5Report Type:Expedited (15-DaCompany Report #001-0945-980919
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG (QHS), Initial or Prolonged PER ORAL;		Blindness Dizziness Headache Histoplasmosis Paraesthesia Sedation Vision Blurred	Consumer	Neurontin Prozac	PS C		ORAL

Date:01/11/99ISR Number: 3179461-4Report Type:Expedited (15-DaCompany Report #111386
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly .5 MG 2 X PER DAY ORAL		Benign Congenital Hypotonia	Foreign Other	Rivotril	PS		ORAL
500 MG 3 X PER DAY ORAL		Breech Presentation Complications Of Maternal Exposure To Therapeutic		Depakine (Valproate Sodium) 500 Mg	SS		ORAL
200 MG 2 X PER DAY ORAL		Drugs Drug Withdrawal Syndrome Neonatal Ear Malformation Facial Dysmorphism Hypospadias Premature Baby Prepuce Redundant Small For Dates Baby Tremor Neonatal		Neurontin (Gabapentin) 100 Mg	SS		ORAL

Date:01/11/99ISR Number: 3179532-2Report Type:Expedited (15-DaCompany Report #LBID002990002

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 600 MG, PER Initial or Prolonged DAY, PER ORAL		Agitation	Consumer	Lithobid	PS		ORAL
PER ORAL		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: 3294263-6Report Type:Periodic Company Report #2972/20771

Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2 MG-2Q1DY;ORAL		Drug Effect Decreased Keratoconjunctivitis	Consumer Company	Detrol Tablets (2 Mg)	PS		ORAL
		Sicca	Representative	Tegretol	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Neurontin SS

Date:01/12/99ISR Number: 3179781-3Report Type:Expedited (15-DaCompany Report #001-0945-980860
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to 1200 MG (600 Prevent Permanent MG, BID), PER Impairment/Damage ORAL		Anaphylactic Shock Bronchospasm Cough Dermatitis Dyspnoea Face Oedema	Health Professional	Neurontin (Gabapentin) Paxil (Paroxetine Hydrochloride)	PS C		ORAL

Date:01/12/99ISR Number: 3179783-7Report Type:Expedited (15-DaCompany Report #001-0945-980923
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 3000 MG, DAILY 200 MG, DAILY 500 MG, DAILY		Abortion Spontaneous	Health Professional	Neurontin (Gabapentin) Lamictal (Lamotrigine) Mysoline (Primidone) Folic Acid	PS SS SS C		

Date:01/14/99ISR Number: 3180681-3Report Type:Expedited (15-DaCompany Report #033-0945-980049
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 6 CAPSULES (,		Rash Papular	Foreign	Neurontin	PS		ORAL

DAILY), PER

ORAL

Study

Health

Professional

Triatec (Ramipril)	C
Lexomil (Bromazepam)	C
Rivotril (Clonazepam)	C
Coversyl (Perindopril)	C
Esberiven Forte (Rutoside, Melilot)	C
Stagib (Metformin Embonate)	C
Glucophagea (Metformin Hydrochloride)	C

Date:01/14/99ISR Number: 3180699-0Report Type:Expedited (15-DaCompany Report #001-0073-980624

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health	Dilantin	PS		
600 MG (, DAILY)			Professional				
900 MG (, DAILY)				Neurontin	SS		
1500 MG (, DAILY)				Zarontin	SS		
				Prenatal Vitamins	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/99ISR Number: 3180838-1Report Type:Expedited (15-DaCompany Report #033-0945-990001

Age:1 DY Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly SEE TEXT, Other PLACENTAL; IN UTERO EXPOSURE; PLACENTAL SEE TEXT, PLACENTAL; IN UTERO EXPOSURE; PLACENTAL SEE TEXT, PLACENTAL; IN UTERO EXPOSURE; PLACENTAL	Duration Complications Of Maternal Exposure To Therapeutic Drugs Facial Dysmorphism Feeling Jittery Hypospadias Hypotonia Premature Baby Skin Hypertrophy	Foreign Health Professional	Neurontin Rivotril Depakine	PS SS SS		

Date:01/19/99ISR Number: 3295967-1Report Type:Periodic Company Report #WAES 97050892

Age:30 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose PO	Duration Crystalluria Hyperbilirubinaemia Renal Failure Acute	Health Professional	Cap Crixivan Unk Neurontin Unk Epivir	PS SS C		ORAL

Date:01/20/99ISR Number: 3182805-0Report Type:Expedited (15-DaCompany Report #001-0945-980923
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health Professional	Neurontin (Gabapentin)	PS		
3000 MG DAILY				Lamictal (Lamotrigine)	SS		
200 MG DAILY				Mysoline (Primidone)	SS		
500 MG DAILY				Folic Acid	C		

Date:01/25/99ISR Number: 3187823-4Report Type:Expedited (15-DaCompany Report #SP-9900050
 Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Cellulitis	Health Professional	Remicade	PS		
INTRA VENOUS	INTRA VENOUS 5						
Intervention to		Convulsion					
MG/KG							
Prevent Permanent Impairment/Damage				Gabapentin	SS		
				Doxepin	SS		
				Oxycodone	SS		
				Azathioprine	C		
				Omeprazole	C		
				Prednisone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/26/99ISR Number: 3185295-7Report Type:Direct
 Age: Gender: I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Liver Function Test		Lisinopril	PS		
Initial or Prolonged	Abnormal		Diflucan	SS		
			Gabapentin	SS		
			Macrochantin	SS		

Date:01/26/99ISR Number: 3186035-8Report Type:Expedited (15-DaCompany Report #001-0945-990025
 Age:1 DY Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Complications Of Maternal Exposure To Therapeutic	Health Professional	Neurontin (Gabapentin)	PS		
TRANSPLACENTAL	SEE TEXT,					
PLACENTAL IN	Drugs					
UTERO	Convulsion					
EXPOSURE			Prenatal Vitamins	C		
			Folic Acid	C		

Date:01/26/99ISR Number: 3197622-5Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #9837475

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Drug Ineffective	Health	Viagra Tablets	PS		ORAL
50.00 MG	Drug Interaction	Professional				
TOTAL:PRN:ORA						
L			Neurontin	SS		ORAL
ORAL						

Date:01/29/99ISR Number: 3188590-0Report Type:Expedited (15-DaCompany Report #033-0945-980049
Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Foreign	Neurontin	PS		ORAL
6 CAPSULE		Eczema	Study				
(DAILY) PER		Toxic Skin Eruption	Health				
ORAL			Professional	Triatec	C		
				Lexomil	C		
				Rivotril	C		
				Coversyl	C		
				Esberiven	C		
				Stagid	C		
				Glucophage	C		

Date:01/29/99ISR Number: 3188593-6Report Type:Expedited (15-DaCompany Report #001-0945-990035
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Asthma	Health	Neurontin	PS		ORAL
2100 MG							
Initial or Prolonged		Bronchitis	Professional				
(DAILY) PER		Dermatitis					
ORAL; 2400 MG		Herpes Zoster					
SEE IMAGE		Pain		Unspecified			
		Pyrexia		Antihypertensive	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/29/99ISR Number: 3188595-XReport Type:Expedited (15-DaCompany Report #001-0945-980783

Age:51 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 400 MG (100 Initial or Prolonged MG, QID) PER ORAL	Diarrhoea Dizziness Orthostatic Hypotension Renal Disorder	Consumer	Neurontin	PS		ORAL
			Cozaar	C		
			Insulin	C		
			Nph Insulin	C		
			Proamatine	C		
			Lasix	C		

Date:02/01/99ISR Number: 3189846-8Report Type:Expedited (15-DaCompany Report #001-0945-990043

Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 2400 MG, Required (DAILY), Intervention to Prevent Permanent Impairment/Damage	Anaphylactic Shock Hemiplegia	Health Professional	Neurontin	PS		
			Lipitor	SS		
			Naprosyn	C		

Date:02/01/99ISR Number: 3301104-7Report Type:Periodic Company Report #A0067798

Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose 25 MG TWICE PER DAY ORAL	Asthenia Dizziness Fatigue Hyperhidrosis Sedation	Consumer	Lamictal Tablet	PS		ORAL
			Gabapentin (Formulation Unknown) Sertraline	SS		

Hydrochloride C
Doxazosin Mesylate C
Moexipril
Hydrochloride C
Lisinopril C
Potassium Chloride C
Frusemide C

Date:02/04/99ISR Number: 3192107-4Report Type:Expedited (15-DaCompany Report #002-0945-990003
Age:69 YR Gender:Male I/FU:I

Outcome PT
Hospitalization - Blood Creatine
Initial or Prolonged Phosphokinase Increased
Blood Creatine
Phosphokinase Mb
Increased
Body Temperature
Increased
Chest Pain
Confusional State
Cough
Diarrhoea
Musculoskeletal Stiffness
Neutrophil Count
Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Paralysis Pleural Effusion Sedation	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		White Blood Cell Count Increased	Foreign Health Professional	Gabapentin	PS		ORAL
20 MG (DAILY) PER ORAL				Lovastatin	SS		ORAL
				Atenolol	C		
				Nitroglycerin	C		
				Norvasc	C		
				Rocaltrol	C		
				Calcium Carbonate	C		
				Losec	C		
				Nitroglycerin	C		
				Restoril	C		
				Dulcolax	C		
				Extra Strength			
				Tylenol	C		
				Lactulose	C		
				Colace	C		
				Atasol	C		
				Eprex	C		
				Penta	C		

Date:02/05/99ISR Number: 3199732-5Report Type:Periodic Company Report #FLUV002980264

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 50 MG, PER ORAL		Dermatitis	Health Professional	Luvox	PS		ORAL
PER ORAL		Drug Interaction		Neurontin (Gabapentin)	SS		ORAL

Date:02/09/99ISR Number: 3194448-3Report Type:Expedited (15-DaCompany Report #034-0945-990002
Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300 MG DAILY	Coordination Abnormal	Foreign	Gabapentin	PS		ORAL
Initial or Prolonged PER ORAL	Muscle Twitching	Health				
50 MG DAILY PER ORAL	Tremor	Professional	(Amitriptyline Hydrochloride)	SS		ORAL
			Doxazosin Mesilate	C		
			Lacidipine	C		
			Calcium Carbonate	C		
			Insulin Human	C		
			Acetylsalicylic Acid	C		

Date:02/09/99ISR Number: 3194450-1Report Type:Expedited (15-DaCompany Report #001-0945-980919
Age:34 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Blindness Unilateral Dizziness Headache Histoplasmosis Paraesthesia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sedation Vision Blurred		Report Source	Product	Role	Manufacturer	Route	
Dose	Duration	Consumer	Neurontin	PS		ORAL	
100 MG(QHS)							
PER ORAL			Prozac	C			
Date:02/09/99ISR Number: 3194989-9Report Type:Direct		Company Report #					
Age:	Gender:Female	I/FU:I					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations	Health	Arthrotec	PS		ORAL
75 MG BID PO							
		Ventricular Extrasystoles	Professional	Neurontin	SS		ORAL
600 MG TID PO	3 MON			Prozac	C		
Date:02/11/99ISR Number: 3195858-0Report Type:Expedited (15-Da		Company Report #049-0945-980025					
Age:73 YR	Gender:Male	I/FU:F					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arrhythmia	Foreign	Neurontin	PS		ORAL
Death							
3200 MG (800		Blood Creatinine	Health				
Hospitalization -							
MG, QID), PER		Increased	Professional				
Initial or Prolonged							
ORAL		Blood Urea Increased	Company	Tegretol	C		
		Cardio-Respiratory Arrest	Representative				
		Dehydration					
		Drug Toxicity					
		Electrolyte Imbalance					
		Pulmonary Embolism					
		Renal Impairment					
		Vomiting					

Date:02/17/99ISR Number: 3200356-1Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Palpitations		Arthrotec	PS		ORAL
75MG BID PO							
		Ventricular Extrasystoles		Neurontin	SS		ORAL
600MG TID PO				Prozac	C		

Date:02/19/99ISR Number: 3204464-0Report Type:Expedited (15-DaCompany Report #199910754DDC
Age:84 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - Initial or Prolonged RESPIRATORY		Atrioventricular Block Status Epilepticus Thrombocytopenia	Foreign	Terbutaline Sulfate (Bricanyl) Aerosol (Solution)	PS		
(INHALATION)	INH	1 WK					
				Methylprednisolone	SS		
				Gabapentin	SS		
				Heparin-Fraction, Sodium Salt	SS		
30 DAY				Phenytoin	SS		
1 DAY				Amoxicillin Trihydrate	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/22/99ISR Number: 3203557-1Report Type:Direct
Age:44 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oedema	Health	Gabapentin	PS		ORAL
300 MG	TID						
		Weight Increased	Professional	Neurontin	SS		ORAL
				Humalog	C		
				Humulin Ultralente	C		
				Asa	C		
				Atenolol	C		
				Captopril	C		
				Propulsid	C		
				Prilosec	C		
				Premarin	C		
				Paxil	C		
				Klonopin	C		
				Valium	C		
				Duragesic	C		
				Oxycodone	C		
				Alupent	C		
				Allegra	C		
				Ntg	C		

Date:02/22/99ISR Number: 3204891-1Report Type:Expedited (15-DaCompany Report #001-0945-990101
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Medication Error	Health	Neurontin	PS		ORAL
1200 MG (X1)							
			Professional				
PER ORAL							

Date:02/22/99ISR Number: 3205029-7Report Type:Expedited (15-DaCompany Report #1274/11153
Age:84 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Thrombocytopenia	Foreign	Medrol	PS		ORAL
16 MG-3Q1DY;							

Other		Consumer			
ORAL		Company Representative	Amoxicillin & Clavulanic Acid	SS	ORAL
2 GM/DAY;ORAL			Phenytoin	SS	
INTRAVENOUS	IV		Terbutaline	SS	NASAL
NASAL			Gabapentine	SS	ORAL
400					
MG-3D1DY;ORAL			Tinzaparine Sodium	SS	
SUBCUTANEOUS	2500				
IU-1Q1DY;SC					

Date:02/23/99ISR Number: 3311544-8Report Type:Periodic Company Report #1998-002917
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Flomax (Tamsulosin)			
0.4 MG/PO		Dysarthria		Capsules / 0.4 Mg	PS		ORAL
1 MG/PO		Gait Disturbance		Ativan	SS		ORAL
100 MG/PO		Paraesthesia		Elavil	SS		ORAL
PO		Sedation		Neurontin	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/25/99ISR Number: 3208681-5Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #9838101

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ecchymosis	Health	Zoloft Tablets	PS		ORAL
100.00 MG			Professional				
TOTAL: DAILY:							
ORAL							
				Gabapentin	SS		
600.00 MG							
TOTAL: BID							

Date:02/25/99ISR Number: 3209965-7Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #9828199

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea	Health	Zoloft Tablets	PS		ORAL
ORAL		Drug Interaction	Professional	Neurontin	SS		
				Ambien	C		
				Hydrocodone /			
				Acetaminophen	C		
				Unknown Rstrogen	C		

Date:02/26/99ISR Number: 3208421-XReport Type:Expedited (15-DaCompany Report #001-0945-990043
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Anaphylactic Reaction	Health	Neurontin			
Required		Condition Aggravated	Professional	(Gabapentin)	PS		
2400 MG							
Intervention to							
(,DAILY),							
Prevent Permanent							
UNKNOWN							
Impairment/Damage				Lipitor			
				(Atorvastatin)	SS		

Date:02/26/99ISR Number: 3208598-6Report Type:Expedited (15-DaCompany Report #044-0945-990016
 Age:1 DY Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Cleft Lip Cleft Palate	Foreign Health	Neurontin (Gabapentin)	PS		
SEE TEXT, PLACENTAL		Complications Of Maternal	Professional				
SEE TEXT, PLACENTAL		Exposure To Therapeutic Drugs		Phenytoin	SS		
		Congenital Megacolon Intestinal Obstruction					

Date:02/26/99ISR Number: 3211129-8Report Type:Periodic Company Report #001-0945-980362
 Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Atrial Fibrillation	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG, TID) PER ORAL				Lasix	C		
				Norvasc	C		
				Iron	C		
				Hytrin	C		
				Pravachol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3211133-XReport Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-980381

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Encephalopathy	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3211135-3Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980413

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Decreased Activity Fatigue	Health Professional	Neurontin (Gabapentin)	PS		
2700 MG (900 MG, TID),		Hypotension		Sertraline	C		
		Paraesthesia		Klonopin	C		
				Lamictal	C		

Date:02/26/99ISR Number: 3211139-0Report Type:Periodic
Age:27 YR Gender:Male I/FU:I

Company Report #001-0945-980441

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3600 MG (1200 MG, TID) PER ORAL		Depression Hostility	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
		Injury					
		Suicide Attempt					
				Zantac	C		

Date:02/26/99ISR Number: 3211142-0Report Type:Periodic
Age:41 YR Gender:Male I/FU:I

Company Report #001-0945-980460

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Deep Vein Thrombosis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Initial or Prolonged							
900 MG (300							
MG, TID) PER							
ORAL							

Naproxen C

Date:02/26/99ISR Number: 3211145-6Report Type:Periodic Company Report #001-0945-980477
 Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Haemorrhage Vitreous Floaters	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG							
(DAILY) PER							
ORAL							

Toprol C
 Casodex C
 Zocor C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3211149-3Report Type:Periodic
Age:63 YR Gender:Female I/FU:I

Company Report #001-0945-980511

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 MG, TID) PER ORAL		Pneumonia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Premarin	C		
				Provera	C		
				Lasix	C		
				Monopril	C		
				Prilosec	C		
				Zoloft	C		
				Immunosuppressive Therapy	C		
				Cyclosporin A	C		
				Prednisone	C		
				Estrace	C		

Date:02/26/99ISR Number: 3211153-5Report Type:Periodic
Age:34 YR Gender:Male I/FU:I

Company Report #001-0945-980523

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2400 MG (800 MG, TID)		Coma Dysphagia Hypoaesthesia	Health Professional	Neurontin (Gabapentin)	PS		
		Reflexes Abnormal Vision Blurred		Unspecified Opiates Methadone	SS C		

Date:02/26/99ISR Number: 3211157-2Report Type:Periodic
Age:66 YR Gender:Female I/FU:I

Company Report #001-0945-980536

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Medication Error	Health	Neurontin Capsules			

Initial or Prolonged Thrombocytopenia Professional 300 Mg (Gabapentin) PS ORAL
 900 MG (300
 MG, TID) PER
 ORAL
 Daypro C

Date:02/26/99ISR Number: 3211161-4Report Type:Periodic Company Report #001-0945-980556
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG DAILY)	2 DAY	Confusional State Medication Error Pyrexia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
PER ORAL				Digoxin	C		
				Accupril	C		
				Tylenol	C		
				Coreg	C		
				Thyroid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3211164-XReport Type:Periodic
Age:52 YR Gender:Female I/FU:I

Company Report #001-0945-980579

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (DAILY) PER ORAL	Facial Palsy Medication Error	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Tegretol Sinequan	C C		

Date:02/26/99ISR Number: 3211167-5Report Type:Periodic
Age:54 YR Gender:Female I/FU:I

Company Report #001-0945-980582

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 MG, TID)	Atrioventricular Block Confusional State Depressed Level Of Consciousness	Health Professional	Neurontin (Gabapentin)	PS		
			Many Unspecified Medications	C		

Date:02/26/99ISR Number: 3211171-7Report Type:Periodic
Age:59 YR Gender:Female I/FU:I

Company Report #001-0945-980591

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2700 MG DAILY PER ORAL	Oedema Peripheral	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Mexiletine Ms Contin Synthroid Insulin	C C C C		

Date:02/26/99ISR Number: 3211177-8Report Type:Periodic
Age:24 YR Gender:Female I/FU:I

Company Report #001-0945-980600

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Cognitive Disorder	Health	Neurontin Capsules			
Initial or Prolonged	Overdose	Professional	400 Mg (Gabapentin)	PS		
			Tylenol (Paracetamol)	SS		

Date:02/26/99ISR Number: 3211181-XReport Type:Periodic
Age:40 YR Gender:Female I/FU:I

Company Report #001-0945-980607

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Cerebrovascular Accident	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG						
(DAILY) PER						
ORAL	2 DAY		Soma	C		
			Morphine Sulfate	C		
			Xanax	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3211184-5Report Type:Periodic
Age:54 YR Gender:Female I/FU:I

Company Report #001-0945-980611

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, TID) PER ORAL		Henoch-Schonlein Purpura Renal Failure Acute	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3211188-2Report Type:Periodic
Age:54 YR Gender:Female I/FU:I

Company Report #001-0945-980615

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (300 MG, TID),		Medication Error Platelet Count Decreased	Health Professional	Neurontin (Gabapentin)	PS		
				Prednisone	C		
				Serzone	C		
				Wellbutrin	C		
				Folic Acid	C		
				Desyrel	C		
				Vicodin	C		
				Syn-Throin	C		

Date:02/26/99ISR Number: 3211193-6Report Type:Periodic
Age:32 YR Gender:Female I/FU:I

Company Report #001-0945-980663

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3600 MG (1200 MG, TID) PER ORAL		Mental Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Multiple Unspecified Medications C

Date:02/26/99ISR Number: 3211197-3Report Type:Periodic Company Report #001-0945-980673
Age:41 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Accidental Overdose	Health	Neurontin			
Initial or Prolonged	Mania	Professional	(Gabapentin) Zyprexa	PS C		

Date:02/26/99ISR Number: 3211201-2Report Type:Periodic Company Report #001-0945-980685
Age:16 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Stevens-Johnson Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL

1200 MG (300

MG, QID) PER

ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3211288-7Report Type:Periodic
Age:26 YR Gender:Female I/FU:I

Company Report #001-0945-980699

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Thrombocytopenia	Health Professional	Neurontin (Gabapentin) Phenobarb	PS C		

Date:02/26/99ISR Number: 3211292-9Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-980702

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2700, DAILY,	Aggression Confusional State	Health Professional	Neurontin (Gabapentin)	PS		

UNKNOWN

Date:02/26/99ISR Number: 3211295-4Report Type:Periodic
Age:28 YR Gender:Female I/FU:I

Company Report #001-0945-980710

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (300	Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL

MG, TID), PER

ORAL

250 MG (TOOK

ONE DOSE),

PER ORAL

Depakote (Valproate Semisodium)	SS	ORAL
------------------------------------	----	------

Buspar	C	
Zoloft	C	

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anger	Consumer	Neurontin			
900 MG (300		Anxiety		(Gabapentin)	PS		ORAL
MG, TID), PER		Arthropathy					
ORAL		Asthma					
		Balance Disorder		Cardizem	C		
		Chest Discomfort		Atenolol	C		
		Headache		Estrogen Nos	C		
		Heart Rate Increased		Flexeril	C		
		Muscle Spasms					
		Pain In Extremity					
		Pyrexia					
		Suicidal Ideation					
		Thinking Abnormal					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Thyroid Stimulating	Health	Neurontin			
Initial or Prolonged		Hormone Decreased	Professional	(Gabapentin)	PS		
		Convulsion		Clonazepam	C		
		Depression		Zoloft	C		
		Lethargy					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3211308-XReport Type:Periodic
 Age:49 YR Gender:Female I/FU:I

Company Report #001-0945-989772

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (300 MG, QID) PER ORAL	Oedema	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
			Lorazepam	C		
			Remeron	C		
			Vicodin	C		
			Synthroid	C		
			Soma	C		

Date:02/26/99ISR Number: 3211311-XReport Type:Periodic
 Age:58 YR Gender:Female I/FU:I

Company Report #001-0945-980831

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG (600 MG, TID), PER ORAL	Pancreatitis	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
			Vicodin	C		
			Prilosec	C		

Date:02/26/99ISR Number: 3211315-7Report Type:Periodic
 Age:17 YR Gender:Male I/FU:I

Company Report #001-0945-980861

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged SEE TEXT, UNKNOWN	Bipolar Disorder Medication Error	Consumer	Neurontin (Gabapentin)	PS		

Wellbutrin C
Trazodone C
Depakote C

Date:02/26/99ISR Number: 3211320-0Report Type:Periodic Company Report #001-0945-980888
Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG, DAILY, PER ORAL	Pancreatitis	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

Divalproex C

Date:02/26/99ISR Number: 3211323-6Report Type:Periodic Company Report #001-0945-980906
Age:24 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged PER ORAL 300 MG (150 MG, BID), PER ORAL	Medication Error Stevens-Johnson Syndrome	Health Professional	Neurontin (Gabapentin) Zyban (Amfebutamone)	PS SS		ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3211328-5Report Type:Periodic
Age:27 YR Gender:Female I/FU:I

Company Report #001-0945-980921

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Breast Engorgement Chills	Consumer	Neurontin (Gabapentin)	PS		
1600 MG (600 MG BID, 400 MG		Dizziness Galactorrhoea					
		Headache Insomnia Medication Error Suicidal Ideation Vomiting		Prozac Valium Xanax	C C C		

Date:02/26/99ISR Number: 3211332-7Report Type:Periodic
Age:35 YR Gender:Male I/FU:I

Company Report #001-0945-990003

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 MG, BID), PER ORAL		Epistaxis Haemoglobin Decreased Headache Photophobia Thrombocytopenia	Health Professional Company Representative	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Aciclovir Prilosec Tylenol Imitrex	C C C C		

Date:02/26/99ISR Number: 3211336-4Report Type:Periodic
Age:23 YR Gender:Female I/FU:F

Company Report #001-0945-973006

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (400		Complications Of Maternal Exposure To Therapeutic	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL

MG TID) PER	Drugs					
ORAL	Porphyria					
	Premature Labour			Desyrel	C	
				Neprogan	C	
				Toprol	C	
				Compazine	C	
Date:02/26/99ISR Number: 3211357-1Report Type:Periodic Company Report #001-0945-973033						
Age:79 YR Gender:Female I/FU:F						
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Activated Partial	Health	Neurontin			
Initial or Prolonged	Thromboplastin Time	Professional	(Gabapentin)	PS		ORAL
900 MG (300	Prolonged					
MG, TID) PER	International Normalised					
ORAL	Ratio Increased		Coumadin	C		
	Phlebitis		Synthroid	C		
	Prothrombin Time		Bumex	C		
	Prolonged		Allopurinol	C		
			Digoxin	C		
			Maxzide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3212580-2Report Type:Periodic
Age:34 YR Gender:Female I/FU:I

Company Report #001-0945-980046

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 900 MG (300 Initial or Prolonged MG, TID), PER		Coordination Abnormal Depression Diplopia Disorientation Drug Level Above Therapeutic Drug Toxicity Dysarthria Fatigue Gait Disturbance Medication Error Stress	Consumer	Neurontin	PS		ORAL
ORAL; 1800 MG, 2400 MG (600 MG,				Cough Medicine Desyrel Klonopin (Clonazepam) Cipro Premarin Luvox Depakote Imitrex Paxil	C C C C C C C C C		

Date:02/26/99ISR Number: 3212589-9Report Type:Periodic
Age:26 YR Gender:Female I/FU:I

Company Report #001-0945-980052

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Deafness	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 MG (600 MG, BID), PER				Depakote (Valproate Semisodium)	C		
ORAL							

Date:02/26/99ISR Number: 3212597-8Report Type:Periodic
Age:23 YR Gender:Female I/FU:I

Company Report #001-0945-980087

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1800 MG 900 Initial or Prolonged MG, BID), PER		Thrombocytopenia	Health Professional	Neurontin	PS		ORAL
ORAL				Sabril (Vigabatrin)	C		
				Diamox (Acetazolamide)	C		
				Contraceptive Pill (Oral Contraceptive Nos)	C		

Date:02/26/99ISR Number: 3212604-2Report Type:Periodic Company Report #001-0945-980115
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2000 MG, Initial or Prolonged DAILY, PER		Anxiety Coordination Abnormal Dysarthria	Health Professional	Neurontin	PS		ORAL
ORAL		Liver Function Test Abnormal Medication Error Orthostatic Hypotension		Accupril (Quinapril Hydrochloride)	C		
				Pravachol (Pravastatin Sodium)	C		
				Fioricet (Caffeine, Butalbital, Paracetamol)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Centrum (Vitamins
Nos, Minerals Nos) C

Date:02/26/99ISR Number: 3212613-3Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-980121

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Renal Failure	Health Professional Company Representative	Neurontin	PS		

Date:02/26/99ISR Number: 3212619-4Report Type:Periodic
Age:50 YR Gender:Male I/FU:I

Company Report #001-0945-980136

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Photophobia	Health	Neurontin	PS		ORAL
2400 MG (800		Vision Blurred	Professional				
MG, TID), PER		Visual Acuity Reduced					
ORAL							

Tegretol
(Carbamazepine) C
Insulin C
Paxil (Paroxetine
Hydrochloride) C
Glucophage
(Metformin
Hydrochloride) C
Toprol Xl
(Metoprolol
Succinate) C
Lasix (Furosemide) C
Zocor (Simvastatin) C
Beconase
(Beclometasone
Dipropionate) C
Asa (Acetylsalicylic
Acid) C

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - PER ORAL	Amnesia	Consumer	Neurontin	PS		ORAL
Initial or Prolonged	Malaise	Health	Vanceril			
	Status Epilepticus	Professional	(Beclometasone)	C		
			Proventil			
			(Salbutamol)	C		
			Beconase			
			(Beclometasone			
			Dipropionate)	C		
			Tegretol			
			(Carbamazepine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3212630-3Report Type:Periodic
Age:43 YR Gender:Male I/FU:I

Company Report #001-0945-980151

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 2700 MG (900 Initial or Prolonged MG, TID), PER	Convulsion	Health	Neurontin	PS		ORAL
ORAL	Diabetes Mellitus	Professional	Depakote	C		

Date:02/26/99ISR Number: 3212658-3Report Type:Periodic
Age:16 YR Gender:Female I/FU:I

Company Report #001-0945-980161

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 4800 MG (1100 MG TID, 1500 MG AT	Clonic Convulsion Drug Level Below Therapeutic Overdose	Health Professional	Neurontin (Gabapentin)	PS		
			Tegretol (Carbamazepine)	C		
			Mysoline	C		
			Tranxene	C		
			Synthroid	C		

Date:02/26/99ISR Number: 3212676-5Report Type:Periodic
Age:70 YR Gender:Male I/FU:I

Company Report #001-0945-980164

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 600 MG (300 Initial or Prolonged MG, BID), PER	Abdominal Pain Dyspnoea	Consumer	Neurontin	PS		ORAL
ORAL			Lipitor			

(Atorvastatin) C
 Lopressor
 (Metoprolol
 Tartrate) C
 Vitamins C

Date:02/26/99ISR Number: 3212678-9Report Type:Periodic Company Report #001-0945-980184
 Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 300 MG, DAILY, PER ORAL		Amblyopia Visual Acuity Reduced	Health Professional	Neurontin	PS		ORAL

Glucotrol
 (Glipizide) C
 Glucophage
 (Metformin
 Hydrochloride) C
 Rezulin
 (Troglitazone) C
 Spironolactone C
 Precose (Acarbose) C
 Mevacor (Lovastatin) C
 Norvasc (Amlodipine
 Besilate) C
 Daypro (Oxaprozin) C
 Humulin L (Insulin
 Human Zinc

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Suspension) C

Date:02/26/99ISR Number: 3212684-4Report Type:Periodic
Age:42 YR Gender:Male I/FU:I

Company Report #001-0945-980189

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness	Health	Neurontin			
		Medication Error	Professional	(Gabapentin)	PS		
1200 MG, AT							
BEDTIME							

Lithobid (Lithium Carbonate) C
Tegretol (Carbamazepine) C
Benadryl (Diphenhydramine Hydrochloride) C

Date:02/26/99ISR Number: 3212688-1Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #001-0945-980206

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract	Health	Neurontin	PS		
1800 MG,		Scotoma	Professional				
DAILY							

Synthroid C

Date:02/26/99ISR Number: 3212694-7Report Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980219

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression	Consumer	Neurontin	PS		ORAL
1800 MG (600		Headache					
MG, TID) PER		Oedema Peripheral					
ORAL							

Pulmonary Oedema

Lithium	C
Premarin (Estrogens Conjugated)	C
Prempro (Medroxyprogesterone)	C
Synthroid (Levothyroxine Sodium)	C
Hytrin (Terazosin Hydrochloride)	C
Klonopin (Clonazepam)	C
Cardizem (Diltiazem Hydrochloride)	C
Trazodone	C

Date:02/26/99ISR Number: 3212699-6Report Type:Periodic
Age:55 YR Gender:Male I/FU:I

Company Report #001-0945-980277

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 900 MG (300 Initial or Prolonged MG, TID)	Duration Abdominal Pain Hyperhidrosis 3 MON Medication Error	Health Professional	Neurontin Oxycontin Prozac (Fluoxetine Hydrochloride)	PS C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3212704-7Report Type:Periodic
Age:46 YR Gender:Male I/FU:I

Company Report #001-0945-980299

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 800 MG (400 Initial or Prolonged MG, BID), PER ORAL		Atrial Fibrillation	Health Professional	Neurontin	PS		ORAL

Risperdal
(Risperidone) C

Date:02/26/99ISR Number: 3212709-6Report Type:Periodic
Age:13 YR Gender:Male I/FU:I

Company Report #001-0945-980307

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2400 MG, DAILY, PER ORAL		Pancreatitis	Health Professional	Neurontin	PS		ORAL

Unspecified
Medications C

Date:02/26/99ISR Number: 3212713-8Report Type:Periodic
Age:24 YR Gender:Female I/FU:I

Company Report #001-0945-980327

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2700 MG, Initial or Prolonged DAILY		Medication Error Pneumonia	Health Professional	Neurontin	PS		
				Amitriptyline	C		
				Ultram	C		
				Oral Contraceptive Nos	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation	Consumer	Neurontin	PS		
1500 MG,		Convulsion					
DAILY		Feeling Abnormal		Tegretol	C		
		Headache					
		Insomnia					
		Sedation					
		Thinking Abnormal					
		Visual Disturbance					
		Visual Field Defect					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error	Consumer	Neurontin	PS		ORAL
50 MG, DAILY,		Pancreatitis					
PER ORAL				Vitamins	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3408310-1Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0073-980355

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Interaction Drug Level Changed	Health Professional	Dilantin Suspension 125 Mg/5 Ml (Phenytoin Sodium) Lamotrigine Neurontin(Gabapentin) Carbamazepine Declomycin(Demeclocy cline Hydrochloride) Risperdal(Risperidon e) Ativan (Lorazepam) Jevity (Potassium Bicarbonate, Potassium Bitartrate, Soya Oil, Corn Oil, Vagal Nerve Stimulator	PS SS SS SS SS SS SS SS		
PER TUBE							

Date:02/26/99ISR Number: 3410096-1Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #001-0945-980670

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1500 MG (300 MG, FIVE TIMES DAILY) PER ORAL		Anxiety Decreased Appetite Disorientation Insomnia Weight Decreased	Consumer	Neurontin Capsules 300 Mg (Gabapenitn)	PS		ORAL

Date:02/26/99ISR Number: 3410097-3Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-980674

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Weight Increased	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3410098-5Report Type:Periodic Company Report #001-0945-980675
 Age:45 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Skin Lesion	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1200 MG (300 MG, QID), PER ORAL				Lamictal (Lamotrigine)	C		

Date:02/26/99ISR Number: 3410099-7Report Type:Periodic Company Report #001-0945-980678
 Age:38 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Galactorrhoea	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG (, AT							

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Freedom Of Information (FOI) Report

BEDTIME), PER

ORAL

Depakote (Valproate
Semisodium) C

Date:02/26/99ISR Number: 3410100-0Report Type:Periodic
Age:46 YR Gender:Female I/FU:I

Company Report #001-0945-980679

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Chest Pain		Neurontin Capsules			
		Depressed Level Of		100 Mg (Gabapentin)	PS		ORAL
50 MG (,		Consciousness					
DAILY), PER		Heart Rate Decreased					

ORAL

Date:02/26/99ISR Number: 3410101-2Report Type:Periodic
Age:56 YR Gender:Female I/FU:I

Company Report #001-0945-980681

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety	Consumer	Neurontin			
		Dizziness		(Gabapentin)	PS		ORAL
600 MG (300							
MG, BID), PER							

ORAL

Nardil (Phenelzine
Sulfate) C
Premarin (Estrogens
Conjugated) C
Daypro (Oxaprozin) C
Axid (Nizatidine) C
Lorazepam C
(Quinidine Sulfate) C
Levo-T
(Levothyroxine) C
Klonopin
(Clonazepam) C

Date:02/26/99ISR Number: 3410102-4Report Type:Periodic
Age:46 YR Gender:Male I/FU:I

Company Report #001-0945-980682

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Pruritus	Consumer	Neurontin (Gabapentin)	PS		ORAL
1500 MG (,							
DAILY), PER							
ORAL							

Prozac (Fluoxetine
Hydrochloride) C
Ativan (Lorazepam) C
Zyprexa (Olanzapine) C

Date:02/26/99ISR Number: 3410103-6Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980686

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Increased	Consumer Health Professional	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410104-8Report Type:Periodic
Age:51 YR Gender:Female I/FU:I

Company Report #001-0945-980687

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL				(Baclofen)	C		
				Daypro (Oxaprozin)	C		
				Bellergal -S (Phenobarbital, Ergotamine Tartrate, Belladonna Alkaloids Ambien (Zolpidem Tartrate)	C C C		
				Estrogen	C		

Date:02/26/99ISR Number: 3410105-XReport Type:Periodic
Age:40 YR Gender:Male I/FU:I

Company Report #001-0945-980688

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Ill-Defined Disorder Overdose	Consumer	Neurontin (Gabapentin)	PS		ORAL
6000 MG (, DAILY), PER ORAL		Suicidal Ideation					

Date:02/26/99ISR Number: 3410106-1Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980695

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia	Health Professional Company	Neurontin (Gabapentin) Haldol (Haloperidol)	PS C		

Representative

Date:02/26/99ISR Number: 3410107-3Report Type:Periodic Company Report #001-0945-980698
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea	Consumer	Neurontin (Gabapentin)	PS		
(, QHS)							

Date:02/26/99ISR Number: 3410108-5Report Type:Periodic Company Report #001-0945-980701
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Health Professional	Neurontin (Gabapentin)	PS		ORAL
100 MG (, HS), PER ORAL 2 DAY							

Oxycontin (Oxycodone
Hydrochloride) C
Thyroid Supplement C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410109-7Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980703

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eye Discharge	Health	Neurontin			
		Visual Disturbance	Professional	(Gabapentin)	PS		
1800 MG (,		Visual Field Defect					
DAILY)							

Date:02/26/99ISR Number: 3410110-3Report Type:Periodic
 Age:35 YR Gender:Female I/FU:I

Company Report #001-0945-980708

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema	Health	Neurontin			
			Professional	(Gabapentin)	PS		ORAL
300 MG (100							
MG, TID), PER							
ORAL							

Lortab (Paracetamol,
 Hydrocone
 Bitartrate) C

Date:02/26/99ISR Number: 3410111-5Report Type:Periodic
 Age:46 YR Gender:Male I/FU:I

Company Report #001-0945-980711

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Engorgement	Health	Neurontin			
		Breast Pain	Professional	(Gabapentin)	PS		ORAL
900 MG (300							
MG, TID), PER							
ORAL							
				(Vitamin Nos)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea Vertigo	Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG (100 MG, BID), PER ORAL				Norflex (Orphenadrine Citrate)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyskinesia Loss Of Consciousness	Consumer	Neurontin (Gabapentin)	PS		ORAL
1800 MG (, DAILY), PER ORAL		Muscle Twitching Tinnitus Vision Blurred		(Baclofen) Prevacid (Lansoprazole)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410114-0Report Type:Periodic
Age:80 YR Gender:Male I/FU:I

Company Report #001-0945-980714

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
PER ORAL		Coordination Abnormal Vision Blurred	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Zantac (Ranitidine Hydrochloride)	C		
				Tenormin (Atenolol)	C		

Date:02/26/99ISR Number: 3410115-2Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980717

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (400 MG, TID)		Drug Interaction Dyskinesia	Consumer Health	Neurontin Capsules 100 Mg (Gabapentin)	PS		
		Insomnia	Professional				
80 MG (, QHS)		Mania Polyuria Tremor		Elavil (Amitriptyline Hydrochloride)	SS		
				Xanax (Alprazolam)	C		
				Klonopin (Clonazepam)	C		
				Phenergan (Promethazine Hydrochloride)	C		
				Imitrex (Sumatriptan)	C		
				(Potassium)	C		

Date:02/26/99ISR Number: 3410565-4Report Type:Periodic
Age:43 YR Gender:Male I/FU:I

Company Report #001-0945-980494

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Condition Aggravated	Consumer	Neurontin			

900 MG (300	Dyspepsia	(Gabapentin)	PS	ORAL
MG, TID) ,	Supraventricular			
PER ORAL	Extrasystoles			
		(Lorazepam)	C	
		Lodine (Etodolac)	C	

Date:02/26/99ISR Number: 3410567-8Report Type:Periodic Company Report #001-0945-980495
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Neurontin			
		Pain		(Gabapentin)	PS		ORAL
2400 MG (,		Paralysis					
DAILY), PER							
ORAL				Wellbutrin			
				(Amfebutamone			
				Hydrochloride)	C		
				Ritalin			
				(Methylphenidate			
				Hydrochloride)	C		
				Premarin (Estrogens			
				Conjugated)	C		
				Klonopin			
				(Clonazepam)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410570-8Report Type:Periodic
Age:52 YR Gender:Female I/FU:I

Company Report #001-0945-980497

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1200 MG (300 MG, QID), PER ORAL							

Prozac (Fluoxetine
Hydrochloride) C
Zyprexa (Olanzapine) C

Date:02/26/99ISR Number: 3410575-7Report Type:Periodic
Age:55 YR Gender:Male I/FU:I

Company Report #001-0945-980499

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pyrexia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
200 MG (,DAILY) , PER ORAL							

Unspecified Blood
Pressure Medication C
Unspecified "Water
Pill" C

Date:02/26/99ISR Number: 3410577-0Report Type:Periodic
Age:53 YR Gender:Female I/FU:I

Company Report #001-0945-980500

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal Feeling Jittery Irritability	Consumer	Neurontin(Gabapentin)	PS		ORAL
900 MG (300 MG, TID) ,							

PER ORAL

Muscle Contractions

Involuntary

Ambien (Zolpidem Tartrate)	C
Valium (Diazepam)	C
(Methadone)	C
Tenormin (Atenolol)	C
Premarin (Dstrogens Conjugated)	C
Docusate Sodium	C
Multivativimins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Calcium)	C
Acetaminophen (Paracetamol)	C

Date:02/26/99ISR Number: 3410579-4Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-980502

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsive Threshold Lowered	Health Professional	Neurontin (Gabapentin)	PS		
		Drug Interaction		Decongestants/Antihistamines	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410581-2Report Type:Periodic
Age:72 YR Gender:Male I/FU:I

Company Report #001-0945-980504

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID) , PER ORAL		Blood Glucose Increased Gastrointestinal Disorder	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				(Insulin)	C		
				(Digoxin)	C		
				(Hydrochlorothiazide)	C		
				(Potassium)	C		
				Coumadin (Warfarin Sodium)	C		
				Ambien (Zolpidem Tartrate)	C		

Date:02/26/99ISR Number: 3410582-4Report Type:Periodic
Age:63 YR Gender:Female I/FU:I

Company Report #001-0945-980505

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG (800 MG, TID), PER ORAL		Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
				(Baclofen)	C		
				Pamelor (Nortriptyline Hydrochloride)	C		
				Xanax (Alprazolam)	C		
				Sinequan (Doxepin Hydrochloride)	C		
				Elavil (Amitriptyline Hydrochloride)	C		

Date:02/26/99ISR Number: 3410584-8Report Type:Periodic
Age:35 YR Gender:Female I/FU:I

Company Report #001-0945-980506

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Movement Disorder	Consumer	Neurontin(Gabapentin)	PS		ORAL
3200 MG (800							
MG, QID) ,							
PER ORAL							

Effexor (Venlafaxine
Hydrochloride) C
Xanax (Alprazolam) C
Unspecified Muscle
Relaxant C

Date:02/26/99ISR Number: 3410586-1Report Type:Periodic
Age:53 YR Gender:Female I/FU:I

Company Report #001-0945-980507

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		White Blood Cell Count Decreased	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
800 - 1200 MG			Professional				
(WHEN NEEDED							
) , PER ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410591-5Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980508

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia Dermatitis	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
1200 MG (400 MG, TID), PER ORAL		Skin Nodule					
				Voltaren(Diclofenac Sodium) Vasoretic (Hydrochlorothiazide , Enalapril Maleate) Glucotrol (Glipizide) Xalatan(Latanoprost)	C C C C		

Date:02/26/99ISR Number: 3410594-0Report Type:Periodic
 Age:18 YR Gender:Female I/FU:I

Company Report #001-0945-980509

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination	Health Professional	Neurontin(Gabapentin)	PS		

Date:02/26/99ISR Number: 3410595-2Report Type:Periodic
 Age:52 YR Gender:Female I/FU:I

Company Report #001-0945-980512

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depressed Level Of Consciousness	Health Professional	Neurontin(Gabapentin)	PS		ORAL
300 MG (100 MG, TID) , PER ORAL		Dysphagia	Company				
		Dyspnoea	Representative				
		Muscular Weakness					

Date:02/26/99ISR Number: 3410596-4Report Type:Periodic Company Report #001-0945-980513
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pruritus Skin Nodule	Consumer	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3410598-8Report Type:Periodic Company Report #001-0945-980514
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Keratoconjunctivitis Sicca	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID) , PER ORAL				Prozac (Fluoxetine Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410600-3Report Type:Periodic
Age:33 YR Gender:Female I/FU:I

Company Report #001-0945-980515

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Balance Disorder Dizziness Feeling Drunk Gait Disturbance	Consumer	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3410602-7Report Type:Periodic
Age:59 YR Gender:Female I/FU:I

Company Report #001-0945-980516

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG (800 MG, TID) , PER ORAL		Dizziness Dysarthria Hypoaesthesia Sedation Vision Blurred	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3410604-0Report Type:Periodic
Age:48 YR Gender:Male I/FU:I

Company Report #001-0945-980519

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG (,DAILY), PER ORAL		Dysarthria Overdose Urinary Retention	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Topamax (Topiramate)	C
Tenormin (Atenolol)	C
Hytrin (Terazosin Hydrochloride)	C
Colace (Docusate Sodium)	C

Date:02/26/99ISR Number: 3410605-2Report Type:Periodic Company Report #001-0945-980520
Age:89 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG (,							
DAILY) , PER							
ORAL							

Date:02/26/99ISR Number: 3410607-6Report Type:Periodic Company Report #001-0945-980521
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG		Gait Disturbance					
(,DAILY), PER							
ORAL				Sulfa (Sulfaguanidine, Papaverine Hydrochloride, Clioquinol, Vitamins Lozol (Indapamide) (Spironolactone)	SS C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Vitamins Nos) C
 Asa (Acetylsalicylic Acid) C

Date:02/26/99ISR Number: 3410608-8Report Type:Periodic Company Report #001-0945-980463
 Age:76 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100-300		Abdominal Pain Upper Nausea	Consumer	Neurontin (Gabapentin)	PS		ORAL
DAILY, PER		Nervousness					
ORAL				Mylanta Prilosec (Omeprazole) Xanax (Alprazolam)	C C C		

Date:02/26/99ISR Number: 3410610-6Report Type:Periodic Company Report #001-0945-980464
 Age:50 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
800 MG		Balance Disorder Blood Pressure	Consumer	Neurontin (Gabapentin)	PS		ORAL
(,QHS), PER		Fluctuation					
ORAL		Dizziness					
		Fatigue Feeling Abnormal Lethargy		Premarin (Estrogens Conjugated) Provera (Medroxyprogesterone Acetate) Darvocet -N (Paracetamol, Dextrepropoxyphene) Lotensin (Benazepril Hydrochloride)	C C C		

Dyazide
(Hydrochlorothiazide
, Triamterene) C

Date:02/26/99ISR Number: 3410612-XReport Type:Periodic
Age:29 YR Gender:Male I/FU:I

Company Report #001-0945-980465

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Constipation Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
4300 MG (, DAILY), PER ORAL		Diarrhoea Dyspepsia Ear Infection Increased Appetite Lip Dry Overdose Tympanic Membrane Perforation Vaginal Candidiasis Weight Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410618-0Report Type:Periodic
 Age:31 YR Gender:Male I/FU:I

Company Report #001-0945-980466

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia Gastritis	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG (,DAILY), PER ORAL		Upper Respiratory Tract Infection	Professional				
				Percocet (Paracetamol, Oxycodone Hydrochloride, Oxycodone Meclizine (Meclozine Hydrochloride) Colace (Docusate Sodium)	SS C C		

Date:02/26/99ISR Number: 3410623-4Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #001-0945-980467

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Drug Interaction	Health Professional	Neurontin (Gabapentin) Amitriptyline (Amitriptyline)	PS SS		

Date:02/26/99ISR Number: 3410626-XReport Type:Periodic
 Age:72 YR Gender:Male I/FU:I

Company Report #001-0945-980468

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia Balance Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG TID), PER		Dizziness					

ORAL

Fall

Insomnia
Peripheral Coldness

Zestril (Lisinopril) C
(Insulin) C
(Atenolol) C
Enteric Coated
Aspirin
(Acetylsalicylic
Acid) C

Date:02/26/99ISR Number: 3410631-3Report Type:Periodic
Age:81 YR Gender:Male I/FU:I

Company Report #001-0945-980471

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG (,		Feelings Of Worthlessness Hyperhidrosis	Consumer	Neurontin (Gabapentin)	PS		ORAL
DAILY), PER		Mental Impairment					
ORAL		Pain In Extremity					
		Sedation		(Digoxin) (Quinidine)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410634-9Report Type:Periodic
Age:46 YR Gender:Female I/FU:I

Company Report #001-0945-980472

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 - 200 MG (,HS PRN)		Cardiac Disorder Feeling Abnormal	Health Professional	Neurontin (Gabapentin)	PS		
				Neurontin (Gabapentin)	SS		
				Estrace (Estradiol Lopressor (Metoprolol Tartrate)	C		
				Motrin (Ibuprofen)	C		
				Zyrtec (Certirizine Hydrochloride)	C		
				Tylenol (Paracetamol)	C		
				Vitamins	C		

Date:02/26/99ISR Number: 3410638-6Report Type:Periodic
Age:84 YR Gender:Male I/FU:I

Company Report #001-0945-980475

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (, ONE DOSE TAKEN) , PER ORAL		Bipolar I Disorder Depression Nervousness Schizophrenia	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3410643-XReport Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #001-0945-980476

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia	Consumer	Neurontin			

1200 MG (300
 MG QID) PER
 ORAL
 (Gabapentin) PS ORAL
 Prozac (Fluoxetine
 Hydrochloride) C
 Xanax (Alprazolam) C

Date:02/26/99ISR Number: 3410649-0Report Type:Periodic Company Report #001-0945-980479
 Age:55 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2700 MG (900 MG, TID), PER ORAL		Cataract Feeling Abnormal Headache Medication Error	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL				Neurontin (Gabapentin)	SS		ORAL
				Vicodin (Parametamol, Hydrocodone Bitartrate) (Nortriptyline) Norco	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Bitolterol) C

Date:02/26/99ISR Number: 3410654-4Report Type:Periodic Company Report #001-0945-980481
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hair Disorder	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3410659-3Report Type:Periodic Company Report #001-0945-980482
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Skin Pigmentation Disorder	Health Professional	Neurontin (Gabapentin)	PS		
				Tegretol (Carbamazepine)	C		

Date:02/26/99ISR Number: 3410665-9Report Type:Periodic Company Report #001-0945-980483
 Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bone Marrow Depression White Blood Cell Count	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (,		Decreased		Unspecified Medication Chemotherapeutic	SS		
DAILY), PER							
ORAL							

Date:02/26/99ISR Number: 3410667-2Report Type:Periodic Company Report #001-0945-980484
 Age:53 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Coordination Abnormal Dizziness	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
3600 MG (1200 MG, TID) PER ORAL		Vision Blurred Weight Increased		(Ibuprofen)	C		

Date:02/26/99ISR Number: 3410669-6Report Type:Periodic Company Report #001-0945-980485
Age:41 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anorgasmia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1500 MG, PER ORAL				Klonopin (Clonazepam)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410670-2Report Type:Periodic
 Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980487

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Neurontin Capsules			
3200 MG (800		Dyspnoea		400 Mg (Gabapentin)	PS		ORAL
MG, QID), PER		Overdose					
ORAL		Stupor					
500 MG (,		Syncope		Peganone (Ethotoin)	SS		ORAL
DAILY), PER							
ORAL				Chlorate	C		
				Haldol (Haloperidol)	C		
				Zoloft (Sertraline			
				Hydrochloride)	C		

Date:02/26/99ISR Number: 3410672-6Report Type:Periodic
 Age:36 YR Gender:Female I/FU:I

Company Report #001-0945-980488

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorgasmia	Consumer	Neurontin Capsules			
300 MG (,				300 Mg (Gabapentin)	PS		ORAL
QHS), PER							
ORAL				Synthroid			
				(Levothyroxine			
				Sodium)	C		
				Depakote (Valproate			
				Semisodium)	C		
				Serzone (Nefazodone			
				Hydrochloride)	C		

Date:02/26/99ISR Number: 3410674-XReport Type:Periodic Company Report #001-0945-980490
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		
450 MG (, 150 MG QAM; 300 MG				Neurontin Capsules 300 Mg (Gabapentin)	SS		ORAL
450 MG (, 150 MG QAM; 300 MG QHS) , PER ORAL							

Date:02/26/99ISR Number: 3410677-5Report Type:Periodic Company Report #001-0945-980492
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Decreased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
400 MG (, DAILY) , PER ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410707-0Report Type:Periodic
Age:75 YR Gender:Male I/FU:I

Company Report #001-0945-980402

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness	Health	Neurontin			
		Dry Mouth	Professional	(Gabapentin)	PS		ORAL
SEE TEXT, PER		Dry Throat					
ORAL		Dysarthria		Many Unspecified			
		Insomnia		Medications	C		
		Vision Blurred					

Date:02/26/99ISR Number: 3410712-4Report Type:Periodic
Age:35 YR Gender:Female I/FU:I

Company Report #001-0945-980403

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Haematuria	Consumer	Neurontin			
			Health	(Gabapentin)	PS		ORAL
1200 MG (300			Professional				
MG, QID), PER							
ORAL				Zoloft	C		
				Nolahist	C		
				Vitamins	C		
				Vitamin E	C		
				Calcium	C		

Date:02/26/99ISR Number: 3410769-0Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980404

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amenorrhoea	Health	Neurontin			
			Professional	(Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG							
(DAILY), PER							
ORAL							
				Calcium	C		
				Quinine	C		
				Elavil	C		
				Zantac	C		
				Nephrocaps	C		
				Sodium Bicarbonate	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Overdose	Health Professional	Neurontin (Gabapentin)	PS		ORAL
3600 MG		Pleural Effusion					
(DAILY), PER		Pneumonitis					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410777-XReport Type:Periodic
 Age:34 YR Gender:Female I/FU:I

Company Report #001-0945-980407

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, BID), PER ORAL		Dysarthria	Health	Neurontin Capsules	PS		ORAL
		Feeling Abnormal	Professional	300 Mg (Gabapentin)			
		Gait Disturbance	Company				
		Urticaria	Representative				
				Prozac	C		
				Klonopin	C		
				Dilantin	C		
				Phenergan	C		
				Indocin	C		

Date:02/26/99ISR Number: 3410779-3Report Type:Periodic
 Age:55 YR Gender:Female I/FU:I

Company Report #001-0945-980408

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (DAILY), PER ORAL		Dizziness	Health	Neurontin	PS		ORAL
		Euphoric Mood	Professional	(Gabapentin)			
		Feeling Drunk					
		Hypertensive Crisis					
				Avapro	C		
				Ticlid	C		
				(Amitriptyline)	C		

Date:02/26/99ISR Number: 3410781-1Report Type:Periodic
 Age:56 YR Gender:Female I/FU:I

Company Report #001-0945-980409

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG (100		Disturbance In Attention	Consumer	Neurontin	PS		ORAL
		Dry Mouth	Company	(Gabapentin)			

MG, BID), PER

Fatigue

Representative

ORAL

Synthroid	C
Paxil	C
Norvasc	C
Lipitor	C
Lasix	C
K-Dur	C
Premarin	C
(Isosorbide)	C
Imdur	C
Vicodin	C
Relafen	C

Date:02/26/99ISR Number: 3410784-7Report Type:Periodic Company Report #001-0945-980410
 Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Dizziness	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
100 MG (XL DOSE), PER		Sedation					

ORAL

Hytrin	C
Pepcid	C
Zestril	C
(Hydrochlorothiazide	
)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410786-0Report Type:Periodic
Age:35 YR Gender:Female I/FU:I

Company Report #001-0945-980411

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion Oedema	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG							
(DAILY), PER							
ORAL							
				Dilantin	C		
				Klonopin	C		

Date:02/26/99ISR Number: 3410790-2Report Type:Periodic
Age:44 YR Gender:Male I/FU:I

Company Report #001-0945-980414

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache Reflexes Abnormal	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900-1200 MG							
(DAILY), PER							
ORAL							
		Sedation					
				Pepcid	C		

Date:02/26/99ISR Number: 3410794-XReport Type:Periodic
Age:56 YR Gender:Female I/FU:I

Company Report #001-0945-980415

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation Increased Appetite	Health Professional	Neurontin (Gabapentin)	PS		ORAL
700 MG							
(DAILY), PER							
ORAL							
		Keratoconjunctivitis					
		Sicca					
		Nausea		Lamictal (Lamotrigine)	SS		

Atarax C
Ativan C

Date:02/26/99ISR Number: 3410799-9Report Type:Periodic Company Report #001-0945-980418
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menstruation Irregular	Health Professional Company Representative	Neurontin (Gabapentin) Contraceptive Nos	PS C		

Date:02/26/99ISR Number: 3410803-8Report Type:Periodic Company Report #001-0945-980419
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Neuropathy Peripheral	Consumer	Neurontin 300 Mg (Gabapentin)	PS		ORAL
2700 MG (900							
MG, TID), PER							
ORAL				Premarin	C		
				Provera	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410807-5Report Type:Periodic
Age:31 YR Gender:Female I/FU:I

Company Report #001-0945-980424

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation Visual Disturbance	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
600 MG (300 MG, BID), PER ORAL							

Proventil	C
Vancenase	C
Deconsal	C

Date:02/26/99ISR Number: 3410819-1Report Type:Periodic
Age:26 YR Gender:Female I/FU:I

Company Report #001-0945-980425

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper Aggression	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (DAILY), PER ORAL		Headache Insomnia Sedation Vomiting					

Date:02/26/99ISR Number: 3410826-9Report Type:Periodic
Age:7 YR Gender:Male I/FU:I

Company Report #001-0945-980428

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Coagulation Time Prolonged	Health Professional	Neurontin (Gabapentin)	PS		ORAL
2400 MG (DAILY), PER ORAL		Drug Interaction Ecchymosis					

Overdose

Robitussin
(Guaifenesin) SS
Triaminic
(Mepyramine Maleate,
Pheniramine Maleate,
Phenylpropanolamine
Hydrochloride) SS

Date:02/26/99ISR Number: 3410833-6Report Type:Periodic
Age:35 YR Gender:Male I/FU:I

Company Report #001-0945-980430

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1000 MG (500 MG, BID), PER ORAL		Neutrophil Count Decreased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Adriamycin	C		

Date:02/26/99ISR Number: 3410839-7Report Type:Periodic
Age:41 YR Gender:Female I/FU:I

Company Report #001-0945-980431

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2700 MG (DAILY), PER ORAL		Abdominal Pain Crying Dyspnoea Hyperhidrosis Irritable Bowel Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Levsin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Clonazepam) C
 Propoxyphene C

Date:02/26/99ISR Number: 3410845-2Report Type:Periodic
 Age:39 YR Gender:Female I/FU:I

Company Report #001-0945-980433

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite Weight Increased	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
100 MG							
(DAILY), PER							
ORAL				Klonopin	C		

Date:02/26/99ISR Number: 3410898-1Report Type:Periodic
 Age:40 YR Gender:Female I/FU:I

Company Report #001-0945-980761

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Abnormal	Health Professional	Neurontin(Gabapentin)	PS		ORAL
2400 MG (800							
MG, TID), PER							
ORAL				Glucophage (Metformin Hydrochloride)	C		
				Restoril (Temazepam)	C		
				Wellbutrin (Amfebutam one Hydrochloride)	C		

Date:02/26/99ISR Number: 3410902-0Report Type:Periodic
 Age:75 YR Gender:Female I/FU:I

Company Report #001-0945-980762

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Joint Swelling	Consumer	Neurontin(Gabapentin			

Paraesthesia

)

PS

ORAL

1100

MG(,DAILY),

PER ORAL

Ultram(Tramadol	
Hydrochloride)	C
Xanax(Alprazolam)	C
Vicodin(Paracetamol,	
Hydrocodone	
Bitartrate)	C

Date:02/26/99ISR Number: 3410907-XReport Type:Periodic

Company Report #001-0945-980763

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Faecal Incontinence	Health Professional	Neurontin(Gabapentin)	PS		ORAL

1200

MG(,DAILY),

PER ORAL

Zoloft(Sertraline	
Hydrochloride)	C
Valium(Diazepam)	C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410911-1Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-980764

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vitamin K Deficiency	Health Professional	Neurontin(Gabapentin)	PS		

Date:02/26/99ISR Number: 3410913-5Report Type:Periodic
Age:61 YR Gender:Male I/FU:I

Company Report #001-0945-980766

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Foot Fracture Hypoaesthesia	Consumer	Neurontin(Gabapentin)	PS		ORAL
1800		Paraesthesia					
MG(,DAILY),							
PER ORAL				(Amoxicillin)	C		
				Minocin(Minocycline)	C		
				Calan(Verapamil Hydrochloride)	C		

Date:02/26/99ISR Number: 3410914-7Report Type:Periodic
Age:71 YR Gender:Female I/FU:I

Company Report #001-0945-980767

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Health Professional	Neurontin(Gabapentin)	PS		ORAL
1800							
MG(,DAILY),							
PER ORAL							

Date:02/26/99ISR Number: 3410917-2Report Type:Periodic
Age:57 YR Gender:Female I/FU:I

Company Report #001-0945-980768

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Consumer	Neurontin Capsules			
		Insomnia		300 Mg(Gabapentin)	PS		ORAL
900 MG(300		Mental Disorder					
MG, TID), PER		Oedema Peripheral					
ORAL				(Clonazepam)	C		

Date:02/26/99ISR Number: 3410920-2Report Type:Periodic Company Report #001-0945-980769
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Neurontin Capsules			
		Drug Interaction		100 Mg(Gabapentin)	PS		
		Dyspnoea		Septra			
		Vertigo		Ds(Sulfamethoxazole, Trimethoprim)	SS		
				Levaquin(Levofloxacin)	C		
				(Meclofenamate Sodium)	C		
				Oxycontin(Oxycodone Hydrochloride)	C		
				Zoloft(Sertraline Hydrochloride)	C		
				Claritin(Loratadine)	C		
				(Alprazolam)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Oxycodone) C

Date:02/26/99ISR Number: 3410922-6Report Type:Periodic Company Report #001-0945-980770
 Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysarthria	Consumer	Neurontin(Gabapentin)	PS		ORAL
300		Fall					
		Sedation					
MG(,DAILY),							
PER ORAL							
				Lopressor(Metoprolol Tartrate)	C		

Date:02/26/99ISR Number: 3410926-3Report Type:Periodic Company Report #001-0945-980771
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Unevaluable Event	Health Professional	Neurontin(Gabapentin)	PS		ORAL
900							
MG(,DAILY),							
PER ORAL							

Date:02/26/99ISR Number: 3410930-5Report Type:Periodic Company Report #001-0945-980773
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination, Visual	Consumer	Neurontin Capsules 300 Mg(Gabapentin)	PS		ORAL
2700 MG (900							
MG, TID), PER							

ORAL

Humulin N(Insulin Human Injection, Isophane)	C
Humulin R(Insulin Human)	C
Calan(Verapamil Hydrochloride)	C
Depakote(Valproate Semisodium)	C
Prozac(Fluoxetine Hydrochloride)	C
(Potassium)	C

Date:02/26/99ISR Number: 3410932-9Report Type:Periodic
 Age:76 YR Gender:Male I/FU:I

Company Report #001-0945-980774

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Faecal Incontinence	Consumer	Neurontin Capsules			
		Gingival Bleeding		100 Mg(Gabapentin)	PS		ORAL
600 MG(300		Gingivitis					
MG, BID), PER		Urinary Incontinence					

ORAL

Asa(Acetylsalicylic Acid)	C
Monopril(Fosinopril Sodium)	C
(Insulin)	C
Glyburide(Glibenclam	

Freedom Of Information (FOI) Report

ide) C

Date:02/26/99ISR Number: 3410936-6Report Type:Periodic
Age:57 YR Gender:Female I/FU:I

Company Report #001-0945-980777

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (300 MG, QID), PER ORAL		Abdominal Pain Upper Faeces Discoloured Gastric Haemorrhage Headache Myalgia Nausea Stomach Discomfort	Consumer	Neurontin Capsules 300 Mg(Gabapentin)	PS		ORAL
				Avonex(Interferon Beta)	C		

Date:02/26/99ISR Number: 3410941-XReport Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980778

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
X2 (, QAM AND QPM), PER ORAL		Convulsion Cushingoid Cyanosis Drug Interaction Peripheral Coldness	Consumer	Neurontin(Gabapentin) Dilantin Infatabs 50 Mg(Phenytoin Sodium)	PS SS		ORAL
				Colace(Docusate Sodium)	SS		

Date:02/26/99ISR Number: 3410945-7Report Type:Periodic
Age:27 YR Gender:Female I/FU:I

Company Report #001-0945-980782

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Paraesthesia	Health	Neurontin Capsules			

600 MG (300

Professional

300 Mg(Gabapentin)

PS

ORAL

MG, BID), PER

ORAL

Zoloft (Sertraline
Hydrochloride)
(Trazodone)
(Lithium)
Ativan(Lorazepam)

C
C
C
C

Date:02/26/99ISR Number: 3410948-2Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980784

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatomyositis	Consumer Health Professional	Neurontin(Gabapentin)	PS		

Date:02/26/99ISR Number: 3410953-6Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980785

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Health Professional	Neurontin(Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3410955-XReport Type:Periodic
 Age:73 YR Gender:Male I/FU:I

Company Report #001-0945-980786

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal	Consumer	Neurontin(Gabapentin)	PS		ORAL
600 MG							
(,QHS), PER							
ORAL							

Hytrin(Terazosin Hydrochloride)	C
Children'S Aspirin(Acetylsalicylic Acid)	C

Date:02/26/99ISR Number: 3410958-5Report Type:Periodic
 Age:54 YR Gender:Female I/FU:I

Company Report #001-0945-980780

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination	Consumer Health	Neurontin Capsules 100 Mg(Gabapentin)	PS		ORAL
400 MG (100			Professional				
MG, QID), PER							
ORAL							

Lopressor(Metoprolol Tartrate)	C
Vasotec(Enalapril Maleate)	C
Zocor(Simvastatin) (Nortriptyline)	C
Zantac(Ranitidine Hydrochloride)	C
Premarin(Estrogens Conjugated)	C
K-Dur(Potassium Chloride)	C
Motrin(Ibuprofen)	C
Lasix(Furosemide)	C

Date:02/26/99ISR Number: 3410963-9Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980781

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Haematuria International Normalised Ratio Increased	Health Professional	Neurontin(Gabapentin) Coumadin(Warfarin Sodium)	PS SS		

Date:02/26/99ISR Number: 3411003-8Report Type:Periodic
Age:61 YR Gender:Female I/FU:I

Company Report #001-0945-980731

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated Hyperthyroidism	Consumer	Neurontin(Gabapentin)	PS		ORAL
1800 MG							
(600MG, TID)							
, PER ORAL				(Estrogen Nos) (Propranolol)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3411006-3Report Type:Periodic
Age:18 YR Gender:Female I/FU:I

Company Report #001-0945-980733

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accommodation Disorder	Consumer	(Gabapentin)	PS		ORAL
600 MG (200		Blepharospasm	Health				
MG, TID) ,		Dystonia	Professional				
PER ORAL				Solu-Medrol (Methylprednisolone)	C		
				Zanaflex (Tizanidine)	C		
				(Clonidine)	C		

Date:02/26/99ISR Number: 3411008-7Report Type:Periodic
Age:48 YR Gender:Female I/FU:I

Company Report #001-0945-980734

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urticaria	Consumer	Neurontin(Gabapentin)	PS		ORAL
SEE TEXT, PER				(Prednisone)	C		
ORAL							

Date:02/26/99ISR Number: 3411009-9Report Type:Periodic
Age:28 YR Gender:Female I/FU:I

Company Report #001-0945-980735

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia Coordination Abnormal	Health Professional	Neurontin (Gabapentin)	PS		
SEE TEXT,		Hallucination		(Seroquel)	C		
UNKNOWN							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Paraesthesia	Health Professional	Neurontin(Gabapentin)	PS		ORAL
300 MG (,							
DAILY) , PER							
ORAL				Pamelor (Nortriptyline Hydrochloride))	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Deafness	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300							
MG, TID) ,							
PER ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3411014-2Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980739

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID)		Oedema	Health Professional Company Representative	Neurontin (Gabapentin)	PS		
				Risperdal (Risperidone)	C		
				Prozac (Fluoxetine Hydrochloride)	C		
				Wellbutrin (Amfebutamone Hydrochloride)	C		

Date:02/26/99ISR Number: 3411019-1Report Type:Periodic
Age:46 YR Gender:Female I/FU:I

Company Report #001-0945-980740

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG (100 MG, BID) , PER ORAL		Gingival Erosion Stomatitis	Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
				Axid (Nizatidine)	C		

Date:02/26/99ISR Number: 3411022-1Report Type:Periodic
Age:35 YR Gender:Male I/FU:I

Company Report #001-0945-980742

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (, DAILY) , PER		Ejaculation Disorder Impaired Healing Libido Decreased	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

ORAL

Luvox (Fluvoxamine
Maleate) C
Klonopin
(Clonazepam) C

Date:02/26/99ISR Number: 3411024-5Report Type:Periodic
Age:40 YR Gender:Female I/FU:I

Company Report #001-0945-980743

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nightmare	Health Professional	Neurontin (Gabapentin)	PS		ORAL
PER ORAL							

Oxycodone C
Restoril (Temazepam) C
Prilosec
(Omeprazole) C
Lorcet (Paracetamol,
Hydrocodone
Bitartrate C

Date:02/26/99ISR Number: 3411026-9Report Type:Periodic
Age:65 YR Gender:Female I/FU:I

Company Report #011-0945-980744

Outcome PT
Dizziness
Night Sweats

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Syncope

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
400 MG (200 MG, BID) , PER ORAL		Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
			St. John'S Wort (Hypericum Extract)	C		
			Dilantin (Phenytoin Sodium)	C		

Date:02/26/99ISR Number: 3411027-0Report Type:Periodic Company Report #001-0945-980745
Age:38 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	300 MG (100MG, TID) , PER ORAL		Bladder Pain Pollakiuria Urethral Pain		Neurontin (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3411029-4Report Type:Periodic Company Report #001-0945-980746
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	600 MG (200 MG, TID)		Arthropathy Fall	Consumer Health	Neurontin (Gabapentin)	PS		
			Injury	Professional				
	2 MG (, QHS)		Sedation Vision Blurred		Zanzflex (Tizanidine)	SS		
					(Clonidine)	C		

Medrol
(Methylprednisolone) C

Date:02/26/99ISR Number: 3411030-0Report Type:Periodic
Age:34 YR Gender:Female I/FU:I

Company Report #001-0945-980750

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bronchospasm Chest Discomfort	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
300 MG (100 MG, TID) , PER ORAL		Dyspnoea					

Klonopin
(Clonazepam) C
Imitrex (Sumatriptan
Succinate) C

Date:02/26/99ISR Number: 3411031-2Report Type:Periodic
Age:91 YR Gender:Female I/FU:I

Company Report #001-0945-980751

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1800 MG (600 MG, THREE TIMES DAILY) , PER ORAL							

(Glyceryl

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Freedom Of Information (FOI) Report

Trinitrate) C
 Xanax (Alprazolam) C
 Pepcid (Famotidine) C

Date:02/26/99ISR Number: 3411032-4Report Type:Periodic
 Age:71 YR Gender:Female I/FU:I

Company Report #001-0945-980753

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (200 MG, BID) , PER ORAL		Dry Mouth Stomatitis	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

Zantac (Ranitidine Hydrochloride) C

Date:02/26/99ISR Number: 3411034-8Report Type:Periodic
 Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980754

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (, NIGHTLY) , PER ORAL		Diarrhoea Flatulence	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

(Trazodone) C

Date:02/26/99ISR Number: 3411035-XReport Type:Periodic
 Age:62 YR Gender:Male I/FU:I

Company Report #001-0945-980755

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300		Arthralgia Joint Stiffness	Consumer	Neurontin (Gabapentin)	PS		ORAL

MG , TID) ,

PER ORAL

Avonex (Interferon Beta)	C
(Baclofen)	C
Xalatan (Latanoptost)	C
Ocupress (Carteolol Hydrochloride)	C
Alphagan (Brimonidine Tartrate)	C
(Amantadine)	C

Date:02/26/99ISR Number: 3411038-5Report Type:Periodic Company Report #001-0945-980756
 Age:13 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (200 MG , BID) ,		Abnormal Behaviour Feeling Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
PER ORAL		Hostility Lethargy Malaise Nausea Paranoia Psychomotor Hyperactivity Thinking Abnormal					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3411042-7Report Type:Periodic
 Age:76 YR Gender:Male I/FU:I

Company Report #001-0945-980757

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Effect Decreased Drug Interaction	Consumer	Neurontin(Gabapentin)	PS		ORAL
300 MG (, HS) , PER		Sleep Disorder					
ORAL				Mavik (Trandolapril)	SS		ORAL
4 MG (, DAILY) , PER							
ORAL							

Date:02/26/99ISR Number: 3412077-0Report Type:Periodic
 Age:79 YR Gender:Female I/FU:I

Company Report #001-0945-980818

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Constipation	Consumer	Neurontin Capsules 400mg (Gabapetin)	PS		ORAL
400 MG (DAILY) PER				Sinemet (Levodopa, Carbidopa)	C		
ORAL				Permax (Pergolide Mesilate)	C		
				Ambien (Zolpidem Tartrate)	C		
				Tylenol (Paracetamol)	C		
				Aspirin (Acetylsalicylic Acid)	C		
				Metamucil (Psyllium Hydrophilic Mucilloid)	C		

Date:02/26/99ISR Number: 3412078-2Report Type:Periodic
Age:74 YR Gender:Male I/FU:I

Company Report #001-0945-980819

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema	Health Professional	Neurontin Capsules 300mg (Gabapetin)	PS		ORAL
2400 MG (600 MG, QID) PER ORAL							

Date:02/26/99ISR Number: 3412080-0Report Type:Periodic
Age:77 YR Gender:Male I/FU:I

Company Report #001-0945-980820

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eye Pain	Consumer	Neurontin (Gabapetin)	PS		ORAL
200 MG (DAILY), PER ORAL							
				Mavik (Trandolapril)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3412082-4Report Type:Periodic
Age:51 YR Gender:Male I/FU:I

Company Report #001-0945-980821

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Ageusia Anosmia	Consumer	Neurontin Capsules 100mg (Gabapetin)	PS		ORAL
100 MG		Deafness					
(DAILY) PER		Ear Infection					
ORAL		Sinus Congestion Tinnitus		Rezulin(Troglitazone) Glucophage (Metformin Hydrochloride) Trazodone	C C C		

Date:02/26/99ISR Number: 3412084-8Report Type:Periodic
Age:61 YR Gender:Female I/FU:I

Company Report #001-0945-980822

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperglycaemia	Consumer	Neurontin (Gabapetin)	PS		ORAL
600 MG							
(DAILY), PER							
ORAL				Nph Insulin (Insulin Injection, Isophane) Humulin Regular(Insulin Human)	C C		

Date:02/26/99ISR Number: 3412086-1Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980823

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence	Consumer	Neurontin Capsules			

Headache
 900 MG (300
 MD, TID) PER
 ORAL
 300mg (Gabapetin) PS ORAL
 Demulen (Mestranol,
 Etnodiol Diacetate) C

Date:02/26/99ISR Number: 3412089-7Report Type:Periodic Company Report #001-0945-980825
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Dyspnoea	Health Professional	Neurontin Capsules 100mg (Gabapetin)	PS		ORAL

400 MG (200
 MG, BID) PER
 ORAL
 Flexeril
 (Cyclobenzaprine
 Hydrochloride) C
 Prozac (Fluoxetine
 Hydrochloride) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3412090-3Report Type:Periodic
Age:37 YR Gender:Female I/FU:I

Company Report #001-0945-980826

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1600 MG (400 MG, QID), PER ORAL		Eczema Pain In Extremity Paraesthesia	Consumer	Neurontin Capsules 400mg (Gabapetin)	PS		ORAL
				Klonopin(Clonazepam)	C		
				Ambien(Zolpidem Tartrate)	C		
				Prozac(Fluoxetine Hydrochloride)	C		
				Oral Contraceptive Nos	C		

Date:02/26/99ISR Number: 3412092-7Report Type:Periodic
Age:18 YR Gender:Male I/FU:I

Company Report #001-0945-980828

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID) PER ORAL		Amnesia Drug Interaction	Consumer	Neurontin Capsules 100mg (Gabapetin)	PS		ORAL
				Feverfew(Herbal Extracts Nos)	SS		
				Nortriptyline	C		

Date:02/26/99ISR Number: 3412093-9Report Type:Periodic
Age:58 YR Gender:Male I/FU:I

Company Report #001-0945-980834

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pancreatitis	Consumer	Neurontin			

800 MG

(Gabapetin)

PS

ORAL

(DAILY), PER

ORAL

Vicodin(Paracetamol,
Hydrocodone
Bitartrate)
Darvocet
(Paracetamol,
Dextropropoxyphene)

C

C

Date:02/26/99ISR Number: 3412095-2Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #001-0945-980835

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Ecchymosis	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
600 MG (300 MG, BID) PER							

ORAL

Zoloft (Sertraline
Hydrochloride)

C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3412096-4Report Type:Periodic
Age:30 YR Gender:Unknown I/FU:I

Company Report #001-0945-980836

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Health Professional	Neurontin Capsules 300mg (Gabapetin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL							

Date:02/26/99ISR Number: 3412098-8Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980837

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation	Health Professional	Neurontin(Gabapentin)	PS		ORAL
1200 MG (300 MG, QID), PER ORAL							

Tylenol W/Codeine
No. 3(Codeine
Phosphate,
Paracetamol) SS

Date:02/26/99ISR Number: 3412100-3Report Type:Periodic
Age:30 YR Gender:Female I/FU:I

Company Report #001-0945-980841

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Increased	Health Professional	Neurontin(Gabapentin)	PS		ORAL
1200 MG (600MG, Q12H), PER							

ORAL

Proventil
Inhaler(Salbutamol) C
Zocor(Simvastatin) C
Trazodone C
Humalin N(Insulin) C

Date:02/26/99ISR Number: 3412102-7Report Type:Periodic
Age:55 YR Gender:Female I/FU:I

Company Report #001-0945-980842

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred	Consumer	Neurontin Capsules 300 Mg(Gabapentin)	PS		ORAL
300 MG							
(,QHS), PER							

ORAL

Rezulin
(Troglitazone) C
Lanoxin(Digoxin) C
Stomach Medicine C
Prednisone C
Glucotrol
(Glipizide) C

Mouth Ulceration

Health Professional

Neurontin (Gabapentin)

PS

Date:02/26/99ISR Number: 3412109-XReport Type:Periodic
Age:42 YR Gender:Female I/FU:I

Company Report #001-0945-980848

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
800 MG (400 MG, BID) PER ORAL		Pruritus Urine Abnormality	Consumer	Neurontin Capsules 400 Mg(Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3412115-5Report Type:Periodic
Age:48 YR Gender:Female I/FU:I

Company Report #001-0945-980787

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID),		Chest Pain Headache Nausea	Consumer	Neurontin (Gabapentin)	PS		
				Monopril (Fosinopril Sodium)	C		
				Klonopin (Clonazepam)	C		
				Evista (Raloxifene)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3412118-0Report Type:Periodic
Age:25 YR Gender:Female I/FU:I

Company Report #001-0945-980788

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chloasma	Consumer Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3412121-0Report Type:Periodic
Age:23 YR Gender:Female I/FU:I

Company Report #001-0945-980789

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Fatigue	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

1200 MG (,
300MG QAM &
QNOON, 600MG
QPM), PER
ORAL

Date:02/26/99ISR Number: 3412123-4Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980791

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Abnormal	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3412124-6Report Type:Periodic
Age:61 YR Gender:Female I/FU:I

Company Report #001-0945-980792

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased	Consumer	Neurontin Capsules			

600 MG (300

300 Mg (Gabapentin) PS

ORAL

MG, BID), PER

ORAL

(Atenolol)	C
Estrogens	C
Zoloft (Sertraline Hydrochloride)	C
Elavil (Amitriptyline Hydrochloride)	C
Valium (Diazepam)	C
(Acetylsalicylic Acid)	C

Date:02/26/99ISR Number: 3412127-1Report Type:Periodic
Age:42 YR Gender:Female I/FU:I

Company Report #001-0945-980793

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Rigidity Myalgia	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
1200 MG							

(,QHS), PER

ORAL

Levothroid (Levothyroxine Sodium)	C
(Lithium)	C
(Chloral Hydrate)	C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3412129-5Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-980797

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema	Health Professional	Neurontin (Gabapentin)	PS		
				Unspecified Analgesics	C		

SEE TEXT

Date:02/26/99ISR Number: 3412131-3Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980799

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blister Dermatitis	Health Professional	Neurontin (Gabapentin)	PS		
				Daypro (Oxaprozin)	SS		
				Dyazide (Hydrochlorothiazide , Triamterene)	SS		

Date:02/26/99ISR Number: 3412133-7Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-980800

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Vasculitis	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3412135-0Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980802

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Feeling Abnormal	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

300 MG,

DAILY, PER

ORAL; 600 MG

(300 MG,

BID), PER

Date:02/26/99ISR Number: 3412136-2Report Type:Periodic

Company Report #001-0945-980804

Age:42 YR Gender:Male

I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea Skin Discolouration	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

300 MG (,

DAILY), PER

ORAL

Vicodin
 (Paracetamol,
 Hydrocodone
 Bitartttrate) C
 Oxycontin (Oxycodone
 Hydrochloride) C
 (Influenza Virus
 Vaccine Polyvalent) C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3412137-4Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980805

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chromaturia	Health Professional	Neurontin (Gabapentin)	PS		
				"Golden Seal" Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Wellbutrin (Amfebutamone Hydrochloride))	SS		
					C		
					C		

Date:02/26/99ISR Number: 3412139-8Report Type:Periodic
Age: Gender:I/FU:I

Company Report #001-0945-980806

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral Overdose	Health Professional	Neurontin (Gabapentin)	PS		
4800 MG (, DAILY),							

Date:02/26/99ISR Number: 3412141-6Report Type:Periodic
Age:65 YR Gender:Male I/FU:I

Company Report #001-0945-980809

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspepsia Feeling Abnormal	Consumer Health	Neurontin(Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL		Oesophagitis	Professional				
				Relafen (Nabumetone) Pravachol (Pravastatin Sodium)	C		
					C		

(Diazepam)

C

Date:02/26/99ISR Number: 3412143-XReport Type:Periodic
Age:52 YR Gender:Female I/FU:I

Company Report #001-0945-980810

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (150 MG, BID), PER ORAL		Tooth Disorder	Consumer	Neurontin(Gabapentin)	PS		ORAL

Klonopin
(Clonazepam)

C

Date:02/26/99ISR Number: 3412145-3Report Type:Periodic
Age:47 YR Gender:Female I/FU:I

Company Report #001-0945-980811

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		Balance Disorder Hyperhidrosis Palpitations Skin Warm Weight Increased	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
				Unspecified Steroid (Acetylsalicylic	C		

Relafen (Nabumetone) C
Synthroid
(Levothyroxine
Sodium) C
(Vitamins Nos) C

Date:02/26/99ISR Number: 3412151-9Report Type:Periodic Company Report #001-0945-980816
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dystonia	Consumer	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3412152-0Report Type:Periodic Company Report #001-0945-980817
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL

1800 MG (600
MG, TID), PER

ORAL

Prempro

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Freedom Of Information (FOI) Report

(Medroxyprogesterone
Acetate, Estrogens
Conjugated) C
Paxil (Paroxetine
Hydrochloride) C

Date:02/26/99ISR Number: 3413091-1Report Type:Periodic
Age:56 YR Gender:Male I/FU:I

Company Report #001-0945-980586

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG (,DAILY), PER ORAL		Anxiety Fatigue Nausea	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Zestril (Lisinopril) Aspirin (Acetylsalicylic Acid) Glyburide (Glibenclamide) Centrum Silver (Ascorbid Acid, Tocopheryl Acetate, Retinol, Zinc, Calcium, Vitamins Vitamin E (Tocopherol)	C C C C C		

Date:02/26/99ISR Number: 3413092-3Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #001-0945-980590

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (,DAILY), PER ORAL		Arrhythmia Electrocardiogram Abnormal Heart Rate Increased	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

Provera (Medroxy
Progesterone
Acetate) C
Humalog (Insulin
Lispro) C
Monopril (Fosinopril
Sodium) C
Propulsid
(Cisapride) C
Premarin (Estrogens
Conjugated) C
Wellbutrin
(Amfebutamone
Hydrochloride) C

Date:02/26/99ISR Number: 3413094-7Report Type:Periodic
Age:52 YR Gender:Female I/FU:I

Company Report #001-0945-980594

Outcome PT
Asthma
Coordination Abnormal
Depression
Disturbance In Attention

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Dizziness Nausea Overdose	Report Source	Product	Role	Manufacturer	Route
4800 MG (UNK), PER ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Unspecified Thyroid Medication	C		
				Unspecified Hormone Medication	C		

Date:02/26/99ISR Number: 3413095-9Report Type:Periodic Company Report #001-0945-980595
Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT Fatigue Feeling Hot Nausea	Report Source	Product	Role	Manufacturer	Route
UNK, UNKNOWN			Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3413096-0Report Type:Periodic Company Report #001-0945-980596
Age:52 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT Coordination Abnormal Fall Muscular Weakness	Report Source	Product	Role	Manufacturer	Route
600 MG (200 MG, TID), PER ORAL			Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
				Vitamin B12 (Cyanocobalamin)	C		
				(Vitamins Nos) Levothroid (Levothyroxine)	C		

Sodium)

C

Date:02/26/99ISR Number: 3413097-2Report Type:Periodic
Age:45 YR Gender:Female I/FU:I

Company Report #001-0945-980598

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Skin Pruritus	Health Professional	Neurontin (Gabapentin)	PS		
1500 MG (300 MG, X 5 DAILY), UNKNOWN				Demerol (Pethidine Hydrochloride)	C		
				Inderal (Propranolol Hydrochloride)	C		
				Valium (Diazepam)	C		
				Lasix (Furosemide)	C		

Date:02/26/99ISR Number: 3413098-4Report Type:Periodic
Age:30 YR Gender:Male I/FU:I

Company Report #001-0945-980599

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
600 MG (300							

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MG, BID), PER

ORAL

(Amikacin)	C
(Imipenem)	C
Morphine	C
Dulcolax (Bisacodyl)	C
Compazine	
(Prochlorperazine	
Edisylate)	C
Colace (Docusate	
Sodium)	C
Multivitamins	
(Ergocalciferol,	
Ascorbid Acid, Folic	
Acid, Thiamine	
Hydrochloride,	C
(Ferrous Sulfate)	C
Folic Acid	C
Enoxaparin Sodium	
(Heparin-Fraction,	
Sodium Salt)	C
Narcan (Naloxone	
Hydrochloride)	C
Tylenol	
(Paracetamol)	C

Date:02/26/99ISR Number: 3413101-1Report Type:Periodic
Age:76 YR Gender:Male I/FU:I

Company Report #001-0945-980601

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Glucose Increased	Health	Neurontin			
		Dizziness	Professional	(Gabapentin)	PS		ORAL
1200 MG (400		Oedema Peripheral					
MG, TID), PER		Weight Increased					

ORAL

Capozide	
(Hydrochlorothiazide	
, Captopril)	C
Glucotrol	
(Glipizide)	C
Persantine	

(Dipyridamole) C
Lanoxin (Digoxin) C
Multivitamins
(Ergocalciferol,
Ascorbid Acid, Folic
Acid, Thiamine
Hydrochloride, C
Hytrin (Terazosin
Hydrochloride) C
Asa (Acetylsalicylic
Acid) C

Date:02/26/99ISR Number: 3413102-3Report Type:Periodic
Age:22 YR Gender:Female I/FU:I

Company Report #001-0945-980602

Outcome PT
Alcohol Intolerance
Asthenia
Blood Glucose Abnormal
Decreased Appetite

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Depression Hyperacusis Irritability	Report Source	Product	Role	Manufacturer	Route
1200 MG	(,DAILY), PER	Nasal Congestion	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:02/26/99ISR Number: 3413104-7Report Type:Periodic Company Report #001-0945-980603
 Age:55 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG	(,DAILY), PER	Bone Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL							

(Thyroid) C
 Unspecified Hormones C
 Unspecified Allergy
 Medication C

Date:02/26/99ISR Number: 3413106-0Report Type:Periodic Company Report #001-0945-980604
 Age:57 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID) PER		Platelet Count Decreased	Consumer Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
ORAL							

Zoloft (Sertraline
Hydrochloride) C

Zantac (Ranitidine Hydrochloride) C
 Cytotec (Misoprostol) C
 Propulsid (Cisapride) C
 Zyloprim (Allopurinol) C
 Ativan (Lorazepam) C
 Lamictal (Lamotrigine) C
 (Colchicine) C

Date:02/26/99ISR Number: 3413107-2Report Type:Periodic Company Report #001-0945-980605
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
600 MG			Professional				
(,DAILY), PER							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3413109-6Report Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980610

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG (100 MG, BID), PER ORAL		Abnormal Behaviour Disorientation Fatigue Headache Vision Blurred	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Paxil (Paroxetine Hydrochloride) Nasalcort	C C		

Date:02/26/99ISR Number: 3413111-4Report Type:Periodic
Age:58 YR Gender:Male I/FU:I

Company Report #001-0945-980612

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (,DAILY), PER ORAL		Urinary Tract Infection	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Procardia Xl (Nifedipine) Aerobid (Flunisolide) Persantine (Dipyridamole)	C C C		

Date:02/26/99ISR Number: 3413112-6Report Type:Periodic
Age:82 YR Gender:Male I/FU:I

Company Report #001-0945-980619

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (, X1		Feeling Abnormal Stupor	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

DOSE), PER

ORAL

Lescol (Fluvastatin Sodium) C
Asa (Acetylsalicylic Acid) C
Hytrin (Terazosin Hydrochloride) C
Norvasc (Amlodipine Besilate) C

Date:02/26/99ISR Number: 3413114-XReport Type:Periodic
Age:72 YR Gender:Male I/FU:I

Company Report #001-0945-980620

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG		Erectile Dysfunction Faecal Incontinence	Health Professional	Neurontin (Gabapentin)	PS		ORAL

(,DAILY), PER

ORAL

Vitamin B12
(Cyanocobalamin-N) C
Ventolin
(Salbutamol) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3413115-1Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980622

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK, UNKNOWN		Unevaluable Event	Consumer	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3413116-3Report Type:Periodic
 Age:48 YR Gender:Female I/FU:I

Company Report #001-0945-980624

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (400 MG,TID), PER ORAL		Vertigo	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3413117-5Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980625

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2000 MG (400 MG, 5 TIMES A DAY), PER ORAL		Pneumonia Weight Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Zoloft (Sertraline
Hydrochloride) C

Date:02/26/99ISR Number: 3413118-7Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #001-0945-980626

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1600 MG (400 MG, QID), PER ORAL		Skin Ulcer Weight Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Darvocet (Paracetamol, Dextropropoxyphene) Soma (Carisoprodol)	C C		

Date:02/26/99ISR Number: 3413119-9Report Type:Periodic Company Report #001-0945-980290
 Age:41 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
90MG (30MG, TID) PER ORAL		Anorgasmia	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3413120-5Report Type:Periodic Company Report #001-0945-980291
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
240MG (DAILY),		Paraesthesia	Consumer Health Professional	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3413121-7Report Type:Periodic
Age:79 YR Gender:Female I/FU:I

Company Report #001-0945-980293

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema Hypoaesthesia Paraesthesia	Consumer	Neurontin (Gabapentin)	PS		ORAL
120MG (40MG, TID) , PER ORAL				Tenormin (Atenolol)	C		

Date:02/26/99ISR Number: 3413122-9Report Type:Periodic
Age:74 YR Gender:Male I/FU:I

Company Report #001-0945-980295

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Herpes Zoster	Consumer Health	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
2400 MG (600 MG, QID), PER ORAL			Professional	Lipitor (Atorvastatin)	C		
				Lopressor (Metoprolol Tartrate)	C		
				Nortriptyline	C		
				Asa (Acetylsalicylic Acid)	C		

Date:02/26/99ISR Number: 3413123-0Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-980296

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea Food Interaction	Health Professional	Neurontin (Gabapentin)	PS		

PER ORAL Nausea Shellfish SS ORAL
 Vomiting

Date:02/26/99 ISR Number: 3413124-2 Report Type:Periodic Company Report #001-0945-980297
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety	Consumer	Neurontin Capsules			
		Balance Disorder		100 Mg (Gabapentin)	PS		ORAL
SEE TEXT, PER		Drug Withdrawal Syndrome					
ORAL		Ear Disorder		Advil (Ibuprofen)	C		
		Fatigue		Floxin (Ofloxacin)	C		
		Nausea		Imitrex			
		Tension		(Sumatriptan)	C		
		Tinnitus		Clozapine	C		
		Vision Blurred		Nervous Tension	C		
				Anxiety	C		
				Nausea	C		
				Withdrawal Symptoms	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3413125-4Report Type:Periodic
Age:12 YR Gender:Female I/FU:I

Company Report #001-0945-980298

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gastrointestinal Disorder Viral Infection Vomiting	Consumer Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3413126-6Report Type:Periodic
Age:44 YR Gender:Female I/FU:I

Company Report #001-0945-980301

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Decreased	Health Professional Company Representative	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
1200 MG (400 MG, TID), PER ORAL				Wellbutrin (Amfebutamone Hydrochloride)	C		

Date:02/26/99ISR Number: 3413127-8Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980302

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue Sedation	Consumer	Neurontin (Gabapentin) Ativan (Lorazepam)	PS C		

Date:02/26/99ISR Number: 3413128-XReport Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980303

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction	Consumer	Neurontin			

1200 MG (DAILY) PER ORAL (Gabapentin) PS ORAL

Date:02/26/99ISR Number: 3413129-1Report Type:Periodic Company Report #001-0945-980305
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Gingivitis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1 GM (DAILY)							
PER ORAL FEW							
WEEKS							

Haldol (Haloperidol) C
 Klonopin (Clonazepam) C
 Depakote (Valproate Semisodium) C

Date:02/26/99ISR Number: 3413130-8Report Type:Periodic Company Report #001-0945-980306
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia Sinusitis	Consumer	Neurontin (Gabapentin)	PS		ORAL
400 MG							

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Freedom Of Information (FOI) Report

(DAILY) PER

ORAL

Carafate
(Sucralfate) C
Prilosec
(Omeprazole) C
Gaviscon (Aluminium
Hydroxide Gel,
Dried, Magnesium
Trisilicate) C

Date:02/26/99ISR Number: 3413131-XReport Type:Periodic Company Report #001-0945-980309
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
500 MG (,SEE TEXT), PER ORAL		Ageusia Amnesia Condition Aggravated Dizziness Flatulence	Consumer Health Professional	Neurontin (Gabapentin) Doxidan (Dantron, Docusate Calcium)	PS C		ORAL

Date:02/26/99ISR Number: 3413132-1Report Type:Periodic Company Report #001-0945-980310
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK, COUPLES OF WEEKS		Abnormal Behaviour Logorrhoea	Consumer	Neurontin (Gabapentin) Baclofen	PS C		

Date:02/26/99ISR Number: 3413133-3Report Type:Periodic Company Report #001-0945-980314
Age:75 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
500 MG (100 MG, FIVE TIMES DAILY), PER ORAL		Condition Aggravated Fatigue Feeling Abnormal Frequent Bowel Movements Nausea Oedema Peripheral Paraesthesia Sedation	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
				Tylenol (Paracetamol) Zantac (Ranitidine Hydrochloride)	C C		

Date:02/26/99ISR Number: 3413134-5Report Type:Periodic Company Report #001-0945-983015
Age:46 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (,DAILY) , PER ORAL SEE TEXT		Vision Blurred Visual Acuity Reduced	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Dilantin (Phenytoin Sodium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3413136-9Report Type:Periodic
Age:25 YR Gender:Female I/FU:I

Company Report #001-0945-980316

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety Insomnia	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
SEE TEXT, PER							
ORAL							
Orgasm Abnormal							
Weight Increased							

Date:02/26/99ISR Number: 3413139-4Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #001-0945-98317

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Balance Disorder Condition Aggravated	Consumer	Neurontin(Gabapentin)	PS		ORAL
900 MG							
(300MG, TID),							
PER ORAL							
Gait Disturbance							
Migraine							
Thinking Abnormal							
Betaseron (Glucose, Albumin Human, Interferon Beta)							
C							

Date:02/26/99ISR Number: 3413140-0Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980319

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Disorientation Drug Withdrawal Syndrome	Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG							
(,DAILY) ,							
PER ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3000 MG (1000 MG, TID) PER ORAL		Convulsion Overdose	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Imuran (Azathioprine)	C		
				Lorazepam	C		
				Isordil (Isosorbide Dinitrate)	C		
				Premarin (Estrogens)	C		
				Lasix (Furosemide)	C		
				Ranitidine	C		
				Lescol (Fluvastatin Sodium)	C		
				Norvasc (Amlodipine Besilate)	C		
				Acyclovir (Aciclovir)	C		
				Albuterol (Salbutamol)	C		
				Atrovent (Fenoterol Hydrobromide, Ipratropium Bromide)	C		
				Proventil (Salbutamol)	C		

Oral Mucosal Blistering
Stomatitis

Health
Professional

Neurontin Capsules
300 Mg (Gabapentin)

PS

ORAL

SEE TEXT, PER

ORAL

Date:02/26/99ISR Number: 3414417-5Report Type:Periodic
Age:51 YR Gender:Male I/FU:I

Company Report #001-0945-980561

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, BID), PER		Abdominal Pain Feeling Drunk Insomnia Vision Blurred	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

ORAL

Date:02/26/99ISR Number: 3414420-5Report Type:Periodic
Age:68 YR Gender:Female I/FU:I

Company Report #001-0945-980563

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (,EVERY NIGHT), PER		Dermatitis Diarrhoea Dizziness Emotional Distress Malaise Sedation Vision Blurred	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Klonopin (Clonazepam)	C		

ORAL

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Freedom Of Information (FOI) Report

Vitamins Nos C
 Asa
 (Acetylsalicyclic
 Acid) C

Date:02/26/99ISR Number: 3414422-9Report Type:Periodic Company Report #001-0945-980564
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Photosensitivity Reaction	Consumer	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3414424-2Report Type:Periodic Company Report #001-0945-980565
 Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3414426-6Report Type:Periodic Company Report #001-0945-980566
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorgasmia	Consumer	Neurontin (Gabapentin)	PS		ORAL

2400 MG
 (,DAILY), PER

ORAL

Rilutek (Riluzole) C

Date:02/26/99ISR Number: 3414427-8Report Type:Periodic Company Report #001-0945-980567
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral Paraesthesia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
100 MG (,X1 DOSE), PER ORAL				Birth Control Pills Tegretol (Carbamazepine)	C C		

Date:02/26/99ISR Number: 3414428-XReport Type:Periodic Company Report #001-0945-980568
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain Medication Error	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
1200 MG (400 MG, TID), PER ORAL							

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Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414432-1Report Type:Periodic
Age:56 YR Gender:Female I/FU:I

Company Report #001-0945-980639

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG DAILY PER ORAL		Blood Pressure Increased Haematuria Medication Error Renal Colic Therapeutic Response Unexpected	Consumer	Neurontin (Gabapentin) (Baclofen) Aspirin (Acetylsalicylic Acid) Synthroid (Levothyroxine Sodium)	PS C C C		ORAL

Date:02/26/99ISR Number: 3414433-3Report Type:Periodic
Age:52 YR Gender:Female I/FU:I

Company Report #001-0945-980569

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Cholesterol Increased Blood Triglycerides Increased	Health Professional	Neurontin (Gabapentin) Insulin 70/30 (Insulin Human, Insulin Human Injection, Isophane) Albuterol (Salbutamol) Digoxin Zestril (Lisinopril) Serevent (Salmeterol Xinafoate) Pravachol (Pracastatin Sodium) Prednisone Rezulin (Troglitazone) Prempro (Medroxyprogesterone Acetate, Estrogens Conjugated) Nasalide	PS C C C C C C C C C C		

(Flunisolide)	C
Lasix (Furosemide)	C
Citracal (Calcium Citrate)	C
Vitamin E (Tocopherol)	C
Zoloft (Sertraline Hydrochloride)	C
Buspar (Buspirone Hydrochloride)	C
Tegretol (Carbamazepine)	C
Trazodone	C
Xanax (Alprazolam)	C
Pulmicort (Budesonide)	C

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Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414435-7Report Type:Periodic
Age:44 YR Gender:Female I/FU:I

Company Report #001-0945-980570

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Liver Function Test Abnormal	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG							
(,DAILY), PER							
ORAL							

Date:02/26/99ISR Number: 3414437-0Report Type:Periodic
Age:47 YR Gender:Female I/FU:I

Company Report #001-0945-980642

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oligomenorrhoea Weight Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300							
MG TID) PER							
ORAL							

Date:02/26/99ISR Number: 3414438-2Report Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980572

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysarthria Ecchymosis	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
300 MG							
(,DAILY), PER		Feeling Drunk					
ORAL							

Amantadine	C
Eskalith (Lithium Carbonate)	C
Klonopin (Clonazepam)	C

Synthroid
 (Levothyroxine
 Sodium) C
 Serax (Oxazepam) C
 Zyprexa (Olanzapine) C

Date:02/26/99ISR Number: 3414440-0Report Type:Periodic
 Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980574

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (,DAILY), PER ORAL		Blindness Visual Disturbance	Consumer	Neurontin(Gabapentin)	PS		ORAL

Prozac (Fluoxetine
 Hydrochloride) C
 Wellbutrin
 (Amfebutamone
 Hydrochloride) C
 Klonopin
 (Clonazepam) C
 Premarin (Estrogens
 Conjugated) C
 Ogen (Estropipate) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414441-2Report Type:Periodic
Age:88 YR Gender:Female I/FU:I

Company Report #001-0945-980643

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anuria	Consumer	Neurontin			
		Increased Appetite		(Gabapentin)	PS		ORAL
400 MG (200		Lethargy					
MG BID) PER		Nervousness					
ORAL							

Date:02/26/99ISR Number: 3414442-4Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980575

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Insomnia	Health	Neurontin Capsules			
		Mania	Professional	100 Mg (Gabapentin)	PS		
400 MG (, 100		Medication Error					
MG QAM; 300		Tachycardia					
QHS), UNKNOWN		Thinking Abnormal		Wellbutrin			
				(Amfebutamone			
				Hydrochloride)	C		

Date:02/26/99ISR Number: 3414444-8Report Type:Periodic
Age:75 YR Gender:Female I/FU:I

Company Report #001-0945-980576

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Consumer	Neurontin			
		Nausea		(Gabapentin)	PS		ORAL
100 MG		Pharyngolaryngeal Pain					
(,DAILY), PER							
ORAL				Fosamax (Alendronate			
				Sodium)	C		

Tylenol #3 (Codeine
Phosphate,
Paracetamol) C

Date:02/26/99ISR Number: 3414445-XReport Type:Periodic Company Report #001-0945-980644
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Creatine Phosphokinase Increased Muscular Weakness	Health Professional	Neurontin (Gabapentin) Rezulin (Troglitazone) (Insulin) Prandin (Repaglinide) (Verapamil)	PS SS C C C		

Date:02/26/99ISR Number: 3414447-3Report Type:Periodic Company Report #001-0945-980580
Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2700 MG (,DAILY),		Arthralgia Myalgia	Health Professional	Neurontin (Gabapentin) Depakote (Valproate Semisodium)	PS C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414448-5Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-980646

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Arthralgia Malaise Myalgia Systemic Lupus Erythematosis	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3414449-7Report Type:Periodic
 Age:42 YR Gender:Male I/FU:I

Company Report #001-0945-980583

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		Skin Odour Abnormal	Consumer Health Professional	Neurontin(Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3414450-3Report Type:Periodic
 Age:40 YR Gender:Female I/FU:I

Company Report #001-0945-980647

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
75 MG DAILY PER ORAL		Asthenia Drug Interaction Syncope	Consumer	Neurontin (Gabapentin) Nardil (Phenelzine Sulfate)	PS SS		ORAL
				Propulsid (Cisapride) Prilosec (Omeprazole) Estratest Hs (Methyltestosterone, Estrogens)	C C		

Esterified) C
 Artane
 (Trihexyphenidyl
 Hydrochloride) C
 Librium
 (Chlordiazepoxide
 Hydrochloride) C
 (Perphenazine) C
 (Oxazepam) C

Date:02/26/99ISR Number: 3414451-5Report Type:Periodic Company Report #001-0945-980584
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		Back Pain Pain In Extremity Sedation	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Atenolol Prinivil (Lisinopril) Voltaren (Diclofenac Sodium) Zantac (Ranitidine	C C C C		

FDA - Adverse Event Reporting System (AERS)

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Hydrochloride) C
 Mevacor (Lovastatin) C
 Premarin (Estrogens
 Conjugated) C
 Elavil
 (Amitriptyline
 Hydrochloride) C
 Diazepam C
 Vicodin
 (Paracetamol,
 Hydrocodone
 Bitartrate) C

Date:02/26/99ISR Number: 3414452-7Report Type:Periodic Company Report #001-0945-980648
 Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Withdrawal Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
2700 MG (300 MG , 9X DAILY) PER ORAL							

Percocet
 (Paracetamol,
 Oxycodone
 Hydrochloride,
 Oxycodone C
 Eye Drops C

Date:02/26/99ISR Number: 3414453-9Report Type:Periodic Company Report #001-0945-980585
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction Sedation	Health Professional Company Representative	Neurontin (Gabapentin) Tegretol (Carbamazepine) Elavil	PS SS		

(Amitriptyline
Hydrochloride) SS

Date:02/26/99ISR Number: 3414454-0Report Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980651

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Health Professional	Neurontin (Gabapentin)	PS		ORAL

1200 MG (600

MG BID) PER

ORAL

Serzone (Nefazodone
Hydrochloride) C
Xanax (Alprazolam) C
Plaquenil
(Hydroxychloroquine
Phosphate) C

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Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414455-2Report Type:Periodic
Age:36 YR Gender:Female I/FU:I

Company Report #001-0945-980652

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG TID) PER ORAL				(Valproate Sodium)	C		

Date:02/26/99ISR Number: 3414457-6Report Type:Periodic
Age:67 YR Gender:Female I/FU:I

Company Report #001-0945-980653

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Neurontin (Gabapentin)	PS		
TID				Unspecified Medications	C		

Date:02/26/99ISR Number: 3414458-8Report Type:Periodic
Age:63 YR Gender:Female I/FU:I

Company Report #001-0945-980654

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia Joint Stiffness	Consumer	Neurontin (Gabapentin)	PS		ORAL
2700 MG (900 MG TID) PER ORAL				Serax (Oxazepam)	C		
				Pamelor (Nortriptyline Hydrochloride)	C		
				(Morphine)	C		

Date:02/26/99ISR Number: 3414459-XReport Type:Periodic
Age:67 YR Gender:Female I/FU:I

Company Report #001-0945-980655

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain Nausea	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG							
(DAILY) PER							
ORAL							

Synthroid (Levothyroxine Sodium)	C
Pravachol (Pravastatin Sodium)	C

Date:02/26/99ISR Number: 3414460-6Report Type:Periodic
Age:36 YR Gender:Female I/FU:I

Company Report #001-0945-980657

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eyelid Oedema Mydriasis	Consumer Company	Neurontin (Gabapentin)	PS		ORAL
300 MG (100							
MG TID) PER							
ORAL							

Representative

Prozac (Fluoxetine Hydrochloride)	C
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414461-8Report Type:Periodic
Age:33 YR Gender:Male I/FU:I

Company Report #001-0945-980658

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4000 MG (DAILY) PER ORAL		Amnesia Chills Diarrhoea Drug Withdrawal Syndrome Dyspnoea Ear Pain Electroencephalogram Abnormal Erectile Dysfunction Fatigue Feeling Abnormal Headache Hypoaesthesia Libido Decreased Night Sweats Overdose Palpitations Pollakiuria Sleep Disorder Tremor Vision Blurred	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Zoloft (Sertraline Hydrochloride)	C		
				Ativan (Lorazepam)	C		
				Luvox (Fluvoxamine Maleate)	C		
				Serzone (Nefazodone Hydrochloride)	C		
				Klonopin (Clonazepam)	C		
				Elavil (Amitriptyline Hydrochloride)	C		

Date:02/26/99ISR Number: 3414462-XReport Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-980659

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Weight Increased	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3414463-1Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-980660

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia Restlessness	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 - 1500							
MG(DAILY)							
PER ORAL				Unspecified Antipsychotic Medications	C		

Date:02/26/99ISR Number: 3414464-3Report Type:Periodic Company Report #001-0945-980662
Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Sleep Walking	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300							
MG TID) PER							
ORAL				(Clonidine)	C		
				(Folic Acid)	C		
				Serevent (Salmeterol Xinafoate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Timoptic (Timolol
Maleate) C

Date:02/26/99ISR Number: 3414465-5Report Type:Periodic
Age:49 YR Gender:Male I/FU:I

Company Report #001-0945-980667

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2500 MG DAILY PER ORAL		Feeling Abnormal Mental Impairment Sedation	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Prilosec
(Omeprazole) C
Unspecified Inhaler C

Date:02/26/99ISR Number: 3414466-7Report Type:Periodic
Age:66 YR Gender:Female I/FU:I

Company Report #001-0945-980668

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG QHS PER ORAL		Oedema Peripheral	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

Vancenase
(Beclometasone
Dipropionate) C
Aerobid
(Flunisolide) C
Atrovent (Fenoterol
Hydrobromide,
Ipratropium Bromide) C
Albuterol
(Salbutamol) C
(Nortriptyline) C
(Cyproheptadine) C
Elmiron (Pentosan
Polysulfate Sodium) C

Date:02/26/99ISR Number: 3414467-9Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980669

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Visual Disturbance	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG TID) PER ORAL				Serzone (Nefazodone Hydrochloride) Buspar (Buspirone Hydrochloride)	C C		

Date:02/26/99ISR Number: 3414468-0Report Type:Periodic
Age:43 YR Gender:Male I/FU:I

Company Report #001-0945-980386

Outcome	PT
	Drug Effect Decreased Euphoric Mood Fatigue Paranoia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sedation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1200 MG (300 MG, QID), PER ORAL		Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
			Klonopin (Clonazepam)	C		
			(Valproic Acid)	C		
			Zyprexa (Olanzapine)	C		

Date:02/26/99ISR Number: 3414469-2Report Type:Periodic Company Report #001-0945-980388
Age:76 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (DAILY) (SEE IMAGE)		Balance Disorder Change Of Bowel Habit Dry Mouth Dysarthria	Consumer	Neurontin (Gabapentin)	PS		
		Fall Flatulence Insomnia Vision Blurred		Zostrix (Capsaicin) Tylenol Pm (Diphenhydramine, Paracetamol) Lodine (Etodolac)	C C C		

Date:02/26/99ISR Number: 3414470-9Report Type:Periodic Company Report #001-0945-980393
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia Anaemia Confusional State Parkinsonian Gait	Consumer	Neurontin (Gabapentin) Dilantin (Phenytoin Sodium)	PS C		

(Felbamate)

C

Date:02/26/99ISR Number: 3414471-0Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980394

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG							
(DAILY), PER							
ORAL							

Dolobid (Diflunisal) C
Vitamins C

Date:02/26/99ISR Number: 3414472-2Report Type:Periodic
Age:77 YR Gender:Male I/FU:I

Company Report #001-0945-980395

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction Nausea	Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG (100							
MG,BID), PER							
ORAL							
				Buspar (Buspirone Hydrochloride)	SS		ORAL
20 MG (10 MG,							

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Freedom Of Information (FOI) Report

BID), PER

ORAL

Procardia Xl (Nifedipine)	C
Tenormin (Atenolol)	C
(Allopurinol)	C
Prilosec (Omeprazole)	C
Coumadin (Warfarin Sodium)	C
Propulsid (Cisapride)	C

Date:02/26/99ISR Number: 3414473-4Report Type:Periodic
Age:76 YR Gender:Female I/FU:I

Company Report #001-0945-980397

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG		Feeling Abnormal Feeling Hot	Consumer	Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER		Pain					
ORAL		Paraesthesia					
		Vasodilatation		Glucophage (Metformin Hydrochloride)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Plendil (Felodipine)	C		
				Coumadin (Warfarin Sodium)	C		
				Xanax (Alprazolam)	C		
				Macrochantin (Nitrofurantoin)	C		
				Vitamin C (Ascorbic Acid)	C		
				Vitamin E (Tocopherol)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Feeling Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG		Sedation					
(DAILY), PER							
ORAL				Adalat (Nifedipine) Aspirin (Acetylsalicylic Acid)	C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Porphyrin Metabolism Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
100-200 MG							
(DAILY), PER							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Herbs C

Date:02/26/99ISR Number: 3414476-XReport Type:Periodic Company Report #001-0945-980400
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion	Health Professional Company Representative	Neurontin (Gabapentin) Dilantin (Phenytoin Sodium)	PS C		

Date:02/26/99ISR Number: 3414477-1Report Type:Periodic Company Report #001-0945-980401
 Age:56 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID), PER ORAL		Lymphadenopathy	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Elavil (Amitriptyline Hydrochloride)	C		

Date:02/26/99ISR Number: 3414478-3Report Type:Periodic Company Report #001-0945-980324
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Neuropathy Peripheral	Health Professional	Neurontin (Gabapentin) Alcohol (Ethanol)	PS SS		

Date:02/26/99ISR Number: 3414479-5Report Type:Periodic Company Report #001-0945-980325
 Age:28 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG		Anorgasmia Libido Decreased	Other	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
(DAILY), PER							
ORAL							

Date:02/26/99ISR Number: 3414480-1Report Type:Periodic Company Report #001-0945-980329
Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		
				Carbocaine (Mepivacaine Hydrochloride)	SS		
				Lamictal (Lamotrigine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414481-3Report Type:Periodic
 Age: Gender: I/FU:I

Company Report #001-0945-980339

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Neuropathy Peripheral	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3414482-5Report Type:Periodic
 Age:54 YR Gender:Female I/FU:I

Company Report #001-0945-980330

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Difficulty In Walking Fatigue	Consumer	Neurontin (Gabapentin)	PS		ORAL

1200 MG (400
 MG, TID), PER

ORAL

Topimax (Topiramate) C

Date:02/26/99ISR Number: 3414483-7Report Type:Periodic
 Age:29 YR Gender:Male I/FU:I

Company Report #001-0945-980340

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Extrapyramidal Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL

1200 MG (,

DAILY--IN

DIVIDED

DOSES) , PER

ORAL

Date:02/26/99ISR Number: 3414484-9Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980333

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
10	DAY	Amnesia	Health Professional	Neurontin (Gabapentin)	PS		
				(Valproate Sodium)	C		
				Pamelor (Nortriptyline Hydrochloride)	C		
				(Nortriptyline)	C		

Date:02/26/99ISR Number: 3414485-0Report Type:Periodic Company Report #001-0945-980343
 Age:62 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
700 MG (,		Arthralgia Myalgia	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
DAILY) , PER							
ORAL				Trazodone	C		

Date:02/26/99ISR Number: 3414486-2Report Type:Periodic Company Report #001-0945-980334
 Age:41 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300		Medication Error Weight Increased	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
MG, TID), PER							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Prozac (Fluoxetine
Hydrochloride) C

Date:02/26/99ISR Number: 3414487-4Report Type:Periodic
Age:4 YR Gender:Male I/FU:I

Company Report #001-0945-980335

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID), PER		Convulsion Insomnia Psychomotor Hyperactivity	Health Professional	Neurontin (Gabapentin)	PS		ORAL

ORAL

(Chloral Hydrate) C
Unisom (Doxylamine
Succinate) C

Date:02/26/99ISR Number: 3414488-6Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980344

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (, DAILY) , PER		Drug Interaction Hallucination	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

ORAL

Nardil Tablets 15 Mg
(Phenelzine Sulfate) SS

45 MG (,
DAILY) , PER

ORAL

(Lithium) C
Plaquenil
(Hydroxy-Chloroquine
Phosphate) C

Ambien (Zolpidem
Tartate) C

Date:02/26/99ISR Number: 3414489-8Report Type:Periodic Company Report #001-0945-980347
Age:56 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (200 MG, BID) , PER ORAL		Blood Pressure Decreased Paraesthesia Speech Disorder	Consumer	Neurontin(Gabapentin) Dilantin (Phenytoin Sodium) Cardizem (Diltiazem Hydrochloride) Prilosec (Omeprazole) Zoloft (Sertraline Hydrochloride) Urispas (Florvoxate Hydrochloride) Ambien (Zolpidem Tartrate) (Atenolol) (Colchicine)	PS C C C C C C C C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414490-4Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980336

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Memory Impairment Weight Increased	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
300 MG (100 MG, TID), PER ORAL							

Date:02/26/99ISR Number: 3414491-6Report Type:Periodic
 Age:22 YR Gender:Female I/FU:I

Company Report #001-0945-980350

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia Chest Pain	Consumer	Neurontin (Gabepentin)	PS		ORAL
1200 MG (, DAILY), PER ORAL							
Drug Level Below Therapeutic							
				Dilantin(Phenytoin Sodium) (Folic Acid)	C C		

Date:02/26/99ISR Number: 3414492-8Report Type:Periodic
 Age:42 YR Gender:Female I/FU:I

Company Report #001-0945-980352

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flat Affect Panic Attack	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
100 MG (, DAILY) , PER ORAL							
				Tremor			
				Klonopin (Clonazepam)	C		

(Meprobamate)

C

Date:02/26/99ISR Number: 3414493-XReport Type:Periodic
Age:62 YR Gender:Female I/FU:I

Company Report #001-0945-980337

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG							
(DAILY), PER							
ORAL							

Date:02/26/99ISR Number: 3414494-1Report Type:Periodic
Age:25 YR Gender:Male I/FU:I

Company Report #001-0945-980353

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error Mydriasis	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
1200 MG (400							
MG, TID) ,							
PER ORAL							

Depakote (Valproate Semisodium)	C
Cogentin (Benzatropine Mesilate)	C
Ativan (Lorazepam)	C
Prolixin (Fluphenazine)	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C

Date:02/26/99ISR Number: 3414495-3Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980338

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
PER ORAL		Drug Level Below Therapeutic	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3414496-5Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980355

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema		Neurontin (Gabapentin)	PS		
				Wellbutrin (Amfebutamone Hydrochloride)	C		

Date:02/26/99ISR Number: 3414497-7Report Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980522

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
SEE TEXT, PER ORAL		Dizziness Nausea Tremor	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				(Insulin)	C		
				Inderal (Propranolol Hydrochloride)	C		
				Ogen (Estropipate)	C		
				Tegretol (Carbamazepine)	C		
				Lasix (Furosemide)	C		
				Betimol (Timolol)	C		
				Pepcid (Famotidine)	C		
				Cardizem (Diltiazem)			

Hydrochloride) C

Date:02/26/99ISR Number: 3414498-9Report Type:Periodic Company Report #001-0945-980524
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Other	Neurontin (Gabapentin) (Cocaine)	PS C		

Date:02/26/99ISR Number: 3414499-0Report Type:Periodic Company Report #001-0945-980525
Age:17 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pollakiuria Polyuria	Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, BID), PER ORAL				Effexor (Venlafaxine Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414500-4Report Type:Periodic
 Age:31 YR Gender:Female I/FU:I

Company Report #001-0945-980361

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (400 MG, TID) , PER ORAL		Clumsiness Disorientation Fall Fatigue Speech Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Tegretol (Carbamazepine)	C		
				Paxil (Paroxetine Hydrochloride)	C		
				Klonopin (Clonazepam)	C		

Date:02/26/99ISR Number: 3414501-6Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #001-0945-980368

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (, DAILY), PER ORAL		Diplopia Dizziness Dry Mouth Dystonia Muscle Spasms Vomiting	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3414502-8Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-980526

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
SEE TEXT		Oedema	Health Professional Company	Neurontin (Gabapentin)	PS		

Representative

Date:02/26/99ISR Number: 3414503-XReport Type:Periodic
 Age:35 YR Gender:Female I/FU:I

Company Report #001-0945-980369

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2100 MG (, DAILY) , PER ORAL		Conjunctival Hyperaemia Ear Pain Food Allergy Keratoconjunctivitis Sicca Lymphadenopathy Medication Error Oral Mucosal Blistering Salivary Gland Enlargement Tongue Oedema	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Pepcid (Famotikine)	C		
				Tylenol # 3 (Codeine Phosphate, Paracetamol)	C		
				Sporanox (Itraconazole)	C		
				Zoloft (Sertraline Hydrochloride)	C		
				Soma (Carisoprodol)	C		
				(Vitamins Nos)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414504-1Report Type:Periodic
Age:31 YR Gender:Male I/FU:I

Company Report #001-0945-980529

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eye Disorder Tic	Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		
300 MG (, 100							
MG QAM, 200							
MG QPM,							
UNKNOWN							

Date:02/26/99ISR Number: 3414505-3Report Type:Periodic
Age:61 YR Gender:Male I/FU:I

Company Report #001-0945-980372

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Weight Increased	Consumer	Neurontin(Gabapentin)	PS		ORAL
900 MG (300							
MG, TID) ,							
PER ORAL							

Date:02/26/99ISR Number: 3414506-5Report Type:Periodic
Age:60 YR Gender:Male I/FU:I

Company Report #001-0945-980530

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoglycaemia	Health Professional	Neurontin (Gabapentin) Diabeta (Glibenclamide)	PS C		

Date:02/26/99ISR Number: 3414507-7Report Type:Periodic
Age:43 YR Gender:Male I/FU:I

Company Report #001-0945-980374

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema Peripheral	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
2400 MG (800							
MG, TID) ,							
PER ORAL							

Date:02/26/99ISR Number: 3414508-9Report Type:Periodic Company Report #001-0945-980531
Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue Lethargy	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG (400							
MG, TID) ,							
PER ORAL							

Coumadin (Warfarin Sodium) C
Hytrin (Terazosin Hydrochloride) C
Vasotec (Enalapril Maleate) C
Zyloprim (Allopurinol) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414509-0Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980375

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia Asthenia	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG,TID) PER ORAL		Coordination Abnormal Flushing					

Date:02/26/99ISR Number: 3414510-7Report Type:Periodic
 Age:45 YR Gender:Female I/FU:I

Company Report #001-0945-980376

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper Constipation	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
450 MG (150 MG, TID) , PER ORAL		Diarrhoea Dyspepsia					
		Fatigue Flatulence Nausea Pyrexia		Xanax (Alprazolam) Klonopin (Clonazepam) (Cortisone)	C C C		

Date:02/26/99ISR Number: 3414511-9Report Type:Periodic
 Age:80 YR Gender:Female I/FU:I

Company Report #001-0945-980532

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Abnormal	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG (X1 DOSE), PER ORAL							

Date:02/26/99ISR Number: 3414512-0Report Type:Periodic Company Report #001-0945-980537
Age:89 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Health Professional	Neurontin (Gabapentin)	PS		
300 MG (,							
DAILY),							

Date:02/26/99ISR Number: 3414513-2Report Type:Periodic Company Report #001-0945-980377
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain Hypoaesthesia	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
100 MG		Insomnia					
(,DAILY), PER		Pain In Extremity					
ORAL		Paraesthesia					

Date:02/26/99ISR Number: 3414514-4Report Type:Periodic Company Report #001-0945-980538
Age:71 YR Gender:Female I/FU:I

Outcome	PT
	Asthenia Condition Aggravated Confusional State Convulsion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Difficulty In Walking Drug Level Above Therapeutic	Report Source	Product	Role	Manufacturer	Route
1200 MG (,DAILY) PER ORAL		Fall Sedation Urinary Incontinence	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG, TID) PER ORAL				Dilantin (Phenytoin Sodium)	SS		ORAL
				Asa (Acetylsalicylic Acid)	C		
				(Calcium)	C		
				(Vitamins Nos)	C		
				Unspecified Spray	C		

Date:02/26/99ISR Number: 3414515-6Report Type:Periodic
Age:80 YR Gender:Female I/FU:I

Company Report #001-0945-980383

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (,300 MG TID), PER ORAL		Flatulence	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Glucophage (Metformin Hydrochloride)	C		
				Paxil (Paroxetine Hydrochloride)	C		
				Glucomide (Metformin Hydrochloride, Glibenclamide)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Dyspnoea	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL		Nervousness		Valium (Diazepam) Darvocet-N (Paracetamol, Dextropropoxyphene) (Thyroid) Lopid(Gemfibrozil) Dyazide (Hydrochlorothiazide , Triamterene) Inderal (Propranolol Hydrochloride)	C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414517-XReport Type:Periodic
Age:55 YR Gender:Male I/FU:I

Company Report #001-0945-980384

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Neurontin (Gabapentin)	PS		ORAL
3600 MG							
(,DAILY), PER							
ORAL							

Zocof (Simvastatin) C
Baby Aspirin C

Date:02/26/99ISR Number: 3414518-1Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980385

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 ,G							
(,DAILY), PER							
ORAL							

Tenormin (Atenolol) C
(Allopurinol) C
Procardia Xl
(Nifedipine) C
Prilosec
(Omeprazole) C
Propulsid
(Cisapride) C
Coumadin (Warfarin
Sodium) C

Date:02/26/99ISR Number: 3414519-3Report Type:Periodic
Age:80 YR Gender:Female I/FU:I

Company Report #001-0945-980721

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

200 MG (100 MG, BID), PER ORAL	Blood Creatinine Increased Oedema Peripheral Weight Increased	Health Professional	Neurontin (Gabapentin)	PS	ORAL
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Depakote (Valproate Semisodium)	C
Prilosec (Omeprazole)	C
Plavix (Clopidogrel)	C
Synthroid (Levothyroxine Sodium)	C
Miacalcin Nasal Spray (Calcitonin, Salmon)	C
Feosol (Ferrous Sulfate)	C
Cardizem (Diltiazem Hydrochloride)	C
Tylenol (Paracetamol)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3414520-XReport Type:Periodic
 Age:62 YR Gender:Male I/FU:I

Company Report #001-0945-980722

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eczema	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG (400 MG, TID, PER ORAL							

Tegretol (Carbamazepine)	C		
Vitamins	C		

Date:02/26/99ISR Number: 3414521-1Report Type:Periodic
 Age:66 YR Gender:Male I/FU:I

Company Report #001-0945-980723

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Dizziness Epistaxis Feeling Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (, DAILY) PER ORAL							

Lethargy	C	Halcion (Triazolam)	
Weight Decreased	C	Viracept	
	C	Azt (Zidovudine)	
	C	Epivir (Lamivudine)	
	C	(Levocarnitine)	

Date:02/26/99ISR Number: 3414522-3Report Type:Periodic
 Age:48 YR Gender:Female I/FU:I

Company Report #001-0945-980724

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Micturition Urgency Muscle Spasms	Consumer	Neurontin (Gabapentin)	PS		ORAL
SEE TEXT, PER							

ORAL

Date:02/26/99ISR Number: 3414523-5Report Type:Periodic Company Report #001-0945-980725
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea Gingivitis	Consumer	Neurontin (Gabapentin)	PS		ORAL
3600 MG		Weight Decreased					
(,DAILY) PER							

ORAL

Dilantin (Phenytoin Sodium) C
Cardizem (Diltiazem Hydrochloride) C
Isordil (Isosorbide Dinitrate) C

Date:02/26/99ISR Number: 3414524-7Report Type:Periodic Company Report #001-0945-980726
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hallucination, Visual	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (300							
MG, TID) PER							

ORAL

(Trazodone) C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vicodin
(Paracetamol,
Hydrocodone
Bitartrate) C

Date:02/26/99ISR Number: 3414525-9Report Type:Periodic Company Report #001-0945-980727
Age:91 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID) PER ORAL		Pancytopenia	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Cardura (Doxazosin
Mesilate) C

Date:02/26/99ISR Number: 3414526-0Report Type:Periodic Company Report #001-0945-980728
Age:35 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG (100 MG, BID), PER ORAL		Constipation Fluid Retention Increased Appetite Insomnia	Consumer	Neurontin (Gabapentin)	PS		ORAL

Ultram (Tramadol
Hydrochloride) C
Carafate
(Sucralfrate) C
Paxil (Paroxetine
Hydrochloride) C

Date:02/26/99ISR Number: 3414527-2Report Type:Periodic Company Report #001-0945-980730
Age:36 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Libido Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
2700 MG (900 MG, TID), PER							
ORAL				Claritin (Loratadine)	C		
				Prilosec (Omeprazole)	C		

Date:02/26/99ISR Number: 3416247-7Report Type:Periodic Company Report #001-0945-980434
 Age:55 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tic	Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG							
(DAILY), PER							
ORAL				Zyprexa	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416251-9Report Type:Periodic
Age:40 YR Gender:Female I/FU:I

Company Report #001-0945-980435

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Swelling Weight Increased	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
400 MG							
(DAILY), PER							
ORAL							

Antivert	C
Percocet	C
Sleeping Pill	C

Date:02/26/99ISR Number: 3416261-1Report Type:Periodic
Age:43 YR Gender:Male I/FU:I

Company Report #001-0945-980436

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gingival Pain Gingivitis	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG (300							
MG, QID), PER							
ORAL							

(Methadone)	C
Valium	C

Date:02/26/99ISR Number: 3416823-1Report Type:Periodic
Age:46 YR Gender:Female I/FU:I

Company Report #001-0945-980086

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
2700 MG							
(DAILY)							

(Trazodone)	SS
Synthroid	

(Levothyroxine Sodium) C
(Insulin) C

Date:02/26/99ISR Number: 3416824-3Report Type:Periodic Company Report #001-0945-980090
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Health	Neurontin Capsules			
1800 MG		Weight Increased	Professional	300 Mg (Gabapentin)	PS		ORAL
(DAILY)				Xanax (Alprazolam)	C		
				Wellbutrin (Amfebutamone Hydrochloride)	C		

Date:02/26/99ISR Number: 3416825-5Report Type:Periodic Company Report #001-0945-980091
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Neurontin Capsules			
1800 MG (600		Feeling Abnormal		300 Mg (Gabapentin)	PS		ORAL
MG, TID)				Serzone (Nefazodone Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416826-7Report Type:Periodic
Age:55 YR Gender:Male I/FU:I

Company Report #001-0945-980092

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Glucose Increased	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
2700 MG (900 MG, TID)							

Date:02/26/99ISR Number: 3416827-9Report Type:Periodic
Age:51 YR Gender:Female I/FU:I

Company Report #001-0945-980094

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Eye Discharge Eyelid Oedema	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
300 MG, (100 MG, TID)							
		Laryngitis					
		Nasal Congestion Vision Blurred		Paxil (Paroxetine Hydrochloride)	C		
				Klonopin (Clonazepam)	C		
				Risperdal (Risperidone)	C		

Date:02/26/99ISR Number: 3416828-0Report Type:Periodic
Age:42 YR Gender:Female I/FU:I

Company Report #001-0945-980095

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Transient Ischaemic Attack	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3416829-2Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980096

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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	Bruxism	Consumer	Neurontin Capsules		
	Tongue Oedema		300 Mg (Gabapentin)	PS	ORAL
1200 MG (600					
MG, BID)					
			Artane		
			(Trihexyphenidylhydr		
			ochloride)	SS	
			Stelazine		
			(Trifluoperazine		
			Hydrochloride)	C	
			Tegretol		
			(Carbamazepine)	C	
			Elavil		
			(Amitriptyline		
			Hydrochloride)	C	
			Tranxene		
			(Clorazepate		
			Dipotassium)	C	

Date:02/26/99ISR Number: 3416830-9Report Type:Periodic Company Report #001-0945-980097
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Increased	Consumer	Neurontin (Gabapentin)	PS		
				(Seroquel)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Konopin (Clonazepam) C
 Premarin (Estrogens
 Conjugated) C
 Vitamins C

Date:02/26/99ISR Number: 3416831-0Report Type:Periodic Company Report #001-0945-980098
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Health Professional	Neurontin (Gabapentin)	PS		
600 MG							
(DAILY)							

Date:02/26/99ISR Number: 3416832-2Report Type:Periodic Company Report #001-0945-980100
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Palpitations	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1800 MG							
(DAILY)							

Date:02/26/99ISR Number: 3416833-4Report Type:Periodic Company Report #001-0945-980104
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amenorrhoea	Consumer	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3416834-6Report Type:Periodic Company Report #001-0945-980105
 Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
600 MG (300 MG, BID)							

Date:02/26/99ISR Number: 3416835-8Report Type:Periodic Company Report #001-0945-980109
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Flatulence Gastrointestinal Pain	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
300 MG (DAILY)				Percocet (Paracetamol, Oxycodone Hydrochloride, Oxycodone Ativan (Lorazepam)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416836-XReport Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-945-980110

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3600 MG (DAILY)		Myopathy	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3416837-1Report Type:Periodic
 Age:36 YR Gender:Female I/FU:I

Company Report #001-0945-980113

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG (600 MG, QID)		Flatulence	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Klonopin (Clonazepam)	C		
				Ultram (Tramadol Hydrochloride)	C		
				Seroquel (Quetiapine Fuarate)	C		
				Midrin (Paracetamol, Dichloralphenazone, Isometheptene)	C		
				Effexor (Venlafaxine Hydrochloride)	C		

Date:02/26/99ISR Number: 3416838-3Report Type:Periodic
 Age:39 YR Gender:Female I/FU:I

Company Report #001-0945-980118

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG		Alopecia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL

(DAILY)

Effexor (Venlafaxine Hydrochloride)	C
Hytrin (Terazosin Hydrochloride)	C
Propulsid (Cisapride)	C
Navane (Tiotixene)	C
Prozac (Fluoxetine Hydrochloride)	C

Date:02/26/99ISR Number: 3416839-5Report Type:Periodic

Company Report #001-0945-980122

Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG		Convulsion Drug Interaction	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
		Tooth Abscess					

(DAILY)

Biaxin (Clarithromycin)	SS
(Phenobarbital)	C
Estrogens)	C
Zantac (Ranitidine Hydrochloride)	C
Reglan	

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Freedom Of Information (FOI) Report

(Metoclopramide) C
 Colace (Docusate Sodium) C

Date:02/26/99ISR Number: 3416840-1Report Type:Periodic Company Report #001-0945-980123
 Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rosacea	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Prinivil (Lisinopril)	C		
				(Allopurinol)	C		
				Lopid (Gemfibrozil)	C		

Date:02/26/99ISR Number: 3416841-3Report Type:Periodic Company Report #001-0945-980124
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperglycaemia	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3416842-5Report Type:Periodic Company Report #001-0945-980125
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pollakiuria Urinary Incontinence	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, BID)				"Norco" Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride,	C		ORAL

Date:02/26/99ISR Number: 3416843-7Report Type:Periodic
Age:11 YR Gender:Female I/FU:I

Company Report #001-0945-980246

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Cold Muscle Twitching	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1500 MG							
(DAILY), PER							
ORAL							

Date:02/26/99ISR Number: 3416844-9Report Type:Periodic
Age:39 YR Gender:Female I/FU:I

Company Report #001-0945-980247

Outcome	PT
	Blister Burning Sensation Dysuria Erythema Eyelid Oedema Gingival Bleeding

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		Hyperhidrosis Urticaria	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
SEE TEXT, PER							
ORAL							
				Effexor (Venlafaxine Hydrochloride)	C		

Date:02/26/99ISR Number: 3416845-0Report Type:Periodic Company Report #001-0945-980248
Age:27 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Pityriasis Rosea	Health Professional	Neurontin Capsules 300 Mg(Gabapentin)	PS		ORAL
Dose							
1200 MG (300							
MG, QID), PER							
ORAL							
				Motrin (Ibuprofen)	C		

Date:02/26/99ISR Number: 3416846-2Report Type:Periodic Company Report #001-0945-980872
Age:40 YR Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Suicidal Ideation	Health Professional	Neurontin (Gabapentin)	PS		
Dose							

Date:02/26/99ISR Number: 3416847-4Report Type:Periodic Company Report #001-0945-980015
Age:50 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration	Hypoaesthesia	Consumer	Neurontin Capsules			
Dose							

3600 MG DAILY

Nervousness

400 Mg (Gabapentin)

PS

ORAL

Oedema Peripheral

PER ORAL

Pain In Extremity

Speech Disorder

Vision Blurred

Weight Increased

Prevacid

(Lansoprazole)

C

Dilantin (Phenytoin

Sodium)

C

(Trazodone)

C

Carafate

(Sucralfate)

C

Propulsid

(Cisapride)

C

Questran

(Colestyramine)

C

Imodium (Loperamide

Hydrochloride)

C

Pancreatic Enzyme

(Unspecified)

C

Estrogens (Estrogen

Nos)

C

Date:02/26/99ISR Number: 3416848-6Report Type:Periodic

Company Report #001-0945-980249

Age:45 YR Gender:Female I/FU:I

Outcome

PT

Report Source

Tremor

Health

Professional

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Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL			Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
			Aerobid (Flunisolide)	C		
			Serevent (Salmeterol Xinafoate)	C		
			Amitriptyline	C		
			Proamatine (Midodrine Hydrochloride)	C		

Date:02/26/99ISR Number: 3416849-8Report Type:Periodic
Age:28 YR Gender:Male I/FU:I

Company Report #001-0945-980027

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (300 MG, QID) PER ORAL		Abdominal Pain Upper Anxiety	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
		Eye Disorder					
		Flushing					
		Headache Hyperhidrosis		Mebaral (Methylphenobarbital) (Allopurinol)	C C		

Date:02/26/99ISR Number: 3416850-4Report Type:Periodic
Age:65 YR Gender:Male I/FU:I

Company Report #001-0945-980250

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Clonic Convulsion	Health	Neurontin Capsules			

300 MG	Sedation	Professional	300 Mg(Gabapentin)	PS	ORAL
(DAILY), PER	Tremor	Company			
ORAL		Representative			

Alprazolam	SS
Diazepam	SS
Fentanyl	C
Prilosec	
(Omeprazole)	C
Humulin Nph (Insulin Human Injection, Isophane)	C
Morphine Sulfate	C
Thiamin (Thiamine Hydrochloride)	C
Human Regular (Insulin Human)	C
Cozaar (Losartan Potassium)	C
Coumadin (Warfarin Sodium)	C
Cefalexin	C
Toprol Xl (Metoprolol Succinate)	C
Aspirin	

FDA - Adverse Event Reporting System (AERS)

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(Acetylsalicylic Acid)	C
Trental	
(Pentoxifylline)	C
Darvocet	
(Paracetamol, Dextropropoxyphene)	C
Xanax (Alprazolam)	C
Dulcolax (Bisacodyl)	C
Nitro (Glyceryl Trinitrate)	C
Propulsid (Cisapride)	C
Nephron Fa	C

Date:02/26/99ISR Number: 3416851-6Report Type:Periodic
Age:13 YR Gender:Male I/FU:I

Company Report #001-0945-980028

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pityriasis Rosea		Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
1600 MG (800 MG BID) PER ORAL							

Date:02/26/99ISR Number: 3416852-8Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-980251

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Withdrawal Syndrome Fatigue	Health Professional	Neurontin(Gabapentin)	PS		
6 MON		Influenza Like Illness Myalgia					

Date:02/26/99ISR Number: 3416853-XReport Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980030

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blister Confusional State Gait Disturbance Vision Blurred	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		

Date:02/26/99ISR Number: 3416854-1Report Type:Periodic Company Report #001-0945-980253
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia Chest Pain	Health Professional Company Representative	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL				Coumadin (Warfarin Sodium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416855-3Report Type:Periodic
Age:81 YR Gender:Female I/FU:I

Company Report #001-0945-980032

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Neurontin Capsules	PS		ORAL
600 MG (300		Embolism		300 Mg (Gabapentin)			
MG, BID) PER		Weight Increased					
ORAL							

Couadin (Warfarin Sodium) C

Date:02/26/99ISR Number: 3416856-5Report Type:Periodic
Age:63 YR Gender:Female I/FU:I

Company Report #001-0945-980257

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hangover	Consumer	Neurontin	PS		ORAL
300 MG (100		Sedation	Health	(Gabapentin0			
MG, TID), PER			Professional				
ORAL							

Buspar (Buspirone Hydrochloride) C
Humulin Nph (Insulin Human Injection, Isophane) C
Humulin Reg. (Insulin Human) C
Moduretic (Hydrochlorothiazide, Amiloride Hydrochloride) C

Date:02/26/99ISR Number: 3416857-7Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #001-0945-980033

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia Hair Growth Abnormal	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
300 MG (100 MG TID) PER ORAL		Hair Texture Abnormal					

Sinemet (Levodopa,
Carbidopa) C
Tegretol
(Carbamazepine) C

Date:02/26/99ISR Number: 3416858-9Report Type:Periodic Company Report #001-0945-980258
Age:51 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pain	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL							

Tylenol W/Codeine
No. 3 (Codeine
Phosphate,
Paracetamol) C
Lescol (Fluvastatin
Sodium) C
Micronase

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Glibenclamide) C
 Potassium Chloride C

Date:02/26/99ISR Number: 3416859-0Report Type:Periodic
 Age:36 YR Gender:Female I/FU:I

Company Report #001-0945-980034

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
SEE TEXT, PER		Diplopia Keratoconjunctivitis	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
ORAL		Sicca		(Phentermine)	C		
		Photophobia Vision Blurred		Accutane (Isotretinoin)	C		
				Soma (Carisoprodol)	C		
				Percocet (Paracetamol, Oxycodone Hydrochloride, Oxycodone	C		

Date:02/26/99ISR Number: 3416860-7Report Type:Periodic
 Age:53 YR Gender:Male I/FU:I

Company Report #001-0945-980259

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG (100 MG, BID), PER		Diplopia	Consumer Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
ORAL				Insulin	C		
				Oxycontin (Oxycodone Hydrochloride)	C		

Date:02/26/99ISR Number: 3416861-9Report Type:Periodic
 Age:68 YR Gender:Female I/FU:I

Company Report #001-0945-980035

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG TID) PER ORAL		Back Pain	Consumer	Neurontin Capsules	PS		ORAL
		Blood Glucose Increased		300 Mg (Gabapentin)			
		Diarrhoea					
		Fatigue					
		Weight Increased		Maalox (Magnesium Hydroxide, Aluminum Hydroxide Gel)	C		
				Rezulin (Troglitazone)			
				(Troglitazone)	C		
				Humulin R (Insulin Human)	C		
				Humulin Nph (Insulin Human Injection, Isophane)	C		
				Glucophage (Metformin Hydrochloride)	C		
				(Furosemide)	C		
				Cycrin (Medroxyprogesterone Acetate)	C		
				Premarin (Estrogens			

Alopecia
 Health Professional
 Neurontin (Gabapentin)
 PS
 ORAL
 900 MG (300 MG, TID) ,
 PER ORAL

Dyazide
 (Hydrochlorothiazide , Triamterene) C
 Hydrochlorothiazide C
 Premarin (Estrogens Conjugated) C
 Synthroid (Levothyroxine Sodium) C

Date:02/26/99ISR Number: 3416865-6Report Type:Periodic Company Report #001-0945-980263
 Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Psychotic Disorder	Health Professional	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416866-8Report Type:Periodic
Age:60 YR Gender:Female I/FU:I

Company Report #001-0945-980039

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anosmia	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		
1200 MG (400 MG TID)			Company Representative				

Date:02/26/99ISR Number: 3416867-XReport Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980264

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Engorgement Breast Tenderness	Health Professional	Neurontin (Gabapentin)	PS		
800 MG (400 MG, BID), UNKNOWN	4 WK		Company Representative				

Date:02/26/99ISR Number: 3416868-1Report Type:Periodic
Age:70 YR Gender:Female I/FU:I

Company Report #001-0945-980040

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Increased Sedation	Consumer	Neurontin (Gabapentin)	PS		ORAL
400 MG (QHS) PER ORAL		Stupor		Insulin (Insulin) Vasotec (Enalapril Maleate) Norvasc (Amlodipine Besilate) Lasix (Furosemide) Epogen (Epoetin Alfa)	C C C C C		

Peri-Colace
 (Docusate Sodium,
 Casanthranol) C
 (Sorbitol) C
 (Calcium) C
 (Iron) C
 Multivitamins
 (Ergocalciferol,
 Ascorbic Acid, Folic
 Acid, Thiamine
 Hydrochloride, C

Date:02/26/99ISR Number: 3416869-3Report Type:Periodic Company Report #001-0945-980265
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Spasms	Health Professional	Neurontin (Gabapentin)	PS		
900 MG							
(DAILY),							
UNKNOWN							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416870-XReport Type:Periodic
Age:48 YR Gender:Female I/FU:I

Company Report #001-0945-980041

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bladder Pain Condition Aggravated	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
PER ORAL				Surmontil (Trimipramine)	C		
				Urispas (Flavoxate Hydrochloride)	C		
				Tylenol (Paracetamol)	C		
				Ativan (Lorazepam)	C		
				Claritin (Loratadine)	C		
				Estraderm (Estradiol)	C		

Date:02/26/99ISR Number: 3416871-1Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-980266

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased Palpitations	Health Professional	Neurontin (Gabapentin)	PS		
SEE TEXT			Company Representative				

Date:02/26/99ISR Number: 3416872-3Report Type:Periodic
Age:36 YR Gender:Female I/FU:I

Company Report #001-0945-980267

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Muscle Spasms	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
200 MG (AT HS), PER ORAL			Professional				

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache Insomnia	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
600 MG (300 MG BID) PER ORAL		Nausea Sedation		Capoten (Captopril) Lanoxin (Digoxin) (Triazolam) Soma (Carisoprodol) Vicodin (Paracetamol, Hydrocodone Bitartrate) Trental (Pentoxifylline)	C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416874-7Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-980268

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Pressure Increased Palpitations	Health Professional Company Representative	Neurontin(Gabapentin)	PS		

SEE TEXT

Date:02/26/99ISR Number: 3416875-9Report Type:Periodic
Age:27 YR Gender:Male I/FU:I

Company Report #001-0945-980047

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG TID) PER ORAL		Eye Movement Disorder Memory Impairment Nystagmus	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Wellbutrin (Amfebutamone Hydrochloride)	C		

Date:02/26/99ISR Number: 3416876-0Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-980269

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Pressure Increased Palpitations	Health Professional Company Representative	Neurontin(Gabapentin)	PS		

SEE TEXT

Date:02/26/99ISR Number: 3416877-2Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-980270

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased Palpitations	Health Professional	Neurontin (Gabapentin)	PS		
SEE TEXT			Company Representative				

Date:02/26/99ISR Number: 3416878-4Report Type:Periodic Company Report #001-0945-980048
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG TID) PER ORAL				Wellbutrin (Amfebutamone Hydrochloride)	C		
				Soma (Carisoprodol) (Lorazepam)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416879-6Report Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980049

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Decreased Drug Withdrawal Syndrome	Health Professional	Neurontin (Gabapentin)	PS		
1800 MG (600 MG TID)		Flushing Headache Malaise					

Date:02/26/99ISR Number: 3416880-2Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980051

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urinary Incontinence	Consumer	Neurontin (Gabapentin)	PS		
200 MG DAILY							

Date:02/26/99ISR Number: 3416881-4Report Type:Periodic
Age:47 YR Gender:Female I/FU:I

Company Report #001-0945-980054

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1800 MG (600 MG TID) PER ORAL ; 200 MG (100 MG BID) PER ORAL				(Morphine)	C		

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG TID) PER ORAL		Asthenia Sedation	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Lanoxin (Digoxin)	C		
				Rythmol (Propafenone)	C		
				Norvasc (Amlodipine Besilate)	C		
				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride,	C		

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Urinary Tract Infection	Consumer Health Professional Company Representative	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416884-XReport Type:Periodic
Age:51 YR Gender:Female I/FU:I

Company Report #001-0945-980058

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		Breast Engorgement Breast Tenderness Night Sweats	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Estrostep (Ethinyl-Estradiol, Norethisterone) Dyazide (Hydrochlorothiazide , Triamterene) Voltaren (Diclofenac Sodium)	C C C		

Date:02/26/99ISR Number: 3416885-1Report Type:Periodic
Age:94 YR Gender:Female I/FU:I

Company Report #001-0945-980059

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error Sleep Disorder	Consumer	Neurontin (Gabapentin) Fioricet (Caffeine, Butalbital, Paracetamol) Xanax (Alprazolam) Klonopin (Clonazepam)	PS C C C		

Date:02/26/99ISR Number: 3416886-3Report Type:Periodic
Age:30 YR Gender:Female I/FU:I

Company Report #001-0945-980060

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300		Alopecia	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

MG TID), PER

ORAL

Remeron	
(Mirtazapine)	C
(Trazodone)	C
Ativan (Lorazepam)	C
(Prednisone0	C
Florinef	
(Fludrocortisone	
Acetate)	C
Dexedrine	
(Dexamfetamine	
Sulfate)	C
Depakote (Valproate	
Semisodium)	C

Date:02/26/99ISR Number: 3416887-5Report Type:Periodic
Age:28 YR Gender:Female I/FU:I

Company Report #001-0945-980061

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cutaneous Lupus Erythematosus	Health Professional	Neurontin (Gabapentin) Tegretol	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Carbamazepine) C

Date:02/26/99ISR Number: 3416888-7Report Type:Periodic
Age:60 YR Gender:Female I/FU:I

Company Report #001-0945-980062

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (, DAILY), PER ORAL		Increased Appetite Weight Increased	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Klonopin (Clonazepam)	C		
				Synthroid (Levothyroxine Sodium)	C		

Date:02/26/99ISR Number: 3416889-9Report Type:Periodic
Age:35 YR Gender:Female I/FU:I

Company Report #001-0945-980066

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, BID), PER ORAL		Medication Error Periodontal Disease Sleep Walking	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				(Amitriptyline)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Capoten (Captopril)	C		
				Macrochantin (Nitrofurantoin)	C		
				Pravachol (Pravastatin Sodium)	C		
				Premarin (Estrogens Conjugated)	C		
				Lasix (Furosemide)	C		

Slow-Mag (Magnesium
Chloride Anhydrous) C
Insulin (Insulin) C

Date:02/26/99ISR Number: 3416890-5Report Type:Periodic
Age:50 YR Gender:Male I/FU:I

Company Report #001-0945-980067

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Toothache	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL				Pamelor (Nortriptyline Hydrochloride)	C		
				Ms Contin (Morphine Sulfate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416891-7Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980126

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction International Normalised Ratio Decreased	Consumer	Neurontin (Gabapentin) Coumadin (Warfarin Sodium)	PS SS		

Date:02/26/99ISR Number: 3416892-9Report Type:Periodic
Age:54 YR Gender:Female I/FU:I

Company Report #001-0945-980068

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia Sedation	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL				Calan (Verapamil Hydrochloride) Synthroid (Levothyroxine Sodium) Zoloft (Sertraline Hydrochloride) Estrace (Estradiol) Vicodin (Paracetamol, Hydrocodone Bitartrate) Elavil (Amitriptyline Hydrochloride) (Carisoprodol)	C C C C C C C		

Date:02/26/99ISR Number: 3416893-0Report Type:Periodic
Age:20 YR Gender:Female I/FU:I

Company Report #001-0945-980070

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Neurontin			
		Hair Growth Abnormal		(Gabapentin)	PS		

Date:02/26/99ISR Number: 3416894-2Report Type:Periodic Company Report #001-0945-980129
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menorrhagia	Health Professional	Neurontin Capsules			
1200 MG				100 Mg (Gabapentin)	PS		ORAL
(DAILY), PER							
ORAL				Asthma Meds	C		

Date:02/26/99ISR Number: 3416895-4Report Type:Periodic Company Report #001-0945-980130
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Consumer	Neurontin			
600 - 1200 MG		Convulsion		(Gabapentin)	PS		ORAL
(DAILY), PER		Malaise					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Tegretol
 (Carbamazepine) C
 Topamax (Topiramate) C
 Ritalin
 (Methylphenidate
 Hydrochloride) C
 Decadron
 (Dexamethasone) C

Date:02/26/99ISR Number: 3416896-6Report Type:Periodic Company Report #001-0945-980071
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, TID), PER		Hypertension Influenza	Health Professional	Neurontin (Gabapentin)	PS		ORAL

ORAL

Olanzapine
 (Olanzapine) C

Date:02/26/99ISR Number: 3416897-8Report Type:Periodic Company Report #001-0945-980073
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Bone Marrow Depression	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3416898-XReport Type:Periodic Company Report #001-0945-980075
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300		Difficulty In Walking Feeling Hot	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

Paralysis

MG, TID), PER

ORAL

Date:02/26/99ISR Number: 3416899-1Report Type:Periodic

Company Report #001-0945-980135

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Neurontin			
4200 MG		Chest Pain		(Gabapentin)	PS		ORAL
(DAILY), PER		Confusional State					
ORAL		Dermatitis					
		Disturbance In Attention		Tylenol With Codeine			
		Dizziness		No. 3 (Codeine			
		Eye Movement Disorder		Phosphate			
		Fatigue		Paracetamol)	C		
		Nervousness					
		Vision Blurred					

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Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416900-5Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980077

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID)		Menstruation Irregular	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		
				(Oral Contraceptive Nos)	SS		
				Duract	SS		

Date:02/26/99ISR Number: 3416901-7Report Type:Periodic
 Age:46 YR Gender:Female I/FU:I

Company Report #001-0945-980078

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG (100 MG, BID), PER ORAL		Alopecia	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
				Zoloft (Sertraline Hydrochloride)	C		
				Prilosec (Omeprazole)	C		
				Ogen (Estropipate)	C		

Date:02/26/99ISR Number: 3416902-9Report Type:Periodic
 Age:62 YR Gender:Female I/FU:I

Company Report #001-0945-980144

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2600 MG (DAILY), PER		Tinnitus	Consumer	Neurontin (Gabapentin)	PS		ORAL

ORAL

Inderal (Propranolol	
Hydrochloride)	C
(Codeine)	C
Tranxene	
(Clorazepate	
Dipotassium)	C
Miacalcin	
(Calcitonin, Salmon)	C

Date:02/26/99ISR Number: 3416903-0Report Type:Periodic
 Age:45 YR Gender:Male I/FU:I

Company Report #001-0945-980145

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
900 MG (300							
MG, TID), PER							

ORAL

Rythmol	
(Propafenone)	C
Baby Aspirin	
(Acetylsalicylic	
Acid)	C
St. John'S Wort	
(Hypericum Extract)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416904-2Report Type:Periodic
Age:53 YR Gender:Male I/FU:I

Company Report #001-0945-980079

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Health	Neurontin Capsules			
		Gingivitis	Professional	300 Mg (Gabapentin)	PS		ORAL
2400 MG (600		Hyperhidrosis					
MG, QID), PER		Myalgia					
ORAL				(Trazodone)	SS		
				(Insulin)	C		
				(Atenolol)	C		
				Vicodin			
				(Paracetamol,			
				Hydrocodone			
				Bitartrate(C		
				Posicor (Mibefradil)	C		
				Valium (Diazepam)	C		

Date:02/26/99ISR Number: 3416905-4Report Type:Periodic
Age:43 YR Gender:Female I/FU:I

Company Report #001-0945-980080

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Health	Neurontin Capsules			
			Professional	300 Mg (Gabapentin)	PS		
900 MG (300							
MG, TID)				(Insulin)	C		
				(Prednisone)	C		
				Cyclosporin			
				(Ciclosporin)	C		
				Imuran			
				(Azathioprine)	C		
				Lasix (Furosemide)	C		
				Normodyne (Labetalol			
				Hydrochloride)	C		
				Cardene (Nicardipine			
				Hydrochloride)	C		
				Coumadin (Warfarin			
				Sodium)	C		

Provera
(Medroxyprogesterone
Acetate) C

Date:02/26/99ISR Number: 3416906-6Report Type:Periodic Company Report #001-0945-980146
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia Dizziness	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL		Fatigue					

Xanax (Alprazolam) C

Date:02/26/99ISR Number: 3416907-8Report Type:Periodic Company Report #001-0945-980082
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Influenza Like Illness Psychotic Disorder	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		
900 MG (300 MG, TID)							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dilantin (Phenytoin Sodium) C
 Haldol (Haloperidol) C
 Cogentin (Benzatropine Mesilate) C
 Antibiotics (Unspecified) C

Date:02/26/99ISR Number: 3416908-XReport Type:Periodic Company Report #001-0945-980147
 Age: Gender: I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Influenza Like Illness	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3416909-1Report Type:Periodic Company Report #001-0945-980084
 Age:41 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG (, DAILY), PER ORAL		Disturbance In Attention Memory Impairment	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Dilantin (Phenytoin Sodium)	C		

Date:02/26/99ISR Number: 3416910-8Report Type:Periodic Company Report #001-0945-980152
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Agitation	Consumer	Neurontin			

PER ORAL	Anxiety	(Gabapentin)	PS	ORAL
	Chest Pain	Effexor (Venlafaxine		
	Feeling Jittery	Hydrochloride)	C	
		(Imipramine)	C	

Date:02/26/99ISR Number: 3416911-XReport Type:Periodic Company Report #001-0945-980153
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Glossodynia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
2700 MG, (900							
MG,TID), PER							
ORAL							

Paxil (Paroxetine Hydrochloride)	C
Synthroid (Levothyroxine Sodium)	C
(Thioridazine)	C
Humibid (Guaifenesin)	C
Decongestant (Unspecified)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416912-1Report Type:Periodic
Age:58 YR Gender:Female I/FU:I

Company Report #001-0945-980085

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG (, DAILY)		Diplopia Eye Disorder Eye Movement Disorder Eye Pain Eyelid Oedema Facial Pain Lacrimation Increased Pain Paraesthesia	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		
				Zoloft (Sertraline Hydrochloride) Indocin (Indometacin) (Verapamil) Nitro Patch (Glyceryl Trinitrate) Thyroid Medication	C C C C C		

Date:02/26/99ISR Number: 3416913-3Report Type:Periodic
Age:46 YR Gender:Male I/FU:I

Company Report #001-0945-980272

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3200 MG (800 MG, QID), PER ORAL		Ventricular Extrasystoles	Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
				Zoloft (Sertraline Hydrochloride) Buspar (Buspirone Hydrochloride)	C C		

Date:02/26/99ISR Number: 3416914-5Report Type:Periodic
Age:39 YR Gender:Female I/FU:I

Company Report #001-0945-980274

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness	Consumer	Neurontin Capsules			

1200 MG (400
 MG, TID), PER
 ORAL

Allegra(Fexofenadine
 Hydrochloride) C
 Imitrex
 (Sumatriptan) C

Date:02/26/99ISR Number: 3416915-7Report Type:Periodic Company Report #001-0945-980276
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG		Dyspnoea Muscle Contractions	Health Professional	Neurontin (Gabapentin)	PS		ORAL
(,DAILY), PER		Involuntary					

ORAL

Premarin(Estrogens
 Conjugated) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416916-9Report Type:Periodic
Age:40 YR Gender:Male I/FU:I

Company Report #001-0945-980279

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety Testicular Pain	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1800 MG							
(,DAILY), PER							
ORAL							

Zoloft(Sertraline
Hydrochloride) C
Klonopin(Clonazepam) C

Date:02/26/99ISR Number: 3416917-0Report Type:Periodic
Age:48 YR Gender:Female I/FU:I

Company Report #001-0945-980280

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dysphagia Pharyngeal Oedema	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1800 MG (300							
MG, SIX TIMES							
DAILY), PER							
ORAL							

Premarin (Estrogens
Conjugated) C
Multivitamins(Ergoca
lciferol, Ascorbic
Acid, Folic Acid,
Thiamine
Hydrochloride, C
Oscal (Calcium
Carbonate) C

Date:02/26/99ISR Number: 3416918-2Report Type:Periodic
Age:39 YR Gender:Female I/FU:I

Company Report #001-0945-980155

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alopecia Hirsutism	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1800 MG (300 MG, X6), PER ORAL							

(Baclofen) C
 Synthroid
 (Levothyroxine
 Sodium) C
 Lasix (Furosemide) C
 (Methadone) C
 Prozac (Fluoxetine
 Hydrochloride) C

Date:02/26/99ISR Number: 3416919-4Report Type:Periodic Company Report #001-0945-980156
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Amylase Increased Lipase Increased	Health Professional	Neurontin (Gabapentin)	PS		
14 DAY				(Lorazepam) Dilantin (Phenytoin Sodium) Phenobarb (Phenobarbital	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sodium) C
(Pyridoxine) C

Date:02/26/99ISR Number: 3416920-0Report Type:Periodic
Age:55 YR Gender:Male I/FU:I

Company Report #001-0945-980282

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
400 MG (100 MG, QID), PER ORAL				Glucotrol (Glipizide)	C		
				Lipitor (Atorvastatin)	C		
				Unspecified Blood Pressure Pill	C		

Date:02/26/99ISR Number: 3416921-2Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980168

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness Hemiparesis	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
1200 MG (400 MG, TID), PER ORAL				Lortab (Paracetamol, Hydrocodone Bitartrate)	C		
				Parafon Forte (Chloroxazone, Paracetamol)	C		
				Dilacor (Digoxin)	C		

Lortab (Paracetamol,
Hydrocodone
Bitartrate) C
Parafon Forte
(Chloroxazone,
Paracetamol) C
Dilacor (Digoxin) C

Date:02/26/99ISR Number: 3416922-4Report Type:Periodic Company Report #001-0945-980284
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
PER ORAL							

Date:02/26/99ISR Number: 3416923-6Report Type:Periodic Company Report #001-0945-980169
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL							
				Rhinocort (Budesonide)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416924-8Report Type:Periodic
Age:78 YR Gender:Male I/FU:I

Company Report #001-0945-980287

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper Vomiting	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG							
(,DAILY), PER							
ORAL							

Lanoxin (Digoxin)	C
Capoten (Captopril)	C
Vitamin C (Ascorbic Acid)	C
Ativan (Lorazepam)	C
Vicodin (Paracetamol, Hydrocodone Bitartrate)	C
(Zinc)	C
(Folic Acid)	C

Date:02/26/99ISR Number: 3416925-XReport Type:Periodic
Age:44 YR Gender:Female I/FU:I

Company Report #001-0945-980171

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Balance Disorder Blister	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600							
MG, TID) PER		Dermatitis					
ORAL		Dry Mouth					
		Ecchymosis		Dilantin (Phenytoin Sodium)	C		
		Headache		Valium (Diazepam)	C		
		Nausea		Fioricet (Caffeine, Butalbital, Paracetamol)	C		
		Pruritus					
		Rash Papular					
		Sedation					
		Skin Ulcer					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nasal Congestion	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
3600 MG							
(,DAILY), PER							
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disorientation Feeling Jittery	Consumer	Neurontin (Gabapentin)	PS		ORAL
THREE		Headache					
CAPSULES (1		Heart Rate Increased					
CAPSULE,		Mental Impairment					
TID), PER		Mood Altered					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416928-5Report Type:Periodic
 Age:66 YR Gender:Female I/FU:I

Company Report #001-0945-980289

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG		Discomfort Oedema	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
(,DAILY), PER ORAL							

Klonopin (Clonazepam)	C
(Estradiol)	C
Baby Asa (Acetylsalicylic Acid)	C

Date:02/26/99ISR Number: 3416929-7Report Type:Periodic
 Age:60 YR Gender:Female I/FU:I

Company Report #001-0945-980177

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID), PER		Bronchospasm Dyspnoea	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
ORAL							

Ativan (Lorazepam)	C
(Prednisone)	C
Calan (Verapamil Hydrochloride)	C
(Skullcap)	C
L-Taurene	C

Date:02/26/99ISR Number: 3416930-3Report Type:Periodic
 Age:24 YR Gender:Female I/FU:I

Company Report #001-0945-980180

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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900 MG (300	900 MG (300	Consumer	Neurontin Capsules	300 Mg (Gabapentin)	PS	ORAL
MG, TID), PER	MG, TID), PER	Hearing Impaired				
ORAL	ORAL	Photopsia				
		Vision Blurred				

Neupogen (Filgrastim)	C
Dextrostat (Dexamfetamine Sulfate)	C

Date:02/26/99ISR Number: 3416931-5Report Type:Periodic Company Report #001-0945-980864
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3416932-7Report Type:Periodic Company Report #001-0945-980181
 Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction Drug Level Above	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300		Therapeutic					
MG, TID), PER							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL/ 600 MG
 (300 MG,
 DAILY), PER
 900 MG (300
 MG, TID), PER
 ORAL

(Lithium) SS ORAL

Depakote (Valproate
 Semisodium) C
 Synthroid
 (Levothyroxine
 Sodium) C

Date:02/26/99ISR Number: 3416933-9Report Type:Periodic Company Report #001-0945-980865
 Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (, ONE DOSE), PER ORAL				Neurontin (Gabapentin)	SS		ORAL
100 MG (, ONE DOSE), PER ORAL				Dyazide (Hydrochlorothiazide , Triamterene) Mellaril (Thioridazine Hydrochloride) Antivert (Nicotinic Acid, Meclozine Hydrochloride)	C C C		

Date:02/26/99ISR Number: 3416934-0Report Type:Periodic
 Age:61 YR Gender:Female I/FU:I

Company Report #001-0945-980182

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1100 MG (DAILY)		Hallucination Hostility	Health Professional	Neurontin (Gabapentin)	PS		
				Elavil (Amitriptyline Hydrochloride)	C		
				Prilosec (Omeprazole)	C		
				Propulsid (Cisapride)	C		
				Peri-Colace (Docusate Sodium, Casanthranol)	C		
				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride,	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416935-2Report Type:Periodic
Age:47 YR Gender:Female I/FU:I

Company Report #001-0945-980866

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
300 MG (,							
QHS), PER							
ORAL							

Elavil (Amitriptyline Hydrochloride)	C
(Salsalate)	C
Ativan (Lorazepam)	C
Lortab (Paracetamol, Hydrocodone Bitartrate)	C
Ogen (Estropipate)	C
Tylenol (Paracetamol)	C

Date:02/26/99ISR Number: 3416936-4Report Type:Periodic
Age:57 YR Gender:Female I/FU:I

Company Report #001-0945-980867

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine Multiple Sclerosis	Consumer Health	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300			Professional				
MG, TID), PER							

Imitrex (Sumatriptan)	C
Fioricet (Caffeine, Butalbital, Paracetamol)	C

ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Coordination Abnormal White Blood Cell Count Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG (100 MG, BID), PER ORAL				(Vitamins Nos)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia Haematuria Joint Swelling Polyuria	Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG (,DAILY), PER ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416939-XReport Type:Periodic
 Age:49 YR Gender:Male I/FU:I

Company Report #001-0945-980873

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (400 MG, TID), PER ORAL		Abdominal Distension Vomiting	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
				Depakote (Valproate Semisodium)	C		

Date:02/26/99ISR Number: 3416940-6Report Type:Periodic
 Age:13 YR Gender:Female I/FU:I

Company Report #001-0945-980875

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID), PER ORAL		Epistaxis	Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
				(Lithium)	C		

Date:02/26/99ISR Number: 3416941-8Report Type:Periodic
 Age:44 YR Gender:Female I/FU:I

Company Report #001-0945-980876

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (,DAILY), PER ORAL		Dizziness Dysarthria Gait Disturbance Sedation	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Flomax			

0.4 MG	(Morniflumate)	SS	ORAL
(,DAILY), PER			
ORAL			
1 MG	Ativan (Lorazepam)	SS	ORAL
(,DAILY), PER			
ORAL			
100 MG	Elavil (Amitriptyline Hydrochloride)	SS	ORAL
(,DAILY), PER			
ORAL			

Date:02/26/99ISR Number: 3416942-XReport Type:Periodic Company Report #001-0945-980877
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Neurontin Capsules			
		Drug Interaction		300 Mg (Gabapentin)	PS		ORAL
1200 MG (300		Libido Decreased					
MG, QID), PER		Nausea					
ORAL				Imitrex (Sumatriptan Succinate)	SS		
INTRAMUSCULAR	WHEN NEEDED,						
INTRAMUSCULAR				Hyzaar (Hydrochlorothiazide , Losartan Potassium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416943-1Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #001-0945-980903

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Coordination Abnormal Lethargy	Consumer	Neurontin (Gabapentin)	PS		
400 MG (,DAILY),		Tremor		Nardil (Phenelzine Sulfate) (Lithium)	C C		

Date:02/26/99ISR Number: 3416944-3Report Type:Periodic
 Age:32 YR Gender:Female I/FU:I

Company Report #001-0945-980905

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hair Growth Abnormal Medication Error	Consumer	Neurontin 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL				Ultram (Tramadol Hydrochloride) Darvocet N (Paracetamol Dextropropoxyphene)	C C		

Date:02/26/99ISR Number: 3416945-5Report Type:Periodic
 Age:53 YR Gender:Male I/FU:I

Company Report #001-0945-980908

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral	Health Professional	Neurontin (Gabapentin)	PS		
3600 MG (,DAILY),							

SEVERAL

MONTHS AGO -

Date:02/26/99ISR Number: 3416946-7Report Type:Periodic Company Report #001-0945-980909
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue Medication Error	Consumer	Neurontin Capsules 100 Mg(Gabapentin)	PS		ORAL
900 MG (300 MG,TID), PER ORAL				Klonopin (Clonazepam)	C		
				Remeron (Mirtazapine)	C		

Date:02/26/99ISR Number: 3416947-9Report Type:Periodic Company Report #001-0945-980910
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error Myalgia	Consumer	Neurontin Capsules 100 Mg(Gabapentin)	PS		ORAL
900 MG (300 MG, TID,) PER ORAL				Remeron (Mirtazapine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Klonopin
(Clonazepam) C

Date:02/26/99ISR Number: 3416948-0Report Type:Periodic Company Report #001-0945-980911
Age:29 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID, PER ORAL)		Chest Pain Fatigue Influenza Like Illness Myalgia	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3416949-2Report Type:Periodic Company Report #001-0945-980913
Age: Gender:Unknown I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tongue Geographic	Health Professional	Neurontin (Gabapentin) Premarin (Estrogens Conjugated) Provera (Medroxyprogesterone Acetate) Synthroid (Levothyroxine Sodium)	PS C C C		

Date:02/26/99ISR Number: 3416951-0Report Type:Periodic Company Report #001-0945-980916
Age:87 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG (, ONE		Asthenia Headache	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

Palpitations

DOSE), PER

ORAL

Date:02/26/99ISR Number: 3416952-2Report Type:Periodic Company Report #001-0945-980917

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Joint Stiffness	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
300 MG, PER				(Nortriptyline)	C		

Dose

Duration

PT

Report Source

Product

Role

Manufacturer

Route

Joint Stiffness

Consumer

Neurontin Capsules
300 Mg (Gabapentin)

PS

ORAL

300 MG, PER

ORAL

(Nortriptyline)

C

Date:02/26/99ISR Number: 3416954-6Report Type:Periodic Company Report #001-0945-980918

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia	Consumer	Neurontin (Gabapentin)	PS		ORAL
800 MG (400				Lamictal (Lamotrigine)	C		

Dose

Duration

PT

Report Source

Product

Role

Manufacturer

Route

Alopecia

Consumer

Neurontin
(Gabapentin)

PS

ORAL

800 MG (400

MG, BID), PER

ORAL

Lamictal
(Lamotrigine)

C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416956-XReport Type:Periodic
Age:78 YR Gender:Female I/FU:I

Company Report #001-0945-990005

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypervigilance	Consumer	Neurontin Capsules 300 Mg(Gabapentin)	PS		ORAL
300 MG, (,QHS), PER ORAL							

Date:02/26/99ISR Number: 3416957-1Report Type:Periodic
Age:72 YR Gender:Female I/FU:I

Company Report #001-0945-990006

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypervigilance	Consumer	Neurontin Capsules 300 Mg(Gabapentin)	PS		ORAL
900 MG (,QHS), PER ORAL							

Synthroid
(Levothyroxine
Sodium) C
Macrochantin
(Nitrofurantoin) C
Zocor (Simvastatin) C
Maxzide
(Hydrochlorothiazide
, Triamterene) C

Date:02/26/99ISR Number: 3416959-5Report Type:Periodic
Age:39 YR Gender:Female I/FU:I

Company Report #001-0945-990007

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia Joint Swelling	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
2800 MG (,							

800 MG TID,
400 MGQHS),
PER ORAL

Triphasil
(Ethinylestradiol,
Levonorgestrel) C
Effexor (Venlafaxine
Hydrochloride) C

Date:02/26/99ISR Number: 3416961-3Report Type:Periodic Company Report #001-0945-990009
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Decreased Sedation	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG							

(,QHS), PER
ORAL

Synthroid
(Levothyroxine
Sodium) C
Klonopin
(Clonazepam) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3416967-4Report Type:Periodic
 Age:44 YR Gender:Female I/FU:I

Company Report #001-0945-990010

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3600 MG (,1200 MG QAM, 2400 MG QHS), PER ORAL		Fatigue	Consumer	Neurontin (Gabapentin)	PS		ORAL

Synthroid
(Levothyroxine Sodium) C
 Klonopin
(Clonazepam) C
 Paxil (Paroxetine Hydrochloride) C

Date:02/26/99ISR Number: 3416968-6Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-990016

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (,DAILY),		Haematochezia Visual Disturbance Vomiting	Consumer	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3416970-4Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-990022

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2700 MG		Abdominal Distension Hyponatraemic Syndrome	Consumer	Neurontin (Gabapentin)	PS		

(,DAILY),

Calan St (Verapamil Hydrochloride)	C
Soma (Carisoprodol)	C
Lasix (Furosemide)	C
Estraderm (Estradiol)	C
Prevacid (Lansoprazole)	C
Naprosyn (Naproxen Sodium)	C

Date:02/26/99ISR Number: 3416973-XReport Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-990092

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Health Professional	Neurontin (Gabapentin)	PS		
				..	C		

Date:02/26/99ISR Number: 3416975-3Report Type:Periodic
 Age: Gender:Male I/FU:F

Company Report #001-0945-950323

Outcome	PT
	Convulsion
	Dizziness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness Postural Epididymitis Haematuria	Consumer	Neurontin Capsules 300 Mg(Gabapentin)	PS		ORAL
SEE TEXT, PER		Headache					
ORAL				Dilantin (Phenytoin Sodium)	SS		ORAL
SEE TEXT, PER							
ORAL							

Date:02/26/99ISR Number: 3416977-7Report Type:Periodic Company Report #001-0945-970439
 Age:58 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG		Fatigue Hypoaesthesia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
(,DAILY), PER		Paraesthesia					
ORAL		Tremor					
				Humulin Nph (Insulin Human Injection, Isophane)	C		
				Lipitor (Atorvastatin)	C		

Date:02/26/99ISR Number: 3417009-7Report Type:Periodic Company Report #001-0945-980183
 Age:41 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2000 MG		Amnesia Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		ORAL
(DAILY) PER		Hyporeflexia					

Pain

ORAL

Lamisil (Terbinafine Hydrochloride) SS

250 MG

(DAILY)

UNKNOWN

Levsin (Hyoscyamine Sulfate) C
Imodium (Loperamide Hydrochloride) C
Elavil (Amitriptyline Hydrochloride) C
Ultram (Tramadol Hydrochloride) C

Date:02/26/99ISR Number: 3417010-3Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980185

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		
UNK, UNKNOWN		Muscular Weakness					
				Magnesium	SS		
UNK, UNKNOWN							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417013-9Report Type:Periodic
 Age:47 YR Gender:Female I/FU:I

Company Report #001-0945-980186

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG QAM, 600 MG QPM), UNKNOWN		Connective Tissue Disorder	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		
				Estrace (Estradiol) Synthroid (Levothyroxine Sodium) Lobutrin	C C C		

Date:02/26/99ISR Number: 3417015-2Report Type:Periodic
 Age:44 YR Gender:Male I/FU:I

Company Report # 001-0945-980187

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (DAILY), PER ORAL		Tinnitus	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3417018-8Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980190

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4000 MG (DAILY)		Cerebrovascular Disorder Dizziness Fall	Consumer	Neurontin (Gabapentin)	PS		

Oral Pain

UNKNOWN

Synthroid
(Levothyroxine
Sodium) C
Effexor (Venlafaxine
Hydrochloride) C

Date:02/26/99ISR Number: 3417019-XReport Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980191

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID) PER ORAL		Blood Glucose Increased Dry Mouth Vision Blurred	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3417021-8Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980192

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID) PER ORAL		Constipation Flatulence	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

Xanax (Alprazolam) C
Synthroid
(Levothyroxine)

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sodium) C
 Premarin (Estrogens
 Conjugated) C
 Cozaar (Losartan
 Potassium) C

Date:02/26/99ISR Number: 3417024-3Report Type:Periodic
 Age:59 YR Gender:Male I/FU:I

Company Report #001-0945-980193

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
SEE TEXT, PER		Feeling Abnormal Sedation	Consumer Health	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
ORAL		Vision Blurred	Professional				

Zocor (Simvastatin) C
 Aspirin
 (Acetylsalicylic
 Acid) C
 Paxil (Paroxetine
 Hydrochloride) C

Date:02/26/99ISR Number: 3417026-7Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-980194

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK, UNKNOWN		Ammonia Increased Drug Ineffective	Health Professional	Neurontin (Gabapentin)	PS		
			Company Representative				

Date:02/26/99ISR Number: 3417028-0Report Type:Periodic
 Age:50 YR Gender:Male I/FU:I

Company Report #001-0945-980195

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG		Dry Skin Skin Disorder	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		

(DAILY) ,
UNKNOWN
Skin Exfoliation
Urinary Incontinence

Phenobarbital C

Date:02/26/99ISR Number: 3417030-9Report Type:Periodic Company Report #001-0945-980196
Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Increased Flushing	Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		
100 MG (TOOK ONE DOSE), UNKNOWN		Headache Nervousness					

Hytrin (Terazosin
Hydrochloride) C
Isoptin Sr
(Verapamil
Hydrochloride) C
Ecotrin
(Acetylsalicylic
Acid) C
Potassium C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417032-2Report Type:Periodic
 Age:56 YR Gender:Female I/FU:I

Company Report #001-0945-980198

Outcome Dose	Duration	PT Drug Interaction	Report Source Consumer	Product	Role	Manufacturer	Route
1200 MG (400 MG , TID) PER ORAL		Dyskinesia		Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
10 MG (UNK) UNKNOWN				Fosamax (Alendronate Sodium)	SS		
				Pamelor (Nortriptyline Hydrochloride)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Premarin (Estrogens Conjugated)	C		
				Provera (Medroxyprogesterone Acetate)	C		
				Tenormin (Atenolol)	C		
				Asa (Acetylsalicylic Acid)	C		

Date:02/26/99ISR Number: 3417033-4Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-980199

Outcome Dose	Duration	PT Drug Interaction	Report Source Health Professional	Product	Role	Manufacturer	Route
UNK, UNKNOWN		International Normalised Ratio Increased		Neurontin (Gabapentin)	PS		
UNK, UNKNOWN				Coumadin (Warfarin Sodium)	SS		

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG	(DAILY),	Alanine Aminotransferase Increased	Health Professional	Neurontin (Gabapentin)	PS		
UNKNOWN		Aspartate Aminotransferase Increased					
UNK, UNKNOWN		Nausea		Depakote (Valproate Semisodium)	SS		
		Pain In Extremity					

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG (QHS),		Dyskinesia	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
PER ORAL				Lorcet (Paracetamol, Hydrocodone)	C		
				Zantac (Ranitidine Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417039-5Report Type:Periodic
Age:28 YR Gender:Male I/FU:I

Company Report #001-0945-980208

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Eyelid Function Disorder Nightmare	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG, TID) PER ORAL		Tremor					

Fioricet (Cafeine
Butalbital,
Paracetamol) C
Dilantin (Phenytoin
Sodium) C
Mysoline (Primidone) C
Propulsid
(Cisapride) C
Prilosec
(Omeprazole) C

Date:02/26/99ISR Number: 3417041-3Report Type:Periodic
Age:53 YR Gender:Female I/FU:I

Company Report #001-0945-980211

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated Dysgeusia	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
300 MG (DAILY) PER ORAL		Urinary Tract Infection					

Arthrotec
(Diclofenac Sodium,
Misoprostol) SS

UNK, UNKNOWN

Axid (Nizatidine) C
Bentyl
(Dicycloverine
Hydrochloride) C
Lactaid (Tilactase) C

Date:02/26/99ISR Number: 3417042-5Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980212

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety Insomnia	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
100 MG							
(DAILY), PER							
ORAL				Xanax (Alprazolam) Premarin (Estrogens Conjugated)	C C		

Date:02/26/99ISR Number: 3417043-7Report Type:Periodic
Age:47 YR Gender:Female I/FU:I

Company Report #001-0945-980214

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vision Blurred Vitreous Floaters	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 - 1200 MG							
(DAILY), PER							
ORAL				Ativan (Lorazepam)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417044-9Report Type:Periodic
 Age:74 YR Gender:Female I/FU:I

Company Report #001-0945-980215

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth Sedation	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
600 MG (200 MG, TID) PER ORAL		Sensation Of Heaviness		Dilatin (Phenytoin Sodium)	C		
				Phenobarb (Phenobarbital Sodium)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Premarin (Estrogens Conjugated)	C		
				Calcium	C		

Date:02/26/99ISR Number: 3417045-0Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980217

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State Delusion	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1800 MG (600 MG, TID), PER ORAL				Neurontin Capsules 300 Mg (Gabapentin)	SS		ORAL
1200 MG (600 MG, BID), PER ORAL				Elavil (Amitriptyline)			

Hydrochloride) C
Valium (Diazepam) C
Laxatives C

Date:02/26/99ISR Number: 3417046-2Report Type:Periodic Company Report #001-0945-980218
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Peripheral Coldness	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
2100 MG (,							
900 MG BID,							
300 MG AT							
NIGHT), PER							
ORAL							

Synthroid
(Levothyroxine
Sodium) C
Zocor (Simvastatin) C
Provera(Medroxyproge
sterone) C
Ogen (Estropipate) C
Levbid (Hyoscyamine
Sulfate) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417047-4Report Type:Periodic
 Age:37 YR Gender:Male I/FU:I

Company Report #001-0945-980220

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, BID), PER ORAL		Erythema	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Prozac (Fluoxetine Hydrochloride)	C		

Date:02/26/99ISR Number: 3417048-6Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980223

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, TID), PER ORAL		Fatigue Muscle Rigidity Palpitations Sedation	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3417049-8Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980226

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		Burning Sensation Drug Interaction Oedema Peripheral	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Zostrix (Capsaicin)	SS		

Date:02/26/99ISR Number: 3417050-4Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980227

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Breast Disorder Calcinosis	Consumer	Neurontin (Gabapentin)	PS		
				Fosamax (Alendronate Sodium)	C		
				Theodur (Theophylline)	C		
				Synthroid (Levothyroxine Sodium)	C		

Date:02/26/99ISR Number: 3417051-6Report Type:Periodic
Age:48 YR Gender:Female I/FU:I

Company Report #001-0945-980229

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia Asthenia	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL				(Tamoxifen)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417053-XReport Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-980230

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperglycaemia	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417054-1Report Type:Periodic
 Age:72 YR Gender:Female I/FU:I

Company Report #001-0945-980231

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction Muscle Twitching	Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

300 MG (100
 MG, TID), PER

ORAL

50 MG (,

QHS), PER

ORAL

Date:02/26/99ISR Number: 3417056-5Report Type:Periodic
 Age:72 YR Gender:Female I/FU:I

Company Report #001-0945-980232

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache Tinnitus	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

900 MG (300

MG, TID), PER

ORAL

Excedrin
 (Acetylsalicylic
 Acid, Caffeine,
 Salicylamide,

Paracetamol) C
 Toprol XL
 (Metoprolol
 Succinate) C
 Prilosec
 (Omeprazole) C
 Lipitor
 (Atorvastatin) C

Date:02/26/99ISR Number: 3417057-7Report Type:Periodic Company Report #001-0945-980233
 Age:51 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID), PER ORAL		Blood Bilirubin Increased Depression	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
		Drug Interaction					
		Gallbladder Disorder					
		Sedation		Risperdal (Risperidone)	SS		
				Paxil (Paroxetine Hydrochloride)	C		
				Klonopin (Clonazepam)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417058-9Report Type:Periodic
Age:19 YR Gender:Female I/FU:I

Company Report #001-0945-980234

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia Pregnancy	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL							

Prenatal Vitamins
(Ergocalciferol,
Ascorbic Acid, Folic
Acid, Thiamine
Hydrochloride, C

Date:02/26/99ISR Number: 3417059-0Report Type:Periodic
Age:50 YR Gender:Male I/FU:I

Company Report #001-0945-980235

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue Steatorrhea Weight Decreased	Health Professional	Neurontin (Gabapentin) Dextroamphetamine-(D examethasone Sulfate)	PS C		

Date:02/26/99ISR Number: 3417060-7Report Type:Periodic
Age:75 YR Gender:Female I/FU:I

Company Report #001-0945-980236

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyskinesia Dystonia	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417061-9Report Type:Periodic
Age:51 YR Gender:Female I/FU:I

Company Report #001-0945-980237

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Joint Swelling	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300							
MG, TID), PER							
ORAL							

(Atenolol) C
 Estrace (Estradiol) C
 Ativan (Lorazepam) C
 Maxzide
 (Hydrochlorothiazide
 , Triamterene) C
 (Trazodone) C
 Klonopin
 (Clonazepam) C
 (Ibuprofen) C
 (Amoxapine) C

Date:02/26/99ISR Number: 3417062-0Report Type:Periodic Company Report #001-0945-980238
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Migraine	Health Professional	Neurontin (Gabapentin)	PS		ORAL
4500 MG							
(,DAILY), PER							

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ORAL

"Many" Unspecified C

Date:02/26/99ISR Number: 3417063-2Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980239

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG (300 MG, EVERY 3 HOURS), PER		Feeling Abnormal Weight Increased	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

ORAL

Date:02/26/99ISR Number: 3417064-4Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980240

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dyskinesia	Health Professional	Neurontin (Gabapentin) Dilantin (Phenytoin Sodium)	PS C		

Date:02/26/99ISR Number: 3417065-6Report Type:Periodic
Age:63 YR Gender:Female I/FU:I

Company Report #001-0945-980244

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, TID), PER		Eye Movement Disorder Nausea Oedema Peripheral	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

ORAL

Roxicet

(Paracetamol,
Oxycodone
Hydrochloride) C
(Calcium) C

Date:02/26/99ISR Number: 3417066-8Report Type:Periodic Company Report #001-0945-980245
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis Urinary Incontinence		Neurontin (Gabapentin)	PS		ORAL
ONE CAPSULE							
IN AM; ONE							
CAPSULE IN							
AM; ONE							
CAPSULE AT HS							

Alphagan
(Brimonidine
Tartrate) C
Timoptic Xe (Timolol
Maleate) C
Risperdal
(Risperidone) C
Ativan (Lorazepam) C
Xalatan
(Latanoprost) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417067-XReport Type:Periodic
Age:65 YR Gender:Male I/FU:F

Company Report #001-0945-970625

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, TID) PER ORAL		Cerebrovascular Accident Sedation	Health Professional	Neurontin 300 Mg (Gabapentin)	PS		ORAL
				Rezulin (Troglitazone)	C		
				Lipitor (Atorvastatin)	C		
				Cozaar (Losartan Potassium)	C		
				Glucophage (Metformin Hydrochloride)	C		
				Toprol (Metoprolol Succinate)	C		
				Norvasc (Amlodipine Besilate)	C		
				(Insulin)	C		
				Ecotrin (Acetylsalicylic Acid)	C		
				Elavil (Amitriptyline Hydrochloride)	C		

Date:02/26/99ISR Number: 3417068-1Report Type:Periodic
Age:40 YR Gender:Male I/FU:F

Company Report #001-0945-970665

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		Clumsiness Disturbance In Attention Visual Disturbance	Consumer	Neurontin 300 Mg (Gabapentin)	PS		ORAL

Azmacort
(Triamcinolone
Acetonide) C

Date:02/26/99ISR Number: 3417069-3Report Type:Periodic Company Report #001-0945-970686
Age:13 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper Gallbladder Disorder	Consumer Health	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
400 MG (, 100 MG QAM, 100 MG QPM, 200 MG QHS), PER ORAL		Polyp	Professional				
				Veetids (Phenoxymethylpenici llin Potassium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417070-XReport Type:Periodic
Age:51 YR Gender:Female I/FU:F

Company Report #001-0945-970703

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue Nausea	Health Professional	Neurontin 300 Mg (Gabapentin)	PS		ORAL
1200 MG (300 MG, QID), PER ORAL		Pancreatitis					
				Prosom (Estazolam)	C		
				Doxepin	C		
				Diazepam	C		
				Hydroxyzine	C		
				Surmontil (Trimipramine)	C		

Date:02/26/99ISR Number: 3417071-1Report Type:Periodic
Age:32 YR Gender:Female I/FU:F

Company Report #001-0945-970708

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG, TID), PER ORAL							
				Amitriptyline	C		
				Verapamil	C		

Date:02/26/99ISR Number: 3417072-3Report Type:Periodic
Age:35 YR Gender:Unknown I/FU:F

Company Report #001-0945-970709

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Facial Palsy	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417073-5Report Type:Periodic Company Report #001-0945-970710
Age:35 YR Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Facial Palsy	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417074-7Report Type:Periodic Company Report #001-0945-970711
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Polymenorrhoea	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417075-9Report Type:Periodic Company Report #001-0945-970712
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menstruation Irregular	Health Professional	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417076-0Report Type:Periodic
 Age: Gender:Female I/FU:F

Company Report #001-0945-970713

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Menstrual Disorder	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417077-2Report Type:Periodic
 Age: Gender:Female I/FU:F

Company Report #001-0945-970714

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Polymenorrhoea	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417078-4Report Type:Periodic
 Age:42 YR Gender:Male I/FU:F

Company Report #001-0945-970716

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Influenza Like Illness	Health Professional	Neurontin (Gabapentin)	PS		
200 MG (AT HS), UNKNOWN							

Date:02/26/99ISR Number: 3417079-6Report Type:Periodic
 Age:31 YR Gender:Male I/FU:F

Company Report #001-0945-970719

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction Vomiting	Health Professional	Neurontin 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID) , PER ORAL							

Novocain (Procaine Hydrochloride)	SS
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Synthroid
(Levothyroxine
Sodium) C
Humalog (Insulin
Lispro) C

Date:02/26/99ISR Number: 3417080-2Report Type:Periodic Company Report #001-0945-973007
Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Delirium Sepsis	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417081-4Report Type:Periodic Company Report #001-0945-973020
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Platelet Count Decreased		Neurontin (Gabapentin)	PS		
300 MG							
(DAILY),							
UNKNOWN				Klonopin (Clonazepam)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vistaril
 (Hydroxyzine
 Embonate) C
 Ativan (Lorazepam) C

Date:02/26/99ISR Number: 3417082-6Report Type:Periodic
 Age:50 YR Gender:Female I/FU:F

Company Report #001-0945-973031

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (,300 MG TID), PER ORAL		Hallucination, Visual	Health Professional Company Representative	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

Elavil
 (Amitriptyline
 Hydrochloride) C

Date:02/26/99ISR Number: 3417083-8Report Type:Periodic
 Age:54 YR Gender:Female I/FU:F

Company Report #001-0945-973036

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (200 MG BID), PER ORAL		Dizziness Epistaxis Nasal Discomfort Throat Irritation	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Fosamax (Alendronate
 Sodium) C
 Prempro
 (Medroxyprogesterone
 Acetate, Estrogens
 Conjugated) C

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, BID)		Constipation Hirsutism	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		
UNKNOWN				Prozac (Fluoxetine Hydrochloride)	C		
				Claritin (Loratadine)	C		
				Nasacort Inhaler (Triamcinolone Acetate)	C		
				Aygestin (Norethisterone Acetate)	C		
				Darvocet (Paracetamol, Dextropropoxyphene)	C		
				Skelaxin (Metaxalone)	C		
				Voltaren (Diclofenac Sodium)	C		
				Ultram (Tramadol)			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C
 Humibid
 (Guaifenesin,
 Dextromethorphan
 Hydrobromide) C

Date:02/26/99ISR Number: 3417085-1Report Type:Periodic
 Age:12 YR Gender:Male I/FU:F

Company Report #001-0945-973044

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Appetite Food Craving	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		
900 MG (300 MG, TID)		Weight Increased					
UNKNOWN							

Prozac (Fluoxetine
 Hydrochloride) C
 Tofranil (Imipramine
 Hydrochloride) C
 Sudafed
 (Pseudoephedrine
 Hydrochloride) C

Date:02/26/99ISR Number: 3417086-3Report Type:Periodic
 Age:55 YR Gender:Female I/FU:F

Company Report #001-0945-980004

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression Oedema	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER							
ORAL							

Zyprexa (Olanzapine) C
 Prozac (Fluoxetine
 Hydrochloride) C
 Depakote (Valproate
 Semisodium) C
 Adderall (Amfetamine

Sulfate)

C

Date:02/26/99ISR Number: 3417087-5Report Type:Periodic
Age:47 YR Gender:Female I/FU:I

Company Report #001-0945-980880

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Burning Sensation	Health	Neurontin Capsules			
		Paraesthesia	Professional	100mg (Gabapentin)	PS		ORAL
100MG DAILY							
PER ORAL		Rash Pruritic					
				Prozac (Fluoxetine Hydrochloride)	C		

Date:02/26/99ISR Number: 3417088-7Report Type:Periodic
Age:8 YR Gender:Female I/FU:I

Company Report #001-0945-980881

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Weight Increased	Consumer	Neurontin Capsules			
				300mg (Gabapentin)	PS		ORAL
2400MG IN							
DIVIDED DOSES							
PER ORAL				Lithium	C		
				Tenex (Guanfactine)			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C

Date:02/26/99ISR Number: 3417089-9Report Type:Periodic
Age:42 YR Gender:Male I/FU:I

Company Report #001-0945-980882

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG X1 DOSE PER ORAL		Burning Sensation Hypoaesthesia Muscle Twitching Paraesthesia Tongue Oedema	Consumer	Neurontin Capsules 300mg (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3417090-5Report Type:Periodic
Age:68 YR Gender:Female I/FU:I

Company Report #001-0945-980883

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
SEE IMAGE		Blood Pressure Increased Dizziness Fatigue Herpes Zoster Keratoconjunctivitis Sicca Migraine	Consumer	Neurontin Capsules 300mg (Gabapentin) Darvocet-N (Paracetamol) Dextropropoxyphene Imitrex (Sumatriptan) Herbal Preparation	PS C C C C		ORAL

Date:02/26/99ISR Number: 3417091-7Report Type:Periodic
Age:61 YR Gender:Female I/FU:I

Company Report #001-0945-980884

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900MG DAILY PER ORAL		Hearing Impaired Tinnitus	Consumer	Neurontin Capsules 300mg (Gabapentin) Estrace (Estradiol) Provera	PS C		ORAL

(Medroxyprogesterone
Acetate) C
Imodium (Loperamide
Hydrochloride) C
Allerga
(Fexofenadine
Hydrochloride) C

Date:02/26/99ISR Number: 3417092-9Report Type:Periodic
Age:44 YR Gender:Female I/FU:I

Company Report #001-0945-980885

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		
SEE TEXT				Lorazepam	SS		ORAL
2MG 1MG BID							
PER ORAL				Rocephin (Ceftriaxone Sodium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417093-0Report Type:Periodic
Age:69 YR Gender:Male I/FU:I

Company Report #001-0945-980887

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400MG DAILY		Condition Aggravated Tremor	Consumer	Neurontin Capsules 400mg (Gabapentin)	PS		ORAL
PER ORAL				Sinemet (Levodopa, Carbidopa)	C		

Date:02/26/99ISR Number: 3417094-2Report Type:Periodic
Age:43 YR Gender:Male I/FU:I

Company Report #001-0945-980889

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG DAILY		Urticaria	Consumer	Neurontin Capsules 300mg (Gabapentin)	PS		ORAL
PER ORAL							

Date:02/26/99ISR Number: 3417095-4Report Type:Periodic
Age:76 YR Gender:Female I/FU:I

Company Report #001-0945-980890

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
800MG (400MG BID) PER ORAL		Constipation Fall	Consumer	Neurontin Capsules 100mg (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3417096-6Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980891

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Diabetes Mellitus	Health	Neurontin			

Professional (Gabapentin) PS
Dilantin (Phenytoin Sodium) C

Date:02/26/99ISR Number: 3417097-8Report Type:Periodic Company Report #001-0945-980892
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Purpura	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417098-XReport Type:Periodic Company Report #001-0945-980893
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Balance Disorder Decreased Activity	Consumer	Neurontin Capsules 100mg (Gabapentin)	PS		ORAL
300MG (100MG TID) PER ORAL		Dizziness Mental Impairment		Aldactone (Spironolactone) K-Dur (Potassium Chloride) Lasix (Furosemide) Aspirin (Acetylsalicylic Acid)	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Magnesium C
 Iso (Isosorbide C
 Mononitrate)

Date:02/26/99ISR Number: 3417099-1Report Type:Periodic Company Report #001-0945-980894
 Age:43 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3600MG (900		Aphonia Chest Pain	Consumer	Neurontin Capsules 300mg (Gabapentin)	PS		ORAL
MG QID) PER		Dyspnoea					
ORAL		Pharyngolaryngeal Pain					
		Weight Increased		Prozac (Fluoxetine Hydrochloride0 Clonazepam	C C		

Date:02/26/99ISR Number: 3417100-5Report Type:Periodic Company Report #001-0945-980895
 Age:52 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
700MG DAILY		Condition Aggravated Fatigue	Health Professional	Neurontin (Gabapentin)	PS		ORAL
PER ORAL				Ultram (Tramadol Hydrochloride)	C		

Date:02/26/99ISR Number: 3417101-7Report Type:Periodic Company Report #001-0945-980897
 Age:9 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2700MG (900MG		Rectal Haemorrhage	Health Professional	Neurontin Capsules 300mg (Gabapentin)	PS		ORAL

TID) PER ORAL

Date:02/26/99ISR Number: 3417102-9Report Type:Periodic Company Report #001-0945-980898
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema Oedema Mouth Stomatitis	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417103-0Report Type:Periodic Company Report #001-0945-980899
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cataract Night Blindness	Consumer Health Professional	Neurontin (Gabapentin)	PS		
1200MG (600MG BID)				Voltaren (Diclofenac Sodium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417104-2Report Type:Periodic
Age:75 YR Gender:Female I/FU:I

Company Report #001-0945-980900

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache	Health Professional	Neurontin Capsules 300mg (Gabapentin)	PS		ORAL
300MG (ONE DOSE) PER ORAL							

Date:02/26/99ISR Number: 3417105-4Report Type:Periodic
Age:69 YR Gender:Female I/FU:I

Company Report #001-0945-980901

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue Increased Appetite	Consumer	Neurontin Capsules 300mg (Gabapentin)	PS		ORAL
300MG (X ONE DOSE) PER ORAL							

Date:02/26/99ISR Number: 3417106-6Report Type:Periodic
Age:80 YR Gender:Female I/FU:I

Company Report #001-0945-980902

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Keratoconjunctivitis Sicca	Consumer	Neurontin Capsules 100mg (Gabapentin)	PS		ORAL
400MG DAILY PER ORAL							
				Cozaar (Losartan Potassium)	C		

Date:02/26/99ISR Number: 3417321-1Report Type:Periodic
Age:17 YR Gender:Female I/FU:F

Company Report #001-0945-980009

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		Alopecia Dry Mouth Fatigue Headache Mood Swings Nausea Sedation Tremor	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Klonopin (Clonazepam) Steroid Injections	SS C		

Date:02/26/99ISR Number: 3417323-5Report Type:Periodic Company Report #001-0945-980010
 Age:22 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, BID), PER ORAL		Asthma	Health Professional	Neurontin Capsules 300 Mg(Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3417324-7Report Type:Periodic Company Report #001-0945-980012
 Age:61 YR Gender:Female I/FU:F

Outcome	PT
	Dry Mouth Eye Irritation Fluid Retention

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
1800 MG	(DAILY), PER	Nervousness Nightmare Oedema Peripheral	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
		Sedation Sleep Disorder					
				Inderal (Codeine)	C		
				Tranxene	C		
				Magnesium	C		
				Estrace (Progesterone)	C		

Date:02/26/99ISR Number: 3417326-0Report Type:Periodic Company Report #001-0945-980019
 Age:43 YR Gender:Male I/FU:F

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
1200 MG (300	MG, QID), PER	Diplopia Dizziness	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
		Flatulence					
		Headache					
		Increased Bronchial Secretion Nasal Congestion Sedation		Dilantin	C		

Date:02/26/99ISR Number: 3417329-6Report Type:Periodic Company Report #001-0945-980020
 Age:57 YR Gender:Male I/FU:F

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
2900 MG		Amnesia Dizziness	Health Professional	Neurontin (Gabapentin)	PS		

(DAILY) Erectile Dysfunction

Fatigue	Glucophage	C
Vision Blurred	Humulin N	C
	Humulin Nph	C

Date:02/26/99ISR Number: 3417332-6Report Type:Periodic Company Report #001-0945-980023
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Platelet Count Decreased	Health Professional	Neurontin (Gabapentin)	PS		
				Toprol	C		
				Ambien	C		

Date:02/26/99ISR Number: 3417337-5Report Type:Periodic Company Report #001-0945-980024
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 MG (400							
MG, TID), PER							
ORAL							
				Lioresal	C		
				Betaseron	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417339-9Report Type:Periodic
 Age: Gender: I/FU:I

Company Report #001-0945-980541

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abdominal Pain Upper	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417342-9Report Type:Periodic
 Age:5 YR Gender:Male I/FU:I

Company Report #001-0945-980542

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Psychomotor Hyperactivity	Consumer	Neurontin (Gabapentin)	PS		ORAL

900 MG (300
 MG, TID), PER
 ORAL

Date:02/26/99ISR Number: 3417344-2Report Type:Periodic
 Age:18 YR Gender:Female I/FU:I

Company Report #001-0945-980544

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Joint Contracture	Health Professional	Neurontin (Gabapentin)	PS		ORAL

900 MG (300
 MG, TID), PER
 ORAL

7 DAY

Atrovent C
 Relafen C

Date:02/26/99ISR Number: 3417345-4Report Type:Periodic
 Age: Gender: I/FU:I

Company Report #001-0945-980546

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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SEE TEXT, PER
ORAL

Anxiety	Health	Neurontin			
Depression	Professional	(Gabapentin)	PS		ORAL
Tremor					

Date:02/26/99ISR Number: 3417347-8Report Type:Periodic Company Report #001-0945-980547
Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypoaesthesia	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL							

Date:02/26/99ISR Number: 3417349-1Report Type:Periodic Company Report #001-0945-980548
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
300 MG (QHS), PER ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3417351-XReport Type:Periodic
 Age:40 YR Gender:Male I/FU:I

Company Report #001-0945-980549

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG IN AM, 600 MG IN PM), PER ORAL		Confusional State Dizziness Hypoaesthesia	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Synthroid	C		
				Serzone	C		

Date:02/26/99ISR Number: 3417360-0Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980551

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Back Pain	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3417361-2Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980552

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (300MG QAM, 300 MG QPM, 600MG QHS)		Disorientation Dizziness Medication Error Pain Sedation	Consumer Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		
				Daypro	C		
				Vitamin E	C		
				Belphenergots	C		

Date:02/26/99ISR Number: 3417363-6Report Type:Periodic Company Report #001-0945-980554
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension Ecchymosis	Consumer	Neurontin (Gabapentin)	PS		
SEE TEXT		Weight Increased					

Date:02/26/99ISR Number: 3423642-9Report Type:Periodic Company Report #001-0945-980849
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urinary Incontinence	Consumer	Neurontin (Gabapentin)	PS		ORAL
1600 MG (400 MG, QID), PER ORAL				Unspecified Antidepressants Vicodin (Paracetamol, Hydrocodone Bitartrate) Tylenol #3 (Codeine Phosphate, Paracetamol)	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3423643-0Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980850

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3423644-2Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980851

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea Myasthenia Gravis	Health Professional Company Representative	Neurontin (Gabapentin) Mestinon (Pyridostigmine Bromide)	PS C		

Date:02/26/99ISR Number: 3423646-6Report Type:Periodic
Age:80 YR Gender:Male I/FU:I

Company Report #001-0945-980852

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension Flatulence	Consumer	Neurontin (Gabapentin)	PS		ORAL

1500 MG (500
MG, TID), PER

ORAL

Coumadin (Warfarin Sodium)	C
Vasotec (Enalapril Maleate)	C
Lasix (Furosemide)	C
K-Dur (Potassium Chloride)	C

Date:02/26/99ISR Number: 3423648-XReport Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-980853

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Gynaecomastia	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3423651-XReport Type:Periodic Company Report #001-0945-980855
 Age:54 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, BID), PER ORAL		Amnesia Coeliac Disease Joint Swelling	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Premarin (Estrogens Conjugated) Unspecified Oral Antihyperglycemic Med	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3423652-1Report Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980857

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Vaginal Haemorrhage	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1400 MG							
(DAILY), PER							
ORAL							

Prozac (Fluoxetine Hydrochloride)	C	
Risperdal (Risperidone)	C	

Date:02/26/99ISR Number: 3423655-7Report Type:Periodic
Age:35 YR Gender:Female I/FU:I

Company Report #001-0945-980858

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea Nausea	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1200 MG							
(DAILY), PER		Sedation					
ORAL		Urticaria					

Paxil (Paroxetine Hydrochloride)	C	
Premarin (Estrogens Conjugated)	C	
Depakote (Valproate Semisodium)	C	
Relafen (Nabumetone)	C	

Date:02/26/99ISR Number: 3423658-2Report Type:Periodic
Age:39 YR Gender:Female I/FU:I

Company Report #001-0945-980862

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia	Consumer	Neurontin Capsules			

1200 MG (400	Health	400 Mg (Gabapentin)	PS	ORAL
MG, TID), PER	Professional			
ORAL		Tegretol (Carbamazepine)	C	
		Paxil (Paroxetine Hydrochloride)	C	
		(Lorazepam)	C	

Date:02/26/99ISR Number: 3423659-4Report Type:Periodic Company Report #001-0945-980863
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Bradykinesia Tremor	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
800 MG (400							
MG, BID), PER							
ORAL				Prolixin (Fluphenazine Hydrochloride)	C		
				Risperdal (Risperidone)	C		
				Depakote (Valproate Semisodium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3424030-1Report Type:Periodic
 Age:53 YR Gender:Female I/FU:I

Company Report #001-0945-980627

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea Nystagmus	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG		Syncope					
(DAILY), PER							
ORAL							
				Asa	C		
				Depakote	C		
				Paxil	C		
				Dalmane	C		

Date:02/26/99ISR Number: 3424032-5Report Type:Periodic
 Age:70 YR Gender:Female I/FU:I

Company Report #001-0945-980628

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper Headache	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG		Stomatitis					
(DAILY), PER							
ORAL							
				Robaxin	C		

Date:02/26/99ISR Number: 3424033-7Report Type:Periodic
 Age:53 YR Gender:Male I/FU:I

Company Report #001-0945-980630

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysarthria Hypoaesthesia	Consumer	Neurontin (Gabapentin)	PS		ORAL
2700 MG							
(DAILY), PER							
ORAL							

Klonopin

C

Date:02/26/99ISR Number: 3424042-8Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980632

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Water Intoxication	Consumer	Neurontin (Gabapentin)	PS		
				Tagamet Hb 200 (Cimetidine)	SS		

Date:02/26/99ISR Number: 3424044-1Report Type:Periodic
Age:52 YR Gender:Female I/FU:I

Company Report #001-0945-980633

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache Myalgia	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

100 MG
(DAILY), PER

ORAL

Zoloft	C
Vistaril	C
Motrin	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3424046-5Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-980634

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
500 MG (DAILY), PER ORAL		Alopecia Burning Sensation Lip Disorder Muscle Twitching Nail Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3424048-9Report Type:Periodic
 Age:35 YR Gender:Male I/FU:I

Company Report #001-0945-980635

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		Anorgasmia Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		ORAL
2 TABS (AS NEEDED)				Propoxyphene Napsylate With Apap (Paracetamol, Dextropropoxyphene Napsilate)	SS		
				Fiorinal	C		

Date:02/26/99ISR Number: 3424050-7Report Type:Periodic
 Age:47 YR Gender:Female I/FU:I

Company Report #001-0945-980636

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia	Consumer	Neurontin			

400 MG Blood Pressure Increased (Gabapentin) PS ORAL
 (DAILY), PER Crying
 ORAL Dyspnoea
 Erythema
 Feeling Abnormal
 Flushing
 Medication Error

Date:02/26/99ISR Number: 3424051-9Report Type:Periodic Company Report #001-0945-980637
 Age:44 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (100 MG, QID) PER		Dizziness Pallor	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Copaxone	C		
				Klonopin	C		
				Cylert	C		
				Pamelor	C		

Date:02/26/99ISR Number: 3424053-2Report Type:Periodic Company Report #001-0945-980638
 Age:41 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG (100		Dizziness Nausea	Consumer	Neurontin (Gabapentin)	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

MG, BID), PER

ORAL

100 MG (QHS),

PER ORAL

Luvox (Fluvoxamine
Maleate)

SS

ORAL

Prozac
Synthroid
Ativan

C
C
C

Date:02/26/99ISR Number: 3424055-6Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-980437

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2800 MG		Dizziness Overdose	Consumer	Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER		Photosensitivity Reaction					
ORAL		Sedation					
		Syncope					

Date:02/26/99ISR Number: 3424057-XReport Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-980438

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG		Drug Interaction Grand Mal Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
(1200 MG,			Company				
BID), PER			Representative				
ORAL				Wellbutrin (Amfebutamone)			

375 (DAILY),
 PER ORAL
 Hydrochloride) SS ORAL
 Prozac C
 Klonopin C

Date:02/26/99ISR Number: 3424059-3Report Type:Periodic Company Report #001-0945-980439
 Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG		Convulsion					
(DAILY), PER		Ear Infection					
ORAL				Tegretol Synthroid	C C		

Date:02/26/99ISR Number: 3424061-1Report Type:Periodic Company Report #001-0945-980442
 Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erectile Dysfunction Loss Of Libido	Consumer	Neurontin (Gabapentin)	PS		ORAL
2100 MG							
(DAILY), PER							
ORAL				Norvasc (Quinapril) (Ibuprofen) Prozac	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Serax C
 (Lithium) C
 Micronase C

Date:02/26/99ISR Number: 3424063-5Report Type:Periodic Company Report #001-0945-980443
 Age:80 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG (HS), PER ORAL		Abdominal Pain Nausea	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

Date:02/26/99ISR Number: 3424064-7Report Type:Periodic Company Report #001-0945-980444
 Age:13 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (DAILY), PER ORAL		Dyskinesia Tremor	Consumer	Neurontin (Gabapentin)	PS		ORAL

Voltaren C
 Zantac C
 (Atenolol) C

Date:02/26/99ISR Number: 3424067-2Report Type:Periodic Company Report #001-0945-980445
 Age:68 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (400 MG, TID), PER		Diarrhoea Uterine Haemorrhage	Health Professional	Neurontin (Gabapentin)	PS		ORAL

ORAL

Date:02/26/99ISR Number: 3424069-6Report Type:Periodic Company Report #001-0945-980446
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension Hair Texture Abnormal	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1800 MG (600 MG, TID), PER ORAL		Increased Appetite Oedema Peripheral Weight Increased					

Date:02/26/99ISR Number: 3424071-4Report Type:Periodic Company Report #001-0945-980448
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Systemic Lupus Erythematosis	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3424072-6Report Type:Periodic
Age:80 YR Gender:Male I/FU:I

Company Report #001-0945-980452

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Liver Function Test Abnormal	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL				Bactrim Ds (Sulfamethoxazole, Trimethoprim)	SS		
(BID)							

Date:02/26/99ISR Number: 3424075-1Report Type:Periodic
Age:43 YR Gender:Male I/FU:I

Company Report #001-0945-980453

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sedation	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG (SEE TEXT), PER ORAL				Cardura	C		
				Klonopin	C		
				Hyzaar	C		
				(Verapamil)	C		
				Effexor Xr	C		
				Atarax	C		
				Dexacort	C		

Date:02/26/99ISR Number: 3424076-3Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-980454

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Urine Analysis Abnormal	Health	Neurontin			

Professional (Gabapentin) PS
Trazodone C
Inderal C

Date:02/26/99ISR Number: 3424077-5Report Type:Periodic Company Report #001-0945-980456
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiac Failure Oedema Peripheral	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3424078-7Report Type:Periodic Company Report #001-0945-980457
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Myalgia	Consumer	Neurontin (Gabapentin)	PS		

Date:02/26/99ISR Number: 3424079-9Report Type:Periodic Company Report #001-0945-980459
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Libido Decreased	Health Professional	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/99ISR Number: 3424080-5Report Type:Periodic
Age:66 YR Gender:Female I/FU:I

Company Report #001-0945-980461

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Increased	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1200 MG (300 MG, QID), PER ORAL				Glyburide (Furosemide) (Potassium) Prozac Anturane	C C C C C		

Date:02/26/99ISR Number: 3424081-7Report Type:Periodic
Age:72 YR Gender:Male I/FU:I

Company Report #001-0945-980462

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Skin Odour Abnormal	Consumer	Neurontin (Gabapentin)	PS		
400 MG (QHS)				Hytrin Mevacor Prozac	C C C		

Date:03/01/99ISR Number: 3208259-3Report Type:Direct
Age:24 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Health Professional	Neurontin	PS	Parke-Davis	ORAL
300 MG / TID / ORAL	1 WK			Haldol Benadryl Cogentin Zyprexa	C C C C		

Date:03/01/99ISR Number: 3218648-9Report Type:Periodic Company Report #A0077155
 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	Stevens-Johnson Syndrome	Health Professional	Zyban Tablet - Zyban	PS		ORAL
			Gabapentin Capsule	SS		ORAL

Date:03/02/99ISR Number: 3210320-4Report Type:Expedited (15-DaCompany Report #001-0945-980783
 Age:51 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 400 MG (100 MG, QID), PER ORAL	Diabetes Mellitus Diarrhoea Dizziness Nervous System Disorder Orthostatic Hypotension Renal Disorder	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
			Cozaar (Losartan Potassium) (Insulin) Nph Insulin (Insulin Injection, Isophane) Proamatine	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Midodrine
Hydrochloride) C
Lasix (Furosemide) C

Date:03/03/99ISR Number: 3210649-XReport Type:Direct
Age:86 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Gabapentin	PS		
				Carbamazepine	SS		
				Lamotrigine	SS		

Date:03/03/99ISR Number: 3211629-0Report Type:Expedited (15-DaCompany Report #001-0945-980676
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Fall Gout	Consumer	Neutrontin (Gabapentin)	PS		ORAL

900 MG (300
MG, TID) PER
Weight Decreased

ORAL

Colchicine C
Coumadin C
Isosorbide C
Norvasc C
Hytrin C
Lanoxin C
Prinivil C
Lasix C
Levodopa C
K-Dur C

Date:03/04/99ISR Number: 3212859-4Report Type:Expedited (15-DaCompany Report #001-0945-990133
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Health	Neurontin Capsules			

900 MG (300
MG, TID),
UNKNOWN
Drug Effect Decreased Professional 300 Mg (Gabapentin) PS

Date:03/05/99ISR Number: 3214125-XReport Type:Direct Company Report #
Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain		Neurontin	PS		ORAL
Other		Convulsion		Dilantin	SS		
PO		Cough					
		Drug Level Below					
		Therapeutic					

Date:03/05/99ISR Number: 3214470-8Report Type:Expedited (15-DaCompany Report #A0082673
Age:44 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Drug Level Above
	Therapeutic

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Feeling Drunk Nausea Tremor	Report Source	Product	Role	Manufacturer	Route
ORAL			Health	Wellbutrin Tablet	PS		ORAL
ORAL			Professional Other	Lithium Carbonate Tablet	SS		ORAL
ORAL				Olanzapine Tablet	SS		ORAL
ORAL				Benzatropine Tablet	SS		ORAL
ORAL				Gabapentin Tablet	SS		ORAL
ORAL				Clonazepam Tablet	SS		ORAL
ORAL				Zolpidem Tartrate Tablet	SS		ORAL
ORAL				Sertraline Hydrochloride Tablet	SS		ORAL

Date:03/08/99ISR Number: 3216166-5Report Type:Direct
Age:72 YR Gender: I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG TID		Coordination Abnormal	Health	Gabapentin	PS		
Initial or Prolonged		Mental Disorder	Professional	Esgic	C		

Date:03/09/99ISR Number: 3216360-3Report Type:Direct
Age:32 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other CAPSULE		Drug Level Below Therapeutic		Neoral (Cyclosporine)	PS	Novartis	
		Graft Versus Host Disease		Neurontin			

Date:03/10/99ISR Number: 3216642-5Report Type:Expedited (15-DaCompany Report #001-0945-990120
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 Other MG, TID), PER		Cardiac Failure Congestive Condition Aggravated Depression	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
ORAL		Haematocrit Decreased Haemoglobin Decreased Personality Disorder Post-Traumatic Stress Disorder Psychotic Disorder Red Blood Cell Count Decreased		Mellaril (Thiorida - Zine Hydrochloride) (Clonidine) Anafranil (Clomipramine Hydrochloride) Robitussin (Guaifenesin) Peri-Colace (Docusate Sodium, Casanthranol) Seroquel (Quetiapine Fumarate) (Trazodone)	C C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/11/99ISR Number: 3218193-0Report Type:Expedited (15-DaCompany Report #001-0945-990126
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Glucose Increased Glycosuria	Consumer	Neurontin Capsules (Gabapentin)	PS		ORAL
900 MG (300 MG, TID) PER ORAL							

Date:03/11/99ISR Number: 3218768-9Report Type:Direct Company Report #
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1200MG PO TID Initial or Prolonged PTA CONT AT DISCHARGE LOWER DOSE		Abnormal Behaviour Appetite Disorder Confusional State Hallucination Movement Disorder		Neurontin Ativan Neutontin Srytin Lortab	PS C C C C		ORAL

Date:03/12/99ISR Number: 3219027-0Report Type:Expedited (15-DaCompany Report #001-0945-990153
 Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Glucose Decreased Convulsion Movement Disorder Tremor	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
300 MG (,DAILY), PER ORAL							

Nph Insulin (Insulin
 Injection, Isophane) C
 (Insulin) C
 Zoloft (Sertraline
 Hydrochloride) C
 Creon (Pancreatin) C
 (Furosemide) C
 Accupril (Quinapril
 Hydrochloride) C
 Miacalcin
 9calcitonin, Salmon) C

Date:03/12/99ISR Number: 3219309-2Report Type:Expedited (15-DaCompany Report #201427

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required		Ecchymosis Haematocoele Female Injury	Foreign Other	Rivotril (Clonazepam) 2.5 Mg/Ml	PS		ORAL
20 DOSE FORM Intervention to DAILY ORAL		Thyroid Disorder					
Prevent Permanent SUBCUTANEOUS	20 MG DAILY			Copaxone (Copaxone)	SS		
Impairment/Damage SUBCUTANEOUS				Mysoline (Primidone) 250 Mg	SS		ORAL
250 MG DAILY ORAL				Atarax (Hydroxyzine Hydrochloride) 25mg	SS		ORAL
50 MG DAILY ORAL				Anafranil			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

75 MG DAILY	(Clomipramine Hydrochloride) 75 Mg	SS	ORAL
ORAL			
900 MG DAILY	Neurontin (Gabapentin) 300 Mg	SS	ORAL
ORAL			
	Xatral	C	
	Topalgic	C	

Date:03/12/99ISR Number: 3219479-6Report Type:Expedited (15-DaCompany Report #033-0945-990018
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG	Ecchymosis Thyroid Disorder	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY) PER			Professional				
ORAL							

Anafranil (Clomipamine Hydrochloride)	C
Rivoril (Clonazepam)	C
Atarax (Hydroxyzine)	C
Mysoline (Primidone)	C
Xatral Lp (Alfuzosin)	C
Copaxone (Copolymer)	C
Topalgic	C

Date:03/15/99ISR Number: 3221115-XReport Type:Expedited (15-DaCompany Report #001-0945-990130
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1800 MG	Abortion Spontaneous	Health	Neurontin	PS		

(,DAILY),

UNKNOWN

Prenatal Vitamins C
Folic Acid C

Date:03/15/99ISR Number: 3221209-9Report Type:Expedited (15-DaCompany Report #001-0991-990448

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abortion Spontaneous	Consumer	Rezulin			
Other		Complications Of Maternal		(Troglitazone)	PS		ORAL
400 MG		Exposure To Therapeutic					
(,DAILY), PER		Drugs					
ORAL		Drug Ineffective		Lipitor			
				(Atorvastatin)	SS		
				Neurontin			
				(Gabapentin)	SS		
				Wellbutrin	C		
				Humulin N	C		
				Lipitor	C		
				Neurontin	C		
				Humulin N	C		
				Humulin R	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/16/99ISR Number: 3222091-6Report Type:Expedited (15-DaCompany Report #001-0945-990140
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Interaction	Health	Neurontin Capsules			
Hospitalization - 900 MG (300 Initial or Prolonged MG, TID)		Hepatic Necrosis	Professional	300 Mg (Gabapentin)	PS		
		Hepatotoxicity		(Ciclosporin)	SS		
		Multi-Organ Failure		(Azathioprine)	SS		
				(Prednisone)	SS		
				(Ranitidine)	SS		
				(Baclofen)	SS		
				(Atorvastatin)	SS		
				(Ketoconazole)	SS		
				Acetaminophen			
				(Paracetamol)	SS		
				Niacin (Nicotinic Acid)	SS		
				Dilaudid			
				(Hydromorphone Hydrochloride)	SS		
				(Alprazolam)	SS		
				(Losartan)	SS		
				(Atenolol)	SS		
				(Diltiazem)	C		

Date:03/17/99ISR Number: 3222507-5Report Type:Expedited (15-DaCompany Report #001-0945-990170
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG QAM; AND 200 MG QPM (, DAILY) , PER ORAL		Medication Error	Health Professional	Neurontin Capsules 100 Mg(Gabapentin)	PS		ORAL

Date:03/17/99ISR Number: 3222551-8Report Type:Expedited (15-DaCompany Report #001-0945-990142
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hyperprolactinaemia	Health	Neurontin	PS		ORAL
1200 MG (300		Pituitary Tumour	Professional				
MG, QID), PER							
ORAL							

Date:03/17/99ISR Number: 3222574-9Report Type:Expedited (15-DaCompany Report #001-0945-990141
Age:8 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Ear Infection	Consumer	Neurontin			
Initial or Prolonged		Influenza Like Illness		(Gabapentin)	PS		ORAL
2400 MG							
(DAILY), PER							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/19/99ISR Number: 3223501-0Report Type:Expedited (15-DaCompany Report #99F--10195

Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 75 MG, DAILY, Initial or Prolonged ORAL		Ecchymosis	Foreign Health	Anafranil	PS		ORAL
250 MG, DAILY, ORAL		Haematocoele	Professional	Mysoline	SS		ORAL
50 MG, DAILY, ORAL			Other	Atarax	SS		ORAL
20 DRP, DAILY, ORAL				Rivotril	SS		ORAL
900 MG, DAILY, ORAL				Gabapentin	SS		ORAL
SUBCUTANEOUS 20 MG, DAILY, SUBCUTANEOUS 5 MON				Copaxone	SS		

Date:03/22/99ISR Number: 3224033-6Report Type:Expedited (15-DaCompany Report #002-0945-990011

Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (DAILY), PER ORAL/600 MG DAILY/300 MG		Dizziness Drug Withdrawal Syndrome Hypoaesthesia Multiple Sclerosis	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

DAILY

Date:03/22/99ISR Number: 3224087-7Report Type:Expedited (15-DaCompany Report #001-0945-990062

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:03/23/99ISR Number: 3224779-XReport Type:Expedited (15-DaCompany Report #034-0945-990005

Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - 400 MG (TAKEN Initial or Prolonged ONCE), PER		Anxiety Coma	Foreign Health Professional	Neurontin Capsules 400mg (Gabapentin)	PS		ORAL
ORAL		Logorrhoea		Fluvoxamine	C		

Date:03/25/99ISR Number: 3226575-6Report Type:Expedited (15-DaCompany Report #001-0945-990217

Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 100MG		Abdominal Pain Dehydration	Consumer	Neurontin Capsules 100mg (Gabapentin)	PS		
		Retching Vomiting		Corgard Estrace Pamelor	C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/99ISR Number: 3229046-6Report Type:Expedited (15-DaCompany Report #JAUSA-36661
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Bronchitis Chills Csf Protein Increased Gingival Bleeding	Health Professional	Sporanox (Itraconazole), Janssen, Capsules 100 Mg	PS	Janssen	ORAL
400 MG PULSE ORAL, 200 MG BID, ONE WEEK ON, THREE WEEKS OFF,		Influenza Like Illness Leukopenia Pyrexia Thrombocytopenia					
ORAL				Neurontin (Gabapentin)	SS		ORAL
				Ibuprofen Skelaxin	C C		

Date:03/30/99ISR Number: 3229662-1Report Type:Periodic Company Report #9818651
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Interaction Unevaluable Event	Health Professional	Diflucan Tablets Zyrtec Neurontin Tegretol	PS SS SS SS		ORAL ORAL

Date:03/30/99ISR Number: 3229952-2Report Type:Expedited (15-DaCompany Report #001-0945-990085
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG		Hypertonia Movement Disorder	Health Professional	Neurontin Capsules (Gabapentin)	PS		ORAL

(, TAKEN

ONCE), PER

ORAL

Baclofen

C

Date:03/31/99ISR Number: 3231005-4Report Type:Expedited (15-DaCompany Report #001-0945-990255

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 600 MG DAILY Initial or Prolonged PER ORAL	Balance Disorder Coordination Abnormal Depression Diplopia Dizziness Fall Fatigue Foot Fracture Speech Disorder Tremor	Consumer	Neurontin Vasotec Procardia Ritalin	PS C C C		ORAL

Date:03/31/99ISR Number: 3418509-6Report Type:Periodic

Company Report #318753

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

Outcome	PT
	Coordination Abnormal Drug Interaction

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sedation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2400 MG		Health Professional	Asacol Tablets, 400 Mg (Mesalamine)	PS		ORAL
DAILY; ORAL			Neurontin (Gabapentin)	SS		ORAL
ORAL			Neurontin (Gabapentin)	C		

Date:04/01/99ISR Number: 3231735-4Report Type:Expedited (15-DaCompany Report #044-0945-990005
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1.2 GM	C-Reactive Protein Increased	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
(DAILY) PER		Clavicle Fracture					
ORAL		Confusional State					
2 GM (DAILY)		Convulsion		Valproate Sodium	SS		ORAL
PER ORAL		Facial Bones Fracture					
YEARS AGO		Hyponatraemia					
		Malaise					
		Nausea					
		Osteoporosis					
		Urinary Tract Infection					
		Weight Increased					

Date:04/01/99ISR Number: 3231738-XReport Type:Expedited (15-DaCompany Report #001-0945-990220
 Age:6 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dermatitis	Health	Neurontin Capsules			
900 MG (300		Haemoglobin Decreased	Professional	300 Mg (Gabapentin)	PS		ORAL
MG, TID) PER		Hyperkalaemia					
ORAL		Hyponatraemia					
		Sepsis		Paxil	C		
		Stevens-Johnson Syndrome		Ritalin	C		
		White Blood Cell Count Decreased		Zantac	C		

Date:04/01/99ISR Number: 3231739-1Report Type:Expedited (15-DaCompany Report #001-0945-990120
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiac Failure	Health	Neurontin Capsules			
1800 MG (600		Congestive	Professional	300 Mg (Gabapentin)	PS		ORAL
Other		Condition Aggravated					
MG, TID), PER		Depression					
ORAL		Haematocrit Decreased		Mellaril	C		
		Haemoglobin Decreased		Clonidine	C		
		Personality Disorder		Anafranil	C		
		Post-Traumatic Stress Disorder		Robitussin	C		
		Psychotic Disorder		Peri-Colace	C		
		Red Blood Cell Count Decreased		Seroquel	C		
				Trazondone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/02/99ISR Number: 3232142-0Report Type:Expedited (15-DaCompany Report #046-0945-990003
 Age:46 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1600 MG (DAILY) PER ORAL	Anaemia Antibody Test Positive Antinuclear Antibody Positive	Foreign Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
PER ORAL	Arthralgia Blood Albumin Decreased		Sabrillex (Vigabatrin)	SS		ORAL
	Dyskinesia Glomerular Filtration Rate Decreased Oedema Peripheral Polyarthritus Spirometry Abnormal Systemic Lupus Erythematosis Tremor		Orfiril Polyfarmaci	C C		

Date:04/02/99ISR Number: 3233035-5Report Type:Direct Company Report #
 Age:57 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 100MG TID PO Intervention to Prevent Permanent Impairment/Damage	Dermatitis Pruritus		Gabapentin 100mg Cap	PS		ORAL
			Amitriptyline Fluoxetine Generic Mylanta Lansoprazole Cetirizine Amoxicillin Acetaminophen Sumatriptan Lorazeparn Buspar Fosmax Naproxen Beconase Nasal Inh	C C C C C C C C C C C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Bronchiolitis Complications Of Maternal	Health Professional	Neurontin (Gabapentin)	PS		
TRANSPLACENTAL Congenital Anomaly PLACENTAL;	SEE TEXT,	Exposure To Therapeutic Drugs					
IN UTERO EXPOSURE		Congenital Eye Disorder					
TRANSPLACENTAL PLACENTAL;	SEE TEXT,	Congenital Hearing Disorder		Gabitril (Tiagabine)	SS		
IN UTERO EXPOSURE		Congenital Tracheomalacia					
TRANSPLACENTAL PLACENTAL;	SEE TEXT,	Diarrhoea					
IN UTERO EXPOSURE		Hypotonia Nasopharyngitis		Tegretol (Carbamazepine)	SS		
TRANSPLACENTAL PLACENTAL;	SEE TEXT,	Prader-Willi Syndrome					
IN UTERO EXPOSURE		Respiratory Disorder					
TRANSPLACENTAL PLACENTAL;	SEE TEXT,	Staring					
IN UTERO EXPOSURE		Visual Disturbance Vomiting Weight Gain Poor		Multivitamins (Ergocalciferol, Ascorbic Acid, Folic			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Acid, Thiamine
Hydrochloride, C

Date:04/08/99ISR Number: 3234626-8Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blister		Gabapentin			
300MG TID		Condition Aggravated		(Neurontin)	PS		ORAL
ORAL		Dermatitis					
		Enterococcal Bacteraemia		Fluoxetine	C		
		Erythema		Methylphenidate	C		
		Escherichia Infection		Codeine	C		
		Mouth Ulceration		Morphine	C		
		Pain		Diphenhydramine	C		
		Rash Pruritic		Ranitidine	C		
		Sepsis		Acetaminophen	C		
		Staphylococcal Infection		Ceftazidime	C		
		Stevens-Johnson Syndrome		Silvadene	C		
		Systemic Candida		Nafcillin	C		
				Prednisolone	C		
				Dopamine	C		
				Amphotericin B	C		

Date:04/12/99ISR Number: 3238493-8Report Type:Direct
Age:52 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dermatitis Exfoliative	Health	Lamotrigine	PS		ORAL
50 MG PO Q		Dry Skin	Professional				
BID	8	MON					
300 MG PO Q		Rash Erythematous		Gabapentin	SS		ORAL
HS	12	DAY					
		Rash Maculo-Papular					
				Gabapentin	C		
				Haldol	C		
				Klonopin	C		
				Valproic Acid	C		

Date:04/13/99ISR Number: 3239116-4Report Type:Expedited (15-DaCompany Report #001-0945-990270
 Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Blood Glucose Increased	Consumer	Neurontin Capsules			
Initial or Prolonged	Weight Increased		300 Mg (Gabapentin)	PS		ORAL
900 MG (300						
Other						
MG TID), PER						
ORAL						

Glucotrol
Lithium

C
C

Date:04/13/99ISR Number: 3239176-0Report Type:Expedited (15-DaCompany Report #001-0945-990134
 Age:20 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Bipolar Disorder
Initial or Prolonged	Convulsion
	Dizziness
	Drug Interaction
	Drug Level Below

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Therapeutic Headache Lethargy Mania Status Epilepticus	Report Source	Product	Role	Manufacturer	Route
2700 MG (900 MG, TID), PER ORAL			Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				(Lithium)	SS		
				(Seroquel)	SS		
				Ambien (Zolpidem Tartrate	C		

Date:04/14/99ISR Number: 3239787-2Report Type:Direct
Age:51 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - PO; FOR A Initial or Prolonged SHORT TIME		Dyspnoea		Neurontin	PS		ORAL
		Fatigue					
		Lethargy		Phenobardital	C		

Date:04/14/99ISR Number: 3239814-2Report Type:Expedited (15-DaCompany Report #9912253
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 20.00 MG Intervention to TOTAL:DAILY:O Prevent Permanent RAL Impairment/Damage 1500.00 TOTAL:DAILY:O		Blood Glucose Increased	Consumer	Feldene Capsules	PS		ORAL
		Blood Triglycerides Increased					
		Haemoglobin Increased		Neurontin	SS		ORAL
		Road Traffic Accident					

RAL

Glyburide	C
Metformin	C
Lopid	C
Misoprostol	C
Tens Unit	C

Date:04/14/99ISR Number: 3240042-5Report Type:Expedited (15-DaCompany Report #001-0945-990266

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 4 TABLET (,UNK)		Ear Pain Headache Hypersensitivity Pain In Jaw	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		

Date:04/15/99ISR Number: 3240719-1Report Type:Expedited (15-DaCompany Report #001-0945-990311

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2800MG (DAILY)		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs	Health Professional	Neurontin (Gabapentin)	PS		
230MG (DAILY)				(Phenobarbital Sodium)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/15/99ISR Number: 3240722-1Report Type:Expedited (15-DaCompany Report #001-0945-990283

Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health	Neurontin			
2800MG		Chromosome Abnormality	Professional	(Gabapentin)	PS		
(DAILY)		Complications Of Maternal					
230MG,		Exposure To Therapeutic		(Phenobarbital			
(DAILY)		Drugs		Sodium)	SS		
				Ergocalciferol	C		
				Ascorbic Acid	C		
				Folic Acid	C		
				Thiamine	C		
				Hydrochloride	C		
				Retinol	C		
				Riboflavin	C		
				Nicotinamide	C		
				Panthenol	C		

Date:04/20/99ISR Number: 3243285-XReport Type:Expedited (15-DaCompany Report #001-0991-990448

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health	Rezulin			
400 MG		Drug Ineffective	Professional	(Troglitazone)	PS		ORAL
(DAILY) PER		Glycosylated Haemoglobin					
ORAL		Increased					
				Lipitor			
				(Atorvastatin)	SS		
				Neurontin			
				(Gabapentin)	SS		
				Wellbutrin	C		
				Humulin N	C		
				Neurontin	C		
				Humulin N	C		

Humulin R

C

Date:04/20/99ISR Number: 3320078-6Report Type:Periodic
Age:42 YR Gender:Male I/FU:I

Company Report #1998SUS0286

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Acne	Consumer	Sustiva	PS		
		Drug Interaction		Celexa	SS		
				Neurontin	SS		

Date:04/22/99ISR Number: 3244510-1Report Type:Direct
Age:42 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour	Consumer	Neurontin	PS		
Other		Aggression					
GRAD		Feeling Abnormal					
		Hypersensitivity					
INCREASED				Carisprodol	C		
DOSES-PROB				Diclofenac	C		
STARTED ON				Midrin	C		
1ST DAY OF							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/22/99ISR Number: 3244511-3Report Type:Direct
 Age:70 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening		Health	Baclofen			
	Encephalopathy					

PS	ORAL	10 MG PO TID 2 DAY	Product	Role
Hospitalization - 300MG MWF 10 DAY Initial or Prolonged	Multiple Sclerosis	Professional	Neurontin	SS
	Respiratory Acidosis		Synthroid	C
	Sepsis		Colace	C
			Phos-Lo	C
			Nephron Caps	C
			Cortisone	C
			Senokot	C
			Prevacid	C
			Restoril	C
			Ventolin	C
			Atrovent	C
			Calcitonin	C
			Vicodin	C

Date:04/23/99ISR Number: 3245256-6Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG QHS 8 MON	Vomiting	Health	Neurontin	PS		
		Professional				

Date:04/26/99ISR Number: 3250747-8Report Type:Periodic
 Age:76 YR Gender:Male I/FU:I

Company Report #9901371

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 50.00 MG	Drug Ineffective	Consumer	Viagra Tablets	PS		ORAL

TOTAL:PRN:ORA
 L
 TID:ORAL
 Neuropathy Peripheral Health
 Professional
 Neurontin SS ORAL
 Serevent C
 Flovent C
 Hydrocodone C
 Temazepam C

Date:04/27/99ISR Number: 3247383-6Report Type:Expedited (15-DaCompany Report #033-0945-990026
 Age:1 DY Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic	Foreign Health	Neurontin (Gabapentin)	PS		
PLACENTAL		Drugs Skull Malformation	Professional	Depakine	C		

Date:04/27/99ISR Number: 3247395-2Report Type:Expedited (15-DaCompany Report #044-0945-990036
 Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anaphylactic Reaction Chest Discomfort	Foreign Health	Neurontin (Gabapentin)	PS		
300 MG		Peak Expiratory Flow Rate	Professional				
(,DAILY),		Decreased					
UNKNOWN		Rash Erythematous		Amitriptyline Tramadol	C C		

Freedom Of Information (FOI) Report

Diclofenac C

Date:04/27/99ISR Number: 3247398-8Report Type:Expedited (15-DaCompany Report #033-0945-990024

Age:1 DY Gender: UNK, I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coarctation Of The Aorta	Foreign	(Gabapentin)	PS		
PARENTERAL Life-Threatening PLACENTAL / Congenital Anomaly IN UTERO EXPOSURE	UNK,	Complications Of Maternal Exposure To Therapeutic Drugs Heart Disease Congenital Small For Dates Baby Ultrasound Antenatal Screen Abnormal Ventricular Septal Defect Acquired	Health Professional				

Date:04/28/99ISR Number: 3248878-1Report Type:Expedited (15-DaCompany Report #001-0945-990153

Age:68 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG DAILY PER ORAL		Clonic Convulsion Convulsion Diabetes Mellitus Inadequate Control Feeling Abnormal Tremor	Health Professional	Neurontin Capsules 300 Mg (Gabapentin) Nph Insulin Insulin Zoloft Creon Furosemide Accurpil Miacalcin	PS C C C C C C C		ORAL

Date:04/29/99ISR Number: 3249873-9Report Type:Direct
Age:55 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 300MG TID; Intervention to 100MG TID Prevent Permanent Impairment/Damage		Angina Pectoris		Neurontin 300mg	PS		

Date:04/29/99ISR Number: 3249986-1Report Type:Expedited (15-DaCompany Report #001-0945-990142
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG (300 MG, QID) PER ORAL		Blood Prolactin Increased Pituitary Tumour	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

Date:04/29/99ISR Number: 3249987-3Report Type:Expedited (15-DaCompany Report #001-0945-990140
Age:55 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

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID)		Drug Interaction Hepatic Necrosis Hepatotoxicity Multi-Organ Failure	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		
				Ciclosporin	SS		
				Azathioprine	SS		
				Prednisone	SS		
				Losartan	SS		
				Atenolol	SS		
				Diltiazem	SS		
				Ranitidine	SS		
				Alprazolam	SS		
				Baclofen	SS		
				Dilaudid (Hydromorphone Hydrochloride)	SS		
				Atorvastatin	SS		
				Niacin (Nicotinic Acid)	SS		
				Ketoconazole	SS		
				Acetaminophen (Paracetamol)	SS		
				Ambien	C		
				Aspirin	C		
				Dicyclomine	C		
				Lasix	C		
				Lonox	C		
				Nizoral	C		
				Vitamin E	C		
				Niaspan	C		
				Erythromycin	C		
				Tylenol	C		

Date:04/29/99ISR Number: 3249988-5Report Type:Expedited (15-DaCompany Report #001-0945-990134
Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Bipolar Disorder	Health	Neurontin Capsules			

Initial or Prolonged 2700 MG (900 MG, TID) PER ORAL	Convulsion Dizziness Drug Interaction Headache Lethargy Status Epilepticus	Professional	300 Mg (Gabapentin)	PS	ORAL
			Eskalith (Lithium Carbonate) Seroquel Ambien	SS SS C	

Date:04/29/99ISR Number: 3250001-4Report Type:Expedited (15-DaCompany Report #001-0073-990173
Age:45 YR Gender:Female I/FU:I

Outcome Disability	PT Alopecia Anxiety Arthralgia Blood Potassium Increased Confusional State Convulsion Disorientation
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Level Below Therapeutic Fall	Consumer	Dilantin Kapseals 100 Mg (Phenytoin Sodium)	PS		ORAL
300 MG (100 MG, TID), PER ORAL		Gingival Hyperplasia Nausea Palpitations Parkinsonian Gait Tinnitus Visual Acuity Reduced		Neurontin (Gabapentin) Prempro	SS C		

Date:05/03/99ISR Number: 3251651-1Report Type:Direct
Age:41 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Coordination Abnormal Dysarthria Gait Disturbance	Health Professional	Gabapentin Morphine Trazodone Fluoxetine	PS SS C C		

Date:05/04/99ISR Number: 3253604-6Report Type:Expedited (15-DaCompany Report #001-0945-990361
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cough Groin Pain	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG, TID) PER ORAL		Headache Myocardial Infarction Nausea Vomiting		Daypro	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Optic Neuritis Retinopathy	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1200 MG (600 MG BID) PER ORAL							
				Elavil	C		
				Betaseron	C		
				Copaxone	C		
				Soma	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Glucose Decreased Coma	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
Required ONE DOSE, PER Intervention to ORAL Prevent Permanent Impairment/Damage							
				(Insulin)	C		
				Lescol (Fluvastatin Sodium)	C		
				(Atenolol)	C		
				Vasotec (Enalapril			

Freedom Of Information (FOI) Report

Maleate) C
 Lasix (Furosemide) C
 Aspirin
 (Acetylsalicylic
 Acid) C

Date:05/06/99ISR Number: 3255699-2Report Type:Expedited (15-DaCompany Report #001-0981-992736
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Lipitor (Atorvastatin)	PS		ORAL
				Neurontin (Gabapentin)	SS		
				Tegretol (Carbamazepine)	SS		

Date:05/07/99ISR Number: 3256537-4Report Type:Expedited (15-DaCompany Report #20615-009
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abdominal Distension	Health Professional	Duraclon Injection (Clonidine Hydrochloride),			
Life-Threatening Required		Agitation		1mg/10ml, Roxane Laboratories, Inc.	PS	Roxane Laboratories, Inc.	
Intervention to Prevent Permanent Impairment/Damage		Apnoea					
3-9 MCG/HR IT		Blood Pressure Increased					
		Coma					
		Condition Aggravated					
		Emotional Distress		Gabapentin 100 Mg Capsules-			
		Failure Of Implant		Parke-Davis	SS	Parke-Davis	ORAL
		Pain					
		Tachycardia					
		Tonic Convulsion					
		Urinary Retention					
200 MG Q8H PO							

Date:05/07/99ISR Number: 3261871-8Report Type:Periodic
 Age:42 YR Gender:Female I/FU:I

Company Report #98USA10702

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG, TID, ORAL		Rash Maculo-Papular	Health Professional	Tegretol Tablet (Carbamazepine)	PS		ORAL
2 MG, DAILY, ORAL				Klonopin Unknown (Clonazepam)	SS		ORAL
SEE IMAGE				Neurontin Capsule (Gabapetin)	SS		ORAL
				Depakote Ganciclovir Solution	C C		

Date:05/11/99ISR Number: 3259107-7Report Type:Expedited (15-DaCompany Report #001-0945-990125
Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3 MON Other		Bile Duct Obstruction Hepatic Enzyme Increased Jaundice	Health Professional	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/11/99ISR Number: 3259108-9Report Type:Expedited (15-DaCompany Report #001-0945-990255

Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Balance Disorder	Consumer	Neurontin			
Hospitalization -		Coma		(Gabapentin)	PS		ORAL
600 MG							
Initial or Prolonged		Coordination Abnormal					
(DAILY), PER							
ORAL		Depression					
		Diplopia		Vasotec	C		
		Dizziness		Procardia	C		
		Drug Toxicity		Ritalin	C		
		Fall					
		Fatigue					
		Foot Fracture					
		Speech Disorder					
		Tremor					

Date:05/11/99ISR Number: 3259109-0Report Type:Expedited (15-DaCompany Report #033-0945-980026

Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Fall	Foreign	Neurontin			
Hospitalization -		Femoral Neck Fracture	Health	(Gabapentin)	PS		ORAL
Initial or Prolonged							
900 MG (300							
MG, TID), PER		Muscular Weakness	Professional				
ORAL				Lioresal	C		

Date:05/12/99ISR Number: 3260114-9Report Type:Expedited (15-DaCompany Report #20615-009

Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abdominal Distension	Health	Duraclon Inection			
Life-Threatening		Agitation	Professional	(Clonidine			
Required		Apnoea		Hydrochloride),			

Intervention to Prevent Permanent Impairment/Damage 3-9 MCG/HR IT	Blood Pressure Increased Clonic Convulsion Coma	1mg/10ml, Roxane Laboratories, Inc.	PS	Roxane Laboratories, Inc.	
	Emotional Distress Mental Disorder Pain	Gabapentin 100 Mg Capsules - Parke Davids	SS	Parke-Davids	ORAL
200MG Q8H PO	Tachycardia Urinary Retention				

Date:05/14/99ISR Number: 3262841-6Report Type:Expedited (15-DaCompany Report #001-0945-990101
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Exposure Drowning	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 MG (X1)							
PER ORAL				Lamictal (Lamotrigine)	SS		
				Dilantin	C		
				Drisdol	C		
				Calcium Carbonate	C		
				Docusate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/17/99ISR Number: 3263532-8Report Type:Expedited (15-DaCompany Report #001-0945-990371
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Optic Ischaemic Neuropathy	Health Professional	Neurontin (Gabapentin)	PS		
180 (600 MG							
TID)							

Date:05/19/99ISR Number: 3265131-0Report Type:Expedited (15-DaCompany Report #001-0945-990120
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Cardiac Failure Congestive	Health Professional	Neurontin Capsules 300mg(Gabapentin)	PS		ORAL
1800 MG (600							
Other		Depression					
MG, TID), PER							
ORAL		Haematocrit Decreased					
		Haemoglobin Decreased		Mellaril(Thioridazine Hydrochloride)	C		
		Major Depression		(Clonidine)	C		
		Personality Disorder		Anafranil(Clomipramine Hydrochloride)	C		
		Post-Traumatic Stress Disorder		Robitussin(Guaifenesin)	C		
		Psychotic Disorder		Peri-Colace(Docusate Sodium, Casanthranol)	C		
		Red Blood Cell Count Decreased		Seroquel (Quetiapine Fumarate)	C		
				(Trazodone)	C		

Date:05/19/99ISR Number: 3330102-2Report Type:Periodic Company Report #S99-USA-00258-01
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

20 MG DAILY	DERMATITIS	CONSUMER	CELEXA	PS	ORAL
PO	DRUG INTERACTION				
30 MG DAILY			CELEXA	SS	ORAL
PO			SUSTIVA	SS	
			NEURONTIN	SS	

Date:05/20/99ISR Number: 3265117-6Report Type:Expedited (15-DaCompany Report #99-05-0121
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Encephalopathy	Health	Baclofen Tablets	PS		ORAL
10MG TID ORAL							
Hospitalization -		Multiple Sclerosis	Professional	Neurontin Tablets	SS		ORAL
300MG MWF							
Initial or Prolonged		Respiratory Acidosis					
ORAL		Sepsis		Synthroid	C		
				Colace	C		
				Nephrocaps	C		
				Cortisone	C		
				Senokot	C		
				Prevacid	C		
				Restoril	C		
				Ventolin	C		
				Vicodin	C		
				Atrovent	C		
				Calcitonin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/20/99ISR Number: 3266113-5Report Type:Expedited (15-DaCompany Report #001-0945-990424

Age:27 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800MG (600MG TID) , PER ORAL	Breast Mass Eosinophil Count Increased Fatigue	Consumer	Neurontin Capsules 300mg (Gabapentin)	PS		ORAL
	Haematoma Malaise Pyrexia White Blood Cell Count Increased		Wellbutrin Desyrel	C C		

Date:05/21/99ISR Number: 3268104-7Report Type:Expedited (15-DaCompany Report #001-0945-990363

Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged (SEE TEXT) 900 MG TWICE A DAY , 600 MG AT BEDTIME1500	Bone Cyst Increased Tendency To Bruise	Health Professional	Neurontin (Gabapentin)	PS		
			Prozac Unspecified Tricyclic Antidepressant	C C		

Date:05/25/99ISR Number: 3269477-1Report Type:Expedited (15-DaCompany Report #001-0945-990170

Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 500 MG(,300 MG QAM AND 200 MG QPM), PER ORAL		Graft Versus Host Disease Medication Error	Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

Date:05/25/99ISR Number: 3269481-3Report Type:Expedited (15-DaCompany Report #001-0073-990173
Age:45 YR Gender:Female I/FU:F

Outcome	PT
Disability	Abdominal Distension Abdominal Pain Alopecia Anxiety Arthralgia Blood Potassium Increased Condition Aggravated Confusional State Convulsion Drug Level Below Therapeutic Fall Gingival Hypertrophy Heart Rate Irregular Nausea Parkinsonian Gait

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID), PER ORAL		Tinnitus Visual Acuity Reduced Weight Increased	Consumer	Dilantin Kapseal 100 Mg (Phenytoin Sodium)	PS		ORAL
				Neurontin (Gabapentin)	SS		
				Prempro (Medroxyprogesterone Acetate, Estrogens Conjugated)	C		

Date: 05/26/99
 ISR Number: 3270270-4
 Report Type: Expedited (15-DaCompany Report #001-0945-990413)
 Age: 79 YR
 Gender: Female
 I/FU: I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1600 MG (400 MG, QID), PER ORAL		Alopecia Asthenia Deafness Decreased Appetite	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
		Depression Dermatitis Dizziness Eructation Flatulence Hypersensitivity Hypoaesthesia Mucosal Ulceration Paraparesis Scab Skin Exfoliation Tremor Ulcer Visual Disturbance		Spinal Epidural	C		

Vulvovaginal Discomfort

Date:05/27/99ISR Number: 3270198-XReport Type:Expedited (15-DaCompany Report #001-0945-990442
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 6400 MG (1600 MG,QID)		Accident Complex Partial Seizures Drug Ineffective Injury Overdose	Consumer	Neurontin (Gabapentin)	PS		
				Dilantin Experimental Seizure Medication	C C		

Date:05/27/99ISR Number: 3270208-XReport Type:Expedited (15-DaCompany Report #001-0945-990412
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 MG TID) PER		Drug Interaction International Normalised Ratio Increased	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

PER ORAL 7.5	Coumadin (Warfarin Sodium)	SS	ORAL
MG	Premarin	C	
	Compazine	C	
	Aciclovir	C	
	Dexamethasone	C	
	Cisplatin	C	
	Bcnu	C	
	Anzemet	C	

Date:05/27/99ISR Number: 3271183-4Report Type:Expedited (15-DaCompany Report #001-0945-990419
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abortion Spontaneous	Health Professional	Neurontin (Gabapentin)	PS		
Other							
600 MG,							
(DAILY),							
(06/ /98 -							
PREGNANCY							
WEEK 5)							

Paxil (Paroxetine Hydrochloride)	C
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Date:05/27/99ISR Number: 3271184-6Report Type:Expedited (15-DaCompany Report #001-0945-990421
 Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Back Pain Encephalopathy	Health Professional	Neurontin (Gabapentin)	PS		
300 MG ON							
MON, WED &		Liver Function Test					

FRI, UNKNOWN	Abnormal			
30 MG (10 MG, TID) PER ORAL	Mental Impairment	(Baclofen)	SS	ORAL
	Respiratory Acidosis			
	Sepsis	Colace (Docusate Sodium)	C	
		Phoslo (Calcium Acetate)	C	
		(Cortisone)	C	
		Nephrocaps (Folic Acid, Vitamins Nos)	C	
		Prevacid (Lansoprazole)	C	
		Restoril (Temazepam)	C	
		(Calcitonin)	C	
		Vicodin (Paracetamol, Hydrocodone Bitartrate)	C	
		Senokot (Senna Fruit)	C	
		Ventolin (Salbutamol)	C	
		Atrovent (Ipratropium Bromide)	C	
		Synthroid (Levothyroxine Sodium)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Cordarone
(Amiodarone
Hydrochloride) C

Date:05/27/99ISR Number: 3271186-XReport Type:Expedited (15-DaCompany Report #JACGBR1999000063
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hypokalaemia	Foreign Health Professional	Hismanal (Unspecified) (Astemizole) Neurontin (Gabapentin)	PS SS		

Date:05/28/99ISR Number: 3271605-9Report Type:Expedited (15-DaCompany Report #002-0945-990014
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 500 MG (DAILY)	4 WK	Anhedonia Catatonia Decreased Appetite Depressed Mood Depression Disturbance In Attention Hypomania Immobile Insomnia Lack Of Spontaneous Speech Muscle Rigidity Staring Tangentiality Tearfulness Thinking Abnormal	Foreign Literature Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:05/28/99ISR Number: 3272789-9Report Type:Expedited (15-DaCompany Report #001-0945-990418
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 MG, TID) PER ORAL		Drug Interaction Drug Level Above Therapeutic Dry Mouth	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
1800 MG (600 MG, TID), PER ORAL		Dyskinesia Parkinsonian Gait Sedation Vision Blurred		(Lithium) Ambien	SS C		ORAL

Date:06/01/99ISR Number: 3274447-3Report Type:Expedited (15-DaCompany Report #12347/20246
Age:30 YR Gender:Female I/FU:I

Outcome	PT
Disability	Diplopia
Other	Dizziness Hysterectomy Iiird Nerve Paralysis 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
		Migraine Mydriasis Nausea					
		Ovarian Cyst Paraesthesia Peripheral Nerve Injury	Health Professional Company	Depo-Provera Contraceptive Injection (150 Mg)	PS		
IM		Sedation Sensation Of Foreign Body Visual Disturbance	Representative	Neurontin Tylenol	SS SS		

Date:06/02/99ISR Number: 3274552-1Report Type:Expedited (15-DaCompany Report #99-05-0121
Age:70 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	10MG TID	ORAL	Encephalopathy	Health	Baclofen Tablets	PS		ORAL
Hospitalization - Initial or Prolonged ORAL	300MG MWF		Mental Impairment Respiratory Acidosis Sepsis	Professional	Neurontin Tablets	SS		ORAL
					Synthroid	C		
					Colace	C		
					Nephrocaps	C		
					Cortisone	C		
					Senokot	C		
					Prevacid	C		
					Restoril	C		
					Ventolin	C		
					Vicodin	C		
					Atrovent	C		
					Calcitonin	C		

Date:06/02/99ISR Number: 3279032-5Report Type:Expedited (15-DaCompany Report #001-0945-990125
Age:75 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Bile Duct Obstruction	Health	Neurontin			
Hospitalization - 3 MON			Bile Duct Stenosis	Professional	(Gabapentin)	PS		

Initial or Prolonged Hepatic Enzyme Increased
Other Jaundice

Date:06/02/99ISR Number: 3283040-8Report Type:Periodic Company Report #98USA10702
Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG, DAILY, ORAL	Rash Maculo-Papular	Health Professional	Tegretol Tablet (Carbamazepine)	PS		ORAL
1800 MG, DAILY, ORAL			Neurontin Capsule (Gabapentin)	SS		ORAL
			Depakote Capsule	C		
			Ganciclovir Solution	C		
			Glipizide Tablet	C		
			Phenobarbital	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/03/99ISR Number: 3276035-1Report Type:Expedited (15-DaCompany Report #001-0981-992736
 Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health Professional	Lipitor (Atorvastatin)	PS		ORAL
Other				Neurontin (Gabapentin)	SS		
PER ORAL				Tegretol (Carbamazepine)	SS		

Date:06/08/99ISR Number: 3277497-6Report Type:Direct Company Report #
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sexual Dysfunction		Gabapentin 300 Mg Cap	PS		
300 MG QID							

Date:06/08/99ISR Number: 3277925-6Report Type:Expedited (15-DaCompany Report #001-0945-990471
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abortion Spontaneous	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
Other		Blood Gonadotrophin					
2400 MG,		Increased					
DAILY, PER				Lamictal	C		
ORAL							

Date:06/09/99ISR Number: 3279012-XReport Type:Direct Company Report #
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -	Accident At Home	Baclofen	PS	ORAL
25MG PO Q 6H				
Initial or Prolonged	Fall	Gabapentin	SS	
300MG PO TID				
	Sedation			

Date:06/10/99ISR Number: 3280402-XReport Type:Expedited (15-DaCompany Report #049-0945-990013
 Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bacterial Infection Dermatitis	Foreign Health	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
200 MG		Sepsis	Professional				
(,DAILY), PER		Toxic Epidermal					
ORAL		Necrolysis		Phenhydan (Phenytoin)	C		
				Novodigal (Digoxin)	C		
				L-Thyroxin (Levothyroxine Sodium)	C		

Date:06/16/99ISR Number: 3284976-4Report Type:Expedited (15-DaCompany Report #001-0945-990491
 Age:88 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

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID), PER ORAL		Atrial Fibrillation Confusional State Delirium Delusion Paranoia Urinary Tract Infection	Health Professional Company Representative	Neurontin Capsules 100 Mg(Gabapentin) Lasix (Furosemide) Coumadin (Warfarin Sodium) (Potassium) (Digoxin) Augmentin(Clavulanate Potassium, Amoxicillin Trihydrate)	PS C C C C C		ORAL

Date:06/16/99ISR Number: 3286392-8Report Type:Expedited (15-DaCompany Report #9925406
Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 200.00MG Intervention to Prevent Permanent Impairment/Damage 300.00MG TOTAL ORAL		Abnormal Behaviour Condition Aggravated Drug Interaction Sleep Disorder Suicidal Ideation	Health Professional	Zoloft Tablets Neurontin	PS SS		ORAL ORAL

Date:06/17/99ISR Number: 3286241-8Report Type:Expedited (15-DaCompany Report #LBID002990002
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Agitation	Consumer	Lithobid Tablets 300			

Initial or Prolonged	Drug Level Above Therapeutic	Mg (Lithium Carbonate)	PS	ORAL
1200 MG, PER				
ORAL	Drug Toxicity			
	Feeling Drunk	Benzotropine (Benzatropine Mesilate)	SS	ORAL
PER ORAL	Nausea			
	Tremor	Neurontin (Gabapentin)	SS	ORAL
PER ORAL				
		Ambien (Zolpidem Tartrate)	SS	ORAL
PER ORAL				
		Buspar	C	
		Lamictal	C	
		Wellbutrin	C	
		Zoloft	C	
		Zyprexa	C	

Date:06/17/99ISR Number: 3286266-2Report Type:Expedited (15-DaCompany Report #A0094748
Age:31 YR Gender:Male I/FU:I

Outcome	PT
Life-Threatening	Aggression
Hospitalization -	Aspiration
Initial or Prolonged	Electrocardiogram Qrs
Other	Complex Prolonged
	Intentional Misuse
	Lung Infiltration

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Suicide Attempt							
Tachypnoea							
Ventricular Tachycardia			Health	Lamictal Tablet	PS		ORAL
Vomiting			Professional	Cabapentin Capsule	SS		ORAL
				Fluoxetine Hydrochloride	SS		ORAL
				Risperidone	SS		ORAL
				Methylphenidate	SS		ORAL
				Thioridazine	SS		
				Ethanol	SS		ORAL

Date:06/21/99ISR Number: 3288053-8Report Type:Expedited (15-DaCompany Report #001-0945-990505
 Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hallucination	Consumer	Neurontin (Gabapentin)	PS		ORAL
3600 MG (2400 Other QAM), 1200 MG QPM) PER ORAL		Logorrhoea					
		Multiple Sclerosis					
				Many Unspecified Medications	C		

Date:06/21/99ISR Number: 3288331-2Report Type:Expedited (15-DaCompany Report #049-0945-990013
 Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dermatitis	Foreign Health	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
200 MG (DAILY) PER ORAL		Epidermolysis Bullosa	Professional				
		Liver Function Test					
		Abnormal					
		Sepsis		Phenhydan	C		

Toxic Epidermal
Necrolysis

Novodigal
L-Thyroxin

C
C

Date:06/21/99ISR Number: 3288377-4Report Type:Expedited (15-DaCompany Report #001-0945-990371
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Optic Ischaemic Neuropathy	Health Professional	Neurontin (Gabapentin)	PS		
1800 MG (600 MG, TID)		Visual Disturbance					

Date:06/22/99ISR Number: 3287414-0Report Type:Direct Company Report #
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema		Neurontin	PS		ORAL
300 MG ONCE PO				Quinapril Hydrochloride	C		
				Hydrochlorothiazide	C		
				Atenolol	C		
				Valproic Acid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/22/99ISR Number: 3288919-9Report Type:Expedited (15-DaCompany Report #001-0945-990508
 Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		International Normalised Ratio Increased	Health Professional	Neurontin (Gabapentin)	PS		
200 MG (AFTER EACH DIALYSIS TREATMENT)		Prothrombin Time Prolonged					
6 MG (AFTER EACH DIALYSIS TREATMENT)				Coumadin (Warfarin Sodium)	SS		
				Erythropoietin	C		
				Insulin	C		
				Cozaar	C		
				Glipizide	C		

Date:06/22/99ISR Number: 3288921-7Report Type:Expedited (15-DaCompany Report #001-0945-990506
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pulmonary Fibrosis	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:06/24/99ISR Number: 3290887-0Report Type:Expedited (15-DaCompany Report #001-0945-990547
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complications Of Maternal Exposure To Therapeutic	Health Professional	Neurontin (Gabapentin)	PS		
2700 MG,							

DAILY

Drugs

Pre-Eclampsia
Premature Baby

Cyclosporine
(Ciclosporin)

C

Date:06/25/99ISR Number: 3296730-8Report Type:Periodic
Age:32 YR Gender:Male I/FU:I

Company Report #9907622

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Alanine Aminotransferase Increased Diabetes Mellitus	Health Professional	Glucotrol Xl Extended Release Tablets	PS		ORAL
10.00 MG		Intentional Misuse					
TOTAL: BID:ORA		Liver Function Test					
L		Abnormal		Gabapentin	SS		ORAL
1200.00 MG		Medication Error					
TOTAL: DAILY:O		Neuralgia					
RAL				Metformin	C		
				Naprosyn	C		
				Multi-Vitamin	C		
				Insulin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/02/99ISR Number: 3342881-9Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0981-992725

Outcome Dose	Duration	PT Drug Ineffective Drug Interaction	Report Source Consumer	Product	Role	Manufacturer	Route
60 MG (20 MG, TID)				Lipitor Tablets 20 Mg (Atorvastatin)	PS		ORAL
				Zestril (Lisinopril)	SS		
				Betapace (Sotalol Hydrochloride)	SS		
				(Salsalate)	SS		
				Cardura (Doxazosin Mesilate)	SS		
				Neurontin (Gabapentin)	SS		
				Plavix (Clopidogrel)	SS		
				K-Dur (Potassium Chloride)	SS		
				(Furosemide)	SS		
				Z-Bec (Zinc Sulfate, Vitamins Nos)	SS		
				Oscal (Calcium Carbonate)	SS		
				(Temazepam)	SS		
				(Ranitidine)	SS		
				Lanoxin (Digoxin)	SS		
				(Amitriptyline)	SS		
				Vitamin E (Tocopherol)	SS		
				Synthroid (Levothyroxine Sodium)	SS		
				Propoxyphene-N (Dextropropoxyphene)	SS		
				Hytrin (Terazosin Hydrochloride)	C		

Date:07/02/99ISR Number: 3351008-9Report Type:Periodic
 Age:78 YR Gender:Female I/FU:I

Company Report #001-0981-992255

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Chills	Consumer	Lipitor	
Diarrhoea	Health	(Atorvastatin)	PS
Malaise	Professional	Neurontin	
Nausea		(Gabapentin)	SS

Date:07/06/99ISR Number: 3298103-0Report Type:Expedited (15-DaCompany Report #049-0945-990013

Age:75 YR Gender:Female I/FU:F

Outcome

PT

Death

Blister

Blood Urea Increased

Cardiac Failure

Conjunctivitis

Dermatitis

Epidermolysis Bullosa

Hyperkalaemia

Leukocytosis

Liver Function Test

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	
600 MG (300 MG, BID), PER ORAL		Mouth Ulceration	Foreign Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL	
		Oral Pain						
		Pyrexia						
		Sedation						
		Sepsis						
		Toxic Epidermal Necrolysis						
		Urinary Tract Infection						
		Phenhydan						C
		Novodigal						C
		L-Thyroxin						C
		Antra						C
		Aspirin						C
		Bronchoretard						C
Zinacef	C							
Catapresan	C							
Paracetamol	C							
Berotec	C							
Pulmicort	C							
Rekawan	C							

Date:07/06/99ISR Number: 3298522-2Report Type:Expedited (15-DaCompany Report #001-0945-990546

Age:1 DY Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Collapse Of Lung Complications Of Maternal	Health Professional	Neurontin (Gabapentin)	PS		
Other							
PLACENTAL	IN UTERO	Exposure To Therapeutic					
EXPOSURE							
PLACENTAL	IN UTERO	Premature Baby		Topamax (Topiramate)	SS		
EXPOSURE							
PLACENTAL	IN UTERO			Felbatol (Felbamate)	SS		
EXPOSURE							

IN UTERO

EXPOSURE

(Phenobarbital) SS

PLACENTAL

IN UTERO

EXPOSURE

(Folic Acid) C
Prenatal Vitamins
(Ergocalciferol,
Acorbic Acid,
Pyridoxine
Hydrochloride, C

Date:07/06/99ISR Number: 3298524-6Report Type:Expedited (15-DaCompany Report #001-0945-990412

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 MG, TID), PER		Drug Interaction International Normalised Ratio Increased	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
ORAL 1800 MG (600 MG, TID), PER				Coumadin (Warfarin Sodium)	SS		ORAL
ORAL				Premarin (Estrogens Conjugated) (Aciclovir)	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Campazine
 (Prochlorperazine
 Edisylate) C
 (Dexamethasone) C
 (Cisplatin) C
 Bcnu (Carmustine) C
 Anzemet (Dolasetron
 Mesilate) C

Date:07/06/99ISR Number: 3298528-3Report Type:Expedited (15-DaCompany Report #001-0945-990592
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health	Neurontin			
		Complications Of Maternal	Professional	(Gabapentin)	PS		
		Exposure To Therapeutic		Felbatol (Felbamate)	SS		
		Drugs					

Date:07/07/99ISR Number: 3298550-7Report Type:Expedited (15-DaCompany Report #044-0945-990044
 Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cachexia	Foreign	Neurontin			
Initial or Prolonged		Cervical Myelopathy	Health	(Gabapentin)	PS		ORAL
1200 MG							
(,QD), PER		Csf Protein Increased	Professional				
ORAL		Nuclear Magnetic					
		Resonance Imaging		Lamotrigine	C		
		Abnormal		Carbamazepine	C		

Date:07/09/99ISR Number: 3299725-3Report Type:Direct Company Report #
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Eye Rolling		Prilosec #3			
Intervention to		Joint Stiffness		Tegretol Susp 100mg			

Prevent Permanent Movement Disorder
8 MON
Impairment/Damage

/5cc PS
Neurontin Capsules SS
Tegretol Susp 100mg
/5cc SS
Eyedrops C

Date:07/09/99ISR Number: 3300470-6Report Type:Expedited (15-DaCompany Report #001-0945-990619
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (, DAILY) , PER ORAL		Abdominal Pain Hepatitis Liver Function Test Abnormal	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Paxil	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/09/99ISR Number: 3300471-8Report Type:Expedited (15-DaCompany Report #001-0945-990421
 Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300 MG ON	Alanine Aminotransferase Increased	Health Professional	Neurontin (Gabapentin)	PS		
	MON, WED, & FRI	Aspartate Aminotransferase Increased		(Baclofen)	SS		ORAL
	30 MG (10 MG, TID), PER ORAL	Encephalopathy					
		Mental Impairment					
		Respiratory Acidosis		Colace	C		
		Sepsis		Phoslo	C		
				Cortisone	C		
				Nephrocaps	C		
				Prevacid	C		
				Restoril	C		
				(Calcitonin)	C		
				Vicodin	C		
				Senokot	C		
				Ventolin	C		
				Atrovent	C		
				Synthroid	C		
				Cordarone	C		

Date:07/12/99ISR Number: 3301817-7Report Type:Expedited (15-DaCompany Report #9912253
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required	20.00 MG	Arthritis	Consumer	Feldene Capsules	PS		ORAL
Intervention to	TOTAL:DAILY:0	Blood Glucose Decreased	Health				
Prevent Permanent	RAL	Blood Glucose Increased	Professional				
Impairment/Damage	900.00 MG	Blood Triglycerides	Company	Neurontin	SS		ORAL

Increased

Representative

Diabetes Mellitus

Non-Insulin-Dependent

Tens Unit

C

Haemoglobin Increased

Hyperlipidaemia

Irritable Bowel Syndrome

Low Density Lipoprotein

Increased

Muscle Spasms

Musculoskeletal Disorder

Pain

Peptic Ulcer

Polytraumatism

Road Traffic Accident

Date:07/13/99ISR Number: 3301729-9Report Type:Direct

Company Report #081177

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Noroxin 400mg (Norfloxacin)	PS	Msd	
				Neurontin 400mg (Gabapentin)	SS	Parke Davis	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/13/99ISR Number: 3302478-3Report Type:Expedited (15-DaCompany Report #001-0945-990363

Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged SEE TEXT	Bone Cyst Ecchymosis	Health Professional	Neurontin (Gabapentin)	PS		
			Prozac Unspecified Tricyclic Antidepressant	C C		

Date:07/14/99ISR Number: 3303992-7Report Type:Expedited (15-DaCompany Report #A0094748

Age:31 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening SINGLE DOSE Hospitalization - ORAL Initial or Prolonged SINGLE DOSE Other ORAL	Aggression Aspiration Electrocardiogram Qrs Complex Prolonged Intentional Misuse Lung Infiltration Suicide Attempt Tachypnoea	Health Professional	Lamictal Tablet	PS		ORAL
	Ventricular Tachycardia Vomiting		Gabapentin Capsule	SS		ORAL
			Fluoxetine Hydrochloride (Formulation Unknown)	SS		ORAL
			Risperidone (Formulation Unknown)	SS		ORAL
			Methylphenidate (Formulation Unknown)	SS		ORAL
			Thioridazine (Formulation Unknown)	SS		ORAL
			Ethanol (Formulation			

ORAL

Unknown) SS ORAL

Date:07/14/99ISR Number: 3304026-0Report Type:Expedited (15-DaCompany Report #A0094619
 Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Aggression	Health	Lamictal Tablet	PS		ORAL
ORAL							
Initial or Prolonged		Aspiration	Professional	Gabapentin	SS		ORAL
SINGLE DOSE/							
Other		Electrocardiogram Qrs					
ORAL							
		Complex Prolonged		Fluoxetine			
		Intentional Misuse		Hydrochloride	SS		
		Lung Infiltration		Risperidone	SS		
		Suicide Attempt		Thioridazine	SS		
		Ventricular Tachycardia		Methylphenidate	SS		
		Vomiting		Ethanol	SS		

Date:07/15/99ISR Number: 3347794-4Report Type:Periodic Company Report #990518-SK796
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200.000MG QD		Headache	Consumer	Celebrex	PS		ORAL
PO		Pain	Health				
5.000MG QD PO		Sedation	Professional	Fosamax	SS		ORAL
1500.000MG QD		Tooth Disorder		Relafen (Usa)	SS		ORAL
PO							
2400.000MG QD				Neurontin	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

PO
 1.000 TB QD
 Premarin SS ORAL

PO
 Medroxyprogesterone C
 Flunisolide C
 Metronidazole C

Date:07/15/99ISR Number: 3349889-8Report Type:Periodic Company Report #990415-SK377
 Age:84 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Myalgia	Consumer	Celebrex	PS		ORAL
QD	PO	3 DAY		Health	Neurontin	SS		ORAL
PO				Professional				

Date:07/15/99ISR Number: 3350821-1Report Type:Periodic Company Report #990426-SK376
 Age:71 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Sedation	Consumer	Celebrex	PS		ORAL
200.000 MG QD				Health				
PO				Professional	Neurontin	SS		ORAL
PO					Soma	SS		ORAL
PO					Trazodone	SS		ORAL
PO					Buspirone	C		
					Clonazepam	C		
					Lansoprazole	C		

Date:07/16/99ISR Number: 3305704-XReport Type:Expedited (15-DaCompany Report #1324/11153
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 160 MG/DAY; Initial or Prolonged		Thrombocytopenia	Foreign	Medrol Tablets	PS		ORAL
ORAL			Consumer				
Other ORAL			Company	Nifedipine	SS		ORAL
ORAL			Representative	Neurontin	SS		ORAL
400 MG-6Q1DY; ORAL				Di-Antalvic (400 Mg)	SS		ORAL
				Vincristine Sulphate	C		
				Carmustine	C		
				Procarbazine	C		
				Cisplatin	C		
				Cytarabine	C		
				Dacarbazine	C		
				Hydroxycarbazine	C		

Date:07/21/99ISR Number: 3308028-XReport Type:Direct
Age:39 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 300MG QID PO Hospitalization - 1200MG QID PO Initial or Prolonged		Aplasia		Carbamazepine	PS		ORAL
		Glossopharyngeal		Gabapentin	SS		ORAL
		Neuralgia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/22/99ISR Number: 3309025-0Report Type:Expedited (15-DaCompany Report #PRIUSA1999002537
 Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Pressure Systolic	Literature	Tylenol With Codeine			
Hospitalization -		Decreased	Health	(Unspecified)			
Initial or Prolonged		Coma	Professional	(Acetaminophen/Codei			
ORAL		Completed Suicide		ne)	PS		ORAL
ORAL		Convulsion		Amitriptyline			
ORAL		Fluid Overload		(Amitriptyline)	SS		ORAL
ORAL		Heart Rate Increased		Valium (Diazepam)	SS		ORAL
OROPHARINGEAL	ORAL	Hypotension		Gabapentin			
		Hypoxia		(Gabapentin)	SS		
		Oxygen Saturation					
		Abnormal					
		Pulmonary Oedema					
		Pupil Fixed					
		Ventricular Tachycardia					

Date:07/22/99ISR Number: 3309155-3Report Type:Expedited (15-DaCompany Report #044-0945-990075
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Drug Interaction	Foreign	Neurontin			
Initial or Prolonged		International Normalised	Health	(Gabapentin)			
300 MG		Ratio Increased	Professional				
DAILY	PER						
ORAL				Warfarin	SS		ORAL
VARIABLE							
PER ORAL				Ciprofloxacin	C		
				Fentanyl	C		
				Co-Amilozide	C		
				Sevredol	C		

Date:07/22/99ISR Number: 3309156-5Report Type:Expedited (15-DaCompany Report #001-0945-990632
Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anal Fissure	Consumer	Neurontin Capsules			
		Colitis Ulcerative	Health	300 Mg (Gabapentin)	PS		ORAL
900 MG (300		Condition Aggravated	Professional				
MG TID)		Enterocolitis					
PER ORAL		Haemorrhagic		Lamictal	C		
		Rectal Haemorrhage					

Date:07/22/99ISR Number: 3309267-4Report Type:Expedited (15-DaCompany Report #10048908
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Creatinine	Study	Omapatrilat	PS		ORAL
40 MILLIGRAM,		Increased	Health				
Initial or Prolonged		Blood Urea Increased	Professional	Neurontin			
2/DAY ORAL		Cardiac Failure		(Gabapentin)	SS		
		Congestive		Vitamins + Minerals	C		
		Condition Aggravated		Lanoxin	C		
		Renal Impairment		Furosemide	C		
				Zaroxolyn	C		
				Quinine	C		
				Isosorbide Dinitrate	C		

Freedom Of Information (FOI) Report

Amlodipine	C
Premarin	C
Alprazolam	C
Aspirin	C
Nph Insulin	C
Insulin Sliding	
Scale	C
Sublingual	
Nitroglycerin	C
Tylenol	C
Cimetidine	C
Allopurinol	C
Famotidine	C

Date:07/22/99ISR Number: 3309750-1Report Type:Expedited (15-DaCompany Report #044-0945-990080
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Congenital Anomaly	Abortion Induced Anal Atresia	Foreign Health	Neurontin (Gabapentin)	PS		
2900 MG (,DAILY), PLACENTAL; IN UTERO EXPOSURE	Complications Of Maternal Exposure To Therapeutic Drugs Congenital Anomaly Congenital Diaphragmatic Eventration Ectropion Exomphalos Kyphoscoliosis Limb Deformity Male Genital Tract Tuberculosis Pulmonary Hypoplasia Talipes	Professional	Lamotrigine Clobazam	C C		

Date:07/22/99ISR Number: 3309754-9Report Type:Expedited (15-DaCompany Report #001-0945-990626
 Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health	Neurontin Capsules			
Hospitalization - SEE TEXT, PER Initial or Prolonged ORAL		Blood Creatinine Increased Hypotension	Professional	300 Mg (Gabapentin)	PS		ORAL

Date:07/22/99ISR Number: 3309757-4Report Type:Expedited (15-DaCompany Report #044-0945-990079
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
2900 MG (,DAILY), PER ORAL			Professional				
				Lamotrigine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/22/99ISR Number: 3309938-XReport Type:Direct
 Age:43 YR Gender:Female I/FU:I

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		White Blood Cell Count		Gabapentin	PS		ORAL
300MG QAM PO;		Decreased					
600MG QHS PO							

Date:07/23/99ISR Number: 3309909-3Report Type:Expedited (15-DaCompany Report #001-0945-990677
 Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiac Failure Congestive	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 Other MG, TID), PER		Influenza Like Illness					
ORAL		International Normalised Ratio Increased		Coumadin (Warfarin Sodium)	SS		ORAL
2.5 MG (,DAILY) , PER ORAL							

Asa	C
Prinivil	C
Lasix	C
Nifedipine	C
Zaroxolyn	C
Lanoxin	C
Mevacor	C
Vitamin B6	C
Vitamin C	C
Vitamin E	C
Centrum	C
Folic Acid	C

Date:07/23/99ISR Number: 3310441-1Report Type:Direct
Age:17 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG QD Initial or Prolonged ORAL		Hangover		Gabapentin	PS		ORAL
1 MG Q 6H ORAL				Lorazepam	SS		ORAL
				Ativan	C		
				Resperidol	C		
				Benadryl	C		

Date:07/26/99ISR Number: 3311025-1Report Type:Expedited (15-DaCompany Report #044-0945-990078
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:07/27/99ISR Number: 3310631-8Report Type:Direct
Age:57 YR Gender:Female I/FU:I

Company Report #

Outcome	PT
Disability	Aggression Amnesia

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Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300MG 4 TIMES	DAY	Burning Sensation Depression Diplopia	Neurontin 300mg Parke Davis	PS	Parke Davis	
200MG 1 TIME		Dysgraphia Fear Formication Heart Rate Irregular Labile Blood Pressure	Celebrex 200mg Searle	SS	Searle	
		Lethargy Nervous System Disorder Nightmare Peripheral Coldness Speech Disorder Thinking Abnormal Tremor Vision Blurred	Armour Thyroid Xanax(Alprazolim)	C C		

Date:07/27/99ISR Number: 3357442-5Report Type:Periodic Company Report #1998SUS0286
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
		Acne Drug Interaction	Sustiva (Efavirenz) Ni (Citalopram) Neurontin (Gabapentin)	PS SS SS		

Date:07/28/99ISR Number: 3313411-2Report Type:Direct Company Report #
 Age: Gender:Male I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Other T QHS X3		Blister	Neurontin	PS		
DAYS;T BID X		Pruritus				
3DAYS;T TID						

THEREAFTER

Lortab	C
Flexeril	C
Trazadone	C
Tylenol	C
Alprazolam	C
Midrin	C
Triazolam	C
Cyanocobalamin	C
Omeprazole	C
Metoclopramide	C
Flonase	C
Fluticasone	C
Salmeterol	C
Albuterol	C
Diltiazem Xr	C
Phenytoin	C
Hydrochlorothiazine	C
Corevert	C
Flovent	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/99ISR Number: 3314137-1Report Type:Expedited (15-DaCompany Report #8250574

Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Bowel Sounds Abnormal Bradycardia Breath Sounds Decreased Coma Drug Level Above Therapeutic Drug Toxicity Haematoma Haemodialysis Hypotension Miosis Pneumonia Aspiration Procedural Site Reaction Respiratory Disorder Shock Suicide Attempt Thrombocytopenia	Literature Health Professional	Abbott-Depakote Gabapentin	PS SS	Abbott	

Date:07/30/99ISR Number: 3315214-1Report Type:Expedited (15-DaCompany Report #049-0945-990017

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 1400 MG (DAILY), PER ORAL		Pneumonia Thrombocytopenia White Blood Cell Count Decreased	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Lopirin Pantozol Fortecortin Morphine Bifiteral	C C C C C		

Date:07/30/99ISR Number: 3315432-2Report Type:Expedited (15-DaCompany Report #USA010182

Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Bradycardia	Other	Vicodin	PS		
		Cardiac Arrest		Digoxin	SS		
		Coma		Percocet	SS		
		Completed Suicide		Co-Trim	SS		
		Hypotension		Ativan	SS		
		Intentional Misuse		Valium	SS		
		Pupil Fixed		Neuroxin	SS		
		Therapeutic Agent		Imipramine	SS		
		Toxicity					
		Ventricular Fibrillation					
		Ventricular Tachycardia					

Date:07/30/99ISR Number: 3316074-5Report Type:Direct
Age:70 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Nausea		Gabapentin	PS		
Initial or Prolonged		Renal Failure Acute		Methocarbamol	C		
Other		Syncope		Cyclobenzaprine	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Paroxetine C
Amitri C

Date:07/30/99ISR Number: 3316093-9Report Type:Direct
Age:29 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Hypersensitivity Respiratory Distress		Gabapentin	PS		

Date:08/02/99ISR Number: 3316204-5Report Type:Expedited (15-DaCompany Report #10053825
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 40 MILLIGRAM, Initial or Prolonged ORAL		Blood Creatinine Increased Blood Urea Increased Cardiac Failure Condition Aggravated Renal Impairment	Study Health Professional	Omapatrilat Neurontin (Gabapentin)	PS SS		ORAL

Date:08/03/99ISR Number: 3317590-2Report Type:Expedited (15-DaCompany Report #001-0945-990442
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3200 MG (800 MG, QID)		Accident Complex Partial Seizures Drug Ineffective Thermal Burn	Health Professional	Neurontin (Gabapentin)	PS		
				Dilantin (Ucb)	C C		

Date:08/04/99ISR Number: 3319658-3Report Type:Expedited (15-DaCompany Report #001-0073-990304
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health Professional	Dilantin (Phenytoin Sodium)	PS		
600 MG							
(,DAILY)							
				Neurontin (Gabapentin)	SS		
600							
MG(,DAILY)							
				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine)	C		

Date:08/05/99ISR Number: 3319851-XReport Type:Expedited (15-DaCompany Report #001-0945-990413
Age:79 YR Gender:Female I/FU:F

Outcome	PT
Other	Alopecia Deafness Decreased Appetite Depression Dermatitis Dizziness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1600 MG (400 MG,QID),PER ORAL		Eructation	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
		Flatulence					
		Hypersensitivity					
		Hypoaesthesia					
		Muscular Weakness					
SCAB		Scab		Spinal Epidural	C		
		Skin Exfoliation					
		Skin Ulcer					
		Stomatitis					
		Tremor					
ORAL		Visual Disturbance					
		Visual Disturbance					

Date:08/05/99ISR Number: 3319859-4Report Type:Expedited (15-DaCompany Report #044-0945-990080

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly TRANSPLACENTAL (,DAILY), PLACENTAL; IN UTERO EXPOSURE	2900 MG	Abortion Induced	Foreign Health Professional	Neurontin (Gabapentin)	PS		
		Anal Atresia					
		Complications Of Maternal Exposure To Therapeutic Drugs					
		Congenital Anomaly					
		Congenital Diaphragmatic Eventration					
		Congenital Ectopic Bladder		(Lamotrigine)	C		
		Exomphalos		(Clobazam)	C		
		Limb Malformation					
		Pulmonary Hypoplasia					
		Spine Malformation					
		Talipes					
		Talipes					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 4800 MG (1200 Initial or Prolonged MG, QID), PER		Anaemia Aplastic Anaemia Bone Marrow Depression Condition Aggravated	Health Professional	Neurontin (Gabapenin)	PS		ORAL
ORAL 1200 MG (300 MG, QID), PER		Glossopharyngeal Neuralgia Haematocrit Decreased Haemoglobin Decreased Neutropenia Pain Pancytopenia Pyrexia Thrombocytopenia Weight Decreased		Tegretol (Carbamazepine)	SS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2 TID		Hypoaesthesia		Neurontin 300mg Parke Davis Neurontin 400 Mg	PS	Parke Davis	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

2 TID

Parke Davis SS Parke Davis

Atenolol C
Hydrocodone/Apap C
5/500

Date:08/06/99ISR Number: 3321590-6Report Type:Direct
Age:45 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required	20MG DAILY	9 MON	Anxiety	Prilosec	PS		
Intervention to Prevent Permanent Impairment/Damage	100MG/5CC		Drug Interaction Eye Rolling Facial Palsy Joint Stiffness Lip Disorder Medication Error Overdose	Neurontin Capsules Tegretol Susp 100mg/5cc	SS SS		

Date:08/09/99ISR Number: 3321332-4Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG PO TID	2 WK	Alopecia	Neurontin 300mg /Parke Davis	PS	Parke Davis	ORAL
				Prozac Xanax Synthroid Micronor Zretropbic-S Alphagan	C C C C C C		

Date:08/09/99ISR Number: 3321335-XReport Type:Direct
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Alopecia	Neurontin 300mg /			
300GM 600 AS		Parke Davis	PS	Parke Davis	ORAL
(PO)	2 WK	Depakote	C		
		Birth Control Pill	C		

Date:08/12/99ISR Number: 3325058-2Report Type:Expedited (15-DaCompany Report #001-0945-990356
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Blindness	Health	Neurontin			
Other		Macular Degeneration	Professional	(Gabapentin)	PS		
600MG BID		Visual Field Defect	Company Representative				

Date:08/17/99ISR Number: 3327673-9Report Type:Expedited (15-DaCompany Report #99USA10855
 Age:39 YR Gender:Male I/FU:I

Outcome	PT
Life-Threatening	Bone Marrow Depression
Hospitalization -	Drug Interaction
Initial or Prolonged	Leukopenia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Neutropenia Pancytopenia Thrombocytopenia	Report Source	Product	Role	Manufacturer	Route
300 MG, QID, ORAL			Health Professional	Tegretol Tablet (Carbamazepine)	PS		ORAL
4800 MG, DAILY, ORAL				Neurontin Capsule (Gabapentin)	SS		ORAL

Date:08/18/99ISR Number: 3328583-3Report Type:Expedited (15-DaCompany Report #033-0945-990051
Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 3600 MG (,DAILY), PER ORAL		Convulsion Hemiplegia	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:08/18/99ISR Number: 3328584-5Report Type:Expedited (15-DaCompany Report #001-0945-990442
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3200 MG (800 MD, QID)		Accident Complex Partial Seizures Condition Aggravated	Health Professional	Neurontin (Gabapentin)	PS		
		Drug Ineffective Simple Partial Seizures		Dilantin (Ucb Lo59)	C C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aplastic Anaemia	Health Professional	Neurontin (Gabapentin)	PS		

Date:08/18/99ISR Number: 3329386-6Report Type:Direct

Company Report #

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Weight Increased		Neurontin 300 Mg	PS		
1 TID				Amitriptyline 75 Mg			
				Qhs(Start 8/98)	C		
				Premarin N0.625 Mg			
				Qd(Long Term Use)	C		
				Imitrex Tabs Prn			
				(Long Term Use)	C		
				Phrenilin Prn(Long			
				Term Use)	C		

Date:08/18/99ISR Number: 3329515-4Report Type:Direct

Company Report #

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

Outcome	PT
Hospitalization -	Chest Pain
Initial or Prolonged	Muscle Twitching

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nervousness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
200 MG TID PO			Neurontin Po	PS		ORAL
, WITH RECENT INCREASE IN DOSE						
			Claritin D	C		
			Desyrel	C		
			Paxil	C		
			Sulindac	C		

Date:08/19/99ISR Number: 3329358-1Report Type:Expedited (15-DaCompany Report #10048908
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 40 Initial or Prolonged MILLIGRAMS, 2/DAY ORAL	Blood Creatinine Increased Blood Urea Increased Cardiac Failure Congestive Condition Aggravated Dialysis Renal Impairment	Study Health Professional	Omapatrilat	PS		ORAL
			Neurontin (Gabapentin)	SS		
			Vitamins + Minerals	C		
			Lanoxin	C		
			Furosemide	C		
			Zaroxolyn	C		
			Quinine	C		
			Isosorbide Dinitrate	C		
			Amlodipine	C		
			Premarin	C		
			Alprazolam	C		
			Aspirin	C		
			Nph Insulin	C		
			Insulin Sliding Scale	C		
			Sublingual Nitroglycerin	C		

Tylenol C
 Cimetidine C
 Allopurinol C
 Famotidine C

Date:08/26/99ISR Number: 3335019-5Report Type:Expedited (15-DaCompany Report #001-0945-990785
 Age:79 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Blood Creatinine	Health	Neurontin Capsules			
Initial or Prolonged	Increased	Professional	300 Mg (Gabapentin)	PS		
Required	Cardiac Arrest		Vasotec	C		
Intervention to	Cyanosis					
Prevent Permanent	Renal Failure Acute					
Impairment/Damage	Syncope					

Date:08/26/99ISR Number: 3335470-3Report Type:Expedited (15-DaCompany Report #001-0945-990792
 Age:45 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Cholecystectomy
Initial or Prolonged	Gallbladder Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Gingival Bleeding Gingivitis Oral Mucosal Blistering	Health Professional	Neurontin (Gabapentin) Bactrim (Sulfamethoxazole, Trimethoprim)	PS C		

Date:08/26/99ISR Number: 3335485-5Report Type:Expedited (15-DaCompany Report #033-0945-990044
Age:53 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG; 1600 MG; 2400 MG; 3600 MG, DAILY PER ORAL			Clostridium Colitis Diarrhoea Intestinal Functional Disorder Intestinal Obstruction	Foreign Health Professional	Neurontin (Gabapentin) Tegretol (Carbamazepine) Depakine (Valproate Sodium) Daflon (Diosmin) Hept-A-Myl (Heptaminol Hydrochloride)	PS C C C C		ORAL

Date:08/27/99ISR Number: 3336220-7Report Type:Expedited (15-DaCompany Report #044-0945-990096
Age:39 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3600 MG,			Dysphagia Oesophageal Carcinoma	Foreign Health	Neurontin (Gabapentin)	PS		ORAL

Required Oesophagectomy Professional

DAILY, PER
Intervention to
ORAL
Prevent Permanent
Impairment/Damage

Tegretol C
Epanutin C
(Doxycycline) C

Date:09/01/99ISR Number: 3338636-1Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis	Health	Neurontin 300mg Po	PS		ORAL
300MG PO		Dyspnoea	Professional				

Date:09/01/99ISR Number: 3338926-2Report Type:Expedited (15-DaCompany Report #001-0945-990799
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Laryngeal Oedema	Health	Neurontin			
Other		Laryngitis	Professional	(Gabapentin)	PS		
		Neoplasm					
		Obstructive Airways					
		Disorder					
		Swelling					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/99ISR Number: 3339071-2Report Type:Expedited (15-DaCompany Report #002-0945-990009
Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG (100 MG, BID), PER ORAL (SEE IMAGE)		Retinal Artery Thrombosis Visual Acuity Reduced	Foreign Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

Diclofenac	C
Acetaminophen	C
Enalapril	C
Humulin N	C
Centrum	C
Vitamin B12	C
Aspirin E.C.	C

Date:09/01/99ISR Number: 3339098-0Report Type:Expedited (15-DaCompany Report #001-0945-990786
Age:8 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (100 MG, 6QD), PER ORAL		Anger Balance Disorder Convulsion Discomfort Fall Fatigue Gingival Bleeding Gingivitis Haematuria Hypotonia Irritability Joint Swelling Muscular Weakness Oral Discomfort Personality Change	Health Professional Company Representative	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

Depakote (Valproate Semisodium) (Zinc)	C
Vitamin E (Tocopherol) (Phenobarbital)	C

Pyrexia
 Sedation
 Swelling
 Systemic Lupus
 Erythematousus
 Tooth Abscess

Date:09/03/99ISR Number: 3341258-XReport Type:Expedited (15-DaCompany Report #001-0945-990798
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Health	Neurontin Capsules			
Hospitalization - 900 MG (300 Initial or Prolonged MG, TID), PER		Encephalopathy	Professional	300 Mg (Gabapentin)	PS		ORAL
		Hypertension					
ORAL		Lethargy					
		Myocardial Fibrosis		Insulin Nph	C		
		Pneumonia		Amitriptyline	C		
		Urinary Tract Infection		Metoclopramide	C		
				Lisinopril	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/07/99ISR Number: 3340968-8Report Type:Direct
Age:71 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State Nightmare	Health Professional	Gabapentin Fentanyl Transdermal System Patch Morphine Sulf Nortriptyline Casanthranol/Docusat e	PS C C C C C		

Date:09/07/99ISR Number: 3342223-9Report Type:Expedited (15-DaCompany Report #001-0945-990800
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 300 MG PER		Cardiac Arrest Hyperkalaemia	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
ORAL (QHS)			Other	Elavil Risperidone Clonidine Zoloft Vitamin Percocet Hydroxyzine	C C C C C C C		

Date:09/07/99ISR Number: 3342225-2Report Type:Expedited (15-DaCompany Report #001-0945-990801
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 1800 MG DAILY		Chest Pain Pain	Consumer	Neurontin (Gabapentin)	PS		
		Pulmonary Oedema Weight Increased					

Date:09/08/99ISR Number: 3342777-2Report Type:Direct
Age:51 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Health	Gabapentin	PS		ORAL
300MG 4 CAPS			Professional				
TID ORAL	209 DAY						
				Valproic Acid	C		
				Phenytoin	C		
				Loperamide Hcl	C		
				Ascorbic Acid	C		
				Vitamin B Complex	C		
				Vitamin E	C		

Date:09/08/99ISR Number: 3343403-9Report Type:Expedited (15-DaCompany Report #033-0945-990055
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Hyponatraemia	Foreign Health	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
1200 MG (400 MG, TID), PER			Professional				
ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

250 MG (250
 MG, DAILY)
 PER ORAL

Primidone(Primidone) SS ORAL

Date:09/09/99ISR Number: 3344154-7Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG PO		Confusional State	Health	Phenytoin	PS		ORAL
Initial or Prolonged TID		Sedation	Professional				
200MG PO				Gabapentin	SS		ORAL
				Doxazosin	C		
				Lisinopril	C		
				Albuterol	C		
				Atrovent	C		
				Zantac	C		
				Isoniazid	C		
				Pyridoxine	C		

Date:09/16/99ISR Number: 3347966-9Report Type:Direct
 Age:57 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG TID	1 DAY	Hallucination		Gabapentin - Neurotin 300 Mg	PS		

Date:09/16/99ISR Number: 3349609-7Report Type:Expedited (15-DaCompany Report #002-0945-990026
 Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG	Blood Creatine Phosphokinase Increased	Foreign Health	Neurontin(Gabapentin)	PS		
Other (DAILY)		Fall	Professional				
	1200 MG	Hepatic Enzyme Increased		Neurontin(Gabapentin)	SS		
	(DAILY)			Tylenol #3	C		
				Tylenol #3	C		
				Hydrochlorothiazide/ Triamterene	C		
				Furosemide	C		
				Allopurinol	C		
				Indomethacin	C		

Date:09/16/99ISR Number: 3349611-5Report Type:Expedited (15-DaCompany Report #033-0945-990056
Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1600 MG	Cholangitis Hepatic Enzyme Increased	Foreign Health	Neurontin(Gabapentin)	PS		ORAL
Required (DAILY), PER		Jaundice	Professional				
Intervention to ORAL		Pyrexia					
Prevent Permanent Impairment/Damage	1500 MG	Sepsis Stupor		Depakine(Valproate Sodium)	SS		ORAL
	(DAILY), PER						

ORAL

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Freedom Of Information (FOI) Report

Caldine C
 Acuitel C
 Glucor C

Date:09/16/99ISR Number: 3349614-0Report Type:Expedited (15-DaCompany Report #001-0073-990173
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Abdominal Distension Abdominal Pain Alopecia	Consumer	Dilantin Kapseals 100 Mg(Phenytoin Sodium)	PS		ORAL
300 MG (100 MG, TID), PER ORAL		Anxiety Arthralgia					
NDA #20-235		Blood Potassium Increased Confusional State		Neurontin(Gabapentin)	SS		
		Convulsion Disorientation Drug Level Below Therapeutic Fall Gingival Hyperplasia Myalgia Palpitations Parkinsonian Gait Tinnitus Visual Acuity Reduced Weight Increased		Prempro	C		

Date:09/17/99ISR Number: 3349410-4Report Type:Direct Company Report #
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1500 MG/DAY 7 YR Initial or Prolonged 200 MG/DAY		Condition Aggravated Grand Mal Convulsion		Neurontin Celebrex	PS SS		

Date:09/20/99ISR Number: 3352296-5Report Type:Expedited (15-DaCompany Report #001-0945-990900
Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Pulmonary Embolism	Health	Neurontin			
Initial or Prolonged		Professional	(Gabapentin)	PS		
			Celexa	C		
			Nortriptyline	C		

Date:09/21/99ISR Number: 3352979-7Report Type:Direct Company Report #
Age:50 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Drug Effect Decreased		Gabapentin	PS		ORAL
400QID PO						
Initial or Prolonged	Lower Limb Fracture					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/21/99ISR Number: 3353364-4Report Type:Direct
 Age:59 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Idiopathic		Neurontin 300mg 2			
Hospitalization - 300MG 2TT Q	Thrombocytopenic Purpura		Qhs	PS		ORAL
Initial or Prolonged HS PO	Platelet Count Decreased					
Other Required Intervention to Prevent Permanent Impairment/Damage	Thrombocytopenia		Oops	C		

Date:09/21/99ISR Number: 3353365-6Report Type:Direct
 Age:47 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Candidiasis		Neurontin	PS		
Initial or Prolonged	Dermatitis		Celebrex	SS		

Date:09/21/99ISR Number: 3353373-5Report Type:Direct
 Age:59 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Abdominal Pain		Gabapentin			
Initial or Prolonged	Duodenitis		(Neurontin)/Unknown			
	Gastritis		/Parke-Davis	PS		
	Nausea		Coumadin	C		
	Vomiting		Reglan	C		
	Weight Decreased		Carafate	C		
			Prilosec	C		

Date:09/22/99ISR Number: 3354867-9Report Type:Expedited (15-DaCompany Report #033-0945-990048
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG		Aspartate Aminotransferase	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER		Increased	Professional				
ORAL		Hepatitis B					

Date:09/22/99ISR Number: 3355464-1Report Type:Direct
Age:59 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Duodenitis Gastritis		Gabapentin (Neurontin) /Unknown /Parke-Davis	PS	Parke-Davis	ORAL
ORAL		Nausea Vomiting Weight Decreased		Coumadin Reglan Carafate Prilosec	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/22/99ISR Number: 3359746-9Report Type:Periodic
Age:78 YR Gender:Female I/FU:I

Company Report #001-0073-980587

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Convulsion Dizziness	Consumer	Dilantin (Phenytoin Sodium)	PS		ORAL
300 MG (DAILY), PER ORAL	Drug Level Above Therapeutic Fatigue Sedation		Neurontin (Gabapentin)	SS		ORAL
			Propranolol	C		

Date:09/23/99ISR Number: 3357467-XReport Type:Expedited (15-DaCompany Report #001-0945-990908
Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG (600 MG, QAM AND QHS), PER ORAL	Grand Mal Convulsion Loss Of Consciousness Movement Disorder Tremor	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
			Doxepin (Doxepin)	C		
			Prozac (Fluoxetine Hydrochloride)	C		
			Ambien (Zolpidem Tartrate)	C		
			Unspecified Hormones	C		

Date:09/24/99ISR Number: 3356788-4Report Type:Direct
Age:53 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Other Required 2-300MG Intervention to 3X/DAY (1800 Prevent Permanent MG) 1 YR Impairment/Damage	Blood Iron Decreased Coeliac Disease Malaise Oedema Peripheral Swelling	Health Professional	Neurontin /300mg /Park/Davis	PS	Parke-Davis
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Date:09/27/99ISR Number: 3358788-7Report Type:Direct
Age:77 YR Gender:Female I/FU:I

Company Report #

Outcome Dose Life-Threatening 250MG TID (A 2 YRS ON DIFFERENT SEIZURE MEDICATIONS)	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion Thrombocytopenia		Dilantin Mysoline	PS SS		
				Neurontin Hydrochlorthiazide	SS SS		

Date:09/28/99ISR Number: 3358820-0Report Type:Direct
Age: Gender: I/FU:I

Company Report #

Outcome
Other
Required
Intervention to

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	Disturbance In Attention	Consumer	Meridia	PS	
	Fatigue	Other	Meridia	SS	ORAL
10 MG OD PO					
	Tremor		Neurotin	SS	ORAL
400 MG OD PO					

Date:09/29/99ISR Number: 3360703-7Report Type:Expedited (15-DaCompany Report #001-0945-990919
 Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Neurontin Capsules 300mg (Gabapentin)	PS		ORAL
3600MG							
(1200MG TID)							
PER ORAL							

Date:09/29/99ISR Number: 3360705-0Report Type:Expedited (15-DaCompany Report #001-0945-990913
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous	Health Professional	Neurontin (Gabapentin)	PS		
1600MG				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Triamine Hydrochloride,	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/29/99ISR Number: 3360707-4Report Type:Expedited (15-DaCompany Report #001-0073-990394

Age:1 DY Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic	Health Professional	Dilantin (Phenytoin Sodium)	PS		
550MG (DAILY)		Drugs					
PLACENTAL		Congenital Anomaly Pregnancy		Depakote (Valproate Semisodium)	SS		
750MG (DAILY)		Skin Disorder					
PLACENTAL	12 WK			Neurontin (Gabapentin)	SS		
2700MG (DAILY)							
PLACENTAL	12 WK			Folic Acid (Folic Acid)	C		

Date:09/29/99ISR Number: 3360998-XReport Type:Expedited (15-DaCompany Report #B0071068A

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Epilepsy	Foreign	Lamictal	PS		ORAL
TWICE PER DAY		Sudden Death					
/ ORAL				Gabapentin (Formulation Unknown) (Gabapentin)	SS		ORAL
400 MG /							
THREE TIMES							
PER DAY /							
ORAL							

Calcium Glubionate C
Ferrous Sulfate C

Date:10/04/99ISR Number: 3363790-5Report Type:Expedited (15-DaCompany Report #9940757
Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent 5.00 MG Impairment/Damage TOTAL; DAILY; ORAL		Convulsion Drug Interaction	Health Professional	Glucotrol Xl Extended Release Tablets	PS		ORAL
100.00 MG TOTAL; DAILY; ORAL				Neurontin	SS		ORAL
				Lipitor	C		

Date:10/08/99ISR Number: 3369321-8Report Type:Expedited (15-DaCompany Report #001-0073-990417
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 MG BID) PER ORAL		Angioplasty Aspiration Convulsion Coronary Artery Disease	Health Professional	Dilantin (Phenytoin Sodium)	PS		ORAL
PER ORAL		Gamma-Glutamyltransferase Increased		Neurontin (Gabaentin)	SS		ORAL
		Grand Mal Convulsion Myocardial Infarction		Lipitor (Atorvastatin)	C		

Freedom Of Information (FOI) Report

Aspirin
 (Acetylsalicylic
 Acid) C
 Lescol (Fluvastatin
 Sodium) C

Date:10/08/99ISR Number: 3369324-3Report Type:Expedited (15-DaCompany Report #001-0945-990792
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Gallbladder Disorder Gingival Bleeding Gingivitis Oral Mucosal Blistering Pyrexia Stomatitis Streptococcal Infection	Health Professional	Neurontin (Gabapentin) Bactrim (Sulfamethoxazole)	PS C		

Date:10/12/99ISR Number: 3370404-7Report Type:Expedited (15-DaCompany Report #001-0991-991427
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200 MG (DAILY)		Biliary Cirrhosis Primary Chest Pain Cholangitis Sclerosing	Consumer Other	Rezulin (Troglitazone)	PS		ORAL
10 MG (DAILY)		Cholestasis Ear Pain		Lipitor (Atorvastatin)	SS		ORAL
400 MG DAILY; ORAL		Gallbladder Disorder Gastroenteritis Helicobacter Glycosylated Haemoglobin Increased Inflammation Lacrimation Increased Liver Function Test		Neurontin (Gabapentin) Neurontin (Gabapentin) Rezulin (Troglitazone) Prandin	SS SS SS SS		ORAL

30 MG (10 MG,	Abnormal	(Repaglinide)	SS	ORAL
AC)	Muscle Spasms			
	Oedema Peripheral	Niacin	C	
	Pain	Glucotrol Xl		
	Pharyngolaryngeal Pain	(Glipizide)	C	
	Productive Cough	Atrovent Inhaler		
	Sinus Congestion	(Ipratropium		
	Sputum Discoloured	Bromide)	C	
	Tricuspid Valve	Proventil Inhaler		
	Incompetence	(Salbutamol)	C	
	Vaginal Candidiasis	Flexeril		
		(Cyclobenzaprine		
		Hydrochloride)	C	
		Citracal (Calcium		
		Citrate)	C	

Date:10/12/99ISR Number: 3371610-8Report Type:Expedited (15-DaCompany Report #001-0073-990458
Age: Gender:Male I/FU:I

Outcome PT
Hospitalization - Alanine Aminotransferase
Initial or Prolonged Increased
Aspartate

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Aminotransferase Increased Condition Aggravated Hepatic Fibrosis	Report Source	Product	Role	Manufacturer	Route
300 MG (HS), PER ORAL			Health Professional	Dilantin Kapseals 100 Mg (Phenytoin Sodium)	PS		ORAL
300 MG (AM), PER ORAL				Neurontin Capsules 300 Mg (Gabapentin)	SS		ORAL

Date:10/13/99ISR Number: 3370370-4Report Type:Direct
Age:65 YR Gender:Male I/FU:I

Company Report #

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300MG PO Q 8		Hypotension		Neurotin	PS		ORAL

HRS

Date:10/13/99ISR Number: 3371607-8Report Type:Expedited (15-DaCompany Report #A0102566A
Age:16 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other	150 MG/SEE		Aggression Coma Grand Mal Convulsion Intentional Misuse	Health Professional Company Representative	Wellbutrin Tablet-Controlled Release (Bupropion Hydrochloride)	PS		ORAL
TEXT/ORAL			Lethargy		Gabapentin (Formulation Unknown) (Gabapentin)	SS		ORAL

SEE TEXT/ORAL

Date:10/13/99ISR Number: 3371688-1Report Type:Expedited (15-DaCompany Report #001-0945-990361
 Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough	Health	Neurontin Capules			
		Groin Pain	Professional	300 Mg (Gabapentin)	PS		ORAL
900 MG (300,		Headache					
MG, TID), PER		Myocardial Infarction					
ORAL		Nausea		Daypro (Oxaprozin)	C		
		Vomiting					

Date:10/13/99ISR Number: 3381528-2Report Type:Expedited (15-DaCompany Report #SP-9901025
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Confusional State	Consumer	Remicade	PS		
Initial or Prolonged		Delusion	Health	Remicade	SS		
		Delusional Disorder,	Professional	Remicade	SS		
		Persecutory Type		Prednisone	SS		
LO-DOSE		Fatigue		Remicade	SS		
INTRAVENOUS	INTRAVENOUS	Pain		Prozac	SS		
ORAL 30 MG		Psychotic Disorder					
DAILY		Vomiting		Demerol	SS		
				Remicade	SS		
				Roxicodone	SS		ORAL
ORAL UP TO 2							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

TABLETS DAILY

ORAL 300 MG

THREE TIMES

DAILY

Neurontin SS ORAL

Pentasa C
 Cipro C
 Blephamide C
 Tetracycline C
 Tenex C
 Furosemide C
 Relafen C
 B-12 C
 Centrum Silver C
 Ms Contin C
 Prevacid C
 Lotrel C
 Ziac C
 Phenobarbital C
 K-Dur C
 Locholest Powder C
 Kerasal C
 Folic Acid C
 B/Complex-50 C
 B/Complex-100 C
 Magnesium Chloride C
 Calcium C
 Actigall C
 Diphenoxylate C
 Tridesilon C
 Rowasa C
 Proctofoam C
 Cortifoam C
 Claritin-D C

Date:10/15/99ISR Number: 3372051-XReport Type:Direct

Company Report #

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema Multiforme	Health	Gabipentin	PS		ORAL
PO			Professional				

Date:10/15/99ISR Number: 3374004-4Report Type:Expedited (15-DaCompany Report #001-0945-990735

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aplastic Anaemia Haemorrhagic Stroke	Health Professional	Neurontin (Gabapentin)	PS		
900 MG (300 MG, Q8H)		Hypotension Leukopenia Staphylococcal Sepsis Thrombocytopenia					

Date:10/18/99ISR Number: 3374188-8Report Type:Expedited (15-DaCompany Report #991012-SK615

Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health	Celebrex	PS		ORAL
200.000 MG QD PO 2000.000 MG		Drug Interaction	Professional	Gabapentin	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

QD PO
 300.000 MG
 TID PO
 Carbamazepine SS ORAL

Date:10/18/99ISR Number: 3374895-7Report Type:Expedited (15-DaCompany Report #001-0945-991004
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anaemia Hypoglycaemia Jaundice Psychotic Disorder	Health Professional	Neurontin (Gabapentin) Unspecified Medications	PS C		

Date:10/19/99ISR Number: 3375252-XReport Type:Expedited (15-DaCompany Report #033-0945-990065
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hepatitis	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:10/20/99ISR Number: 3376430-6Report Type:Expedited (15-DaCompany Report #001-0945-991001
 Age:1 DY Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 1200 MG DAILY PLACENTAL, CONCEPTION- WEEK 20 OF PREGNANCY-WEE		Complications Of Maternal Exposure To Therapeutic Drugs Stillbirth	Health Professional	Neurontin (Gabapentin)	PS		

1500 MG DAILY

Depakote (Valproate
Semisodium) SS

PLACENTAL,

CONCEPTION -

WEEK 20 OF

PREGNANCY-

Prenatal Vitamins
(Ergocalciferol,
Ascorbic Acid,
Pyridoxine
Hydrochloride, C
(Folic Acid) C

Date:10/20/99ISR Number: 3376552-XReport Type:Expedited (15-DaCompany Report #001-0945-991040
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health Professional	Neurontin (Gabapentin)	PS		
				Depakote (Valproate Semisodium)	C		

Freedom Of Information (FOI) Report

Date:10/20/99ISR Number: 3376554-3Report Type:Expedited (15-DaCompany Report #033-0945-990068
 Age:54 YR Gender:Male I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
PER ORAL	6 MON	Dermatitis Rash Pustular	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
			Professional	Daonil (Glibenclamide)	C		
				Glucophage (Metformin Hydrochloride)	C		
				Creon (Pancreatin)	C		

Date:10/21/99ISR Number: 3378259-1Report Type:Expedited (15-DaCompany Report #046-0945-990009
 Age: Gender: I/FU:I

Outcome Dose Hospitalization - Initial or Prolonged 300 MG EVERY SECOND DAY, PER ORAL	Duration	PT	Report Source	Product	Role	Manufacturer	Route
6 WK		Blister Blood Creatinine Increased Oedema Peripheral	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
			Professional	(Insulin)	C		

Date:10/25/99ISR Number: 3381834-1Report Type:Expedited (15-DaCompany Report #JRFBEL1999001677
 Age:46 YR Gender:Female I/FU:I

Outcome Dose Death	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 MG IN 1 DAY(S) ORAL	2	Cardiac Disorder Cardio-Respiratory Arrest Coma Coronary Artery Disease Toxicologic Test Abnormal	Literature Health Professional	Risperidone (Unspecified) (Risperidone)	PS		ORAL
				Gabapentin			

300 MG		(Gabapentin)	SS	
2 IN 1 DAY(S)				
10 MG	1	Clonazepam (Clonazepam)	SS	
IN 1 DAY(S)				
10 MG		Paroxetine (Paroxetine)	SS	
1 IN 1 DAY(S)				
15 MG	3	Remeron (Mirtazapine)	SS	ORAL
IN 1 DAY(S)				
ORAL				

Date:10/26/99ISR Number: 3382531-9Report Type:Expedited (15-DaCompany Report #001-0945-990908
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion Loss Of Consciousness Movement Disorder Tremor	Health Professional	Neurontin Gabapentin)	PS		ORAL
1200 MG (600 MG, QAM AND QHS), PER							
ORAL				Doxepin (Doxepin) Prozac (Fluoxetine Hydrochloride) Ambien (Zolpidem)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tartrate) C
 Unspecified Hormones C

Date:10/26/99ISR Number: 3382534-4Report Type:Expedited (15-DaCompany Report #033-0945-990069
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER ORAL		Balance Disorder Tremor	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Tegretol (Carbamazepine) Alepsal (Phenobarbital, Caffeine, Belladonna Extract)	C C		

Date:10/26/99ISR Number: 3382855-5Report Type:Periodic Company Report #9937604
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 25.00 MG TOTAL:PRN:ORA		Anorgasmia	Consumer	Viagra Tablets	PS		ORAL

L

				Neurontin	SS		
				Benadryl	C		
				Lithium	C		

Date:10/28/99ISR Number: 3383859-9Report Type:Expedited (15-DaCompany Report #001-0945-991048
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other Abortion Spontaneous Health Neurontin Capsules
Benign Hydatidiform Mole Professional 400 Mg(Gabapentin) PS

400 MG

(DAILY), 1 MON

Date:10/28/99ISR Number: 3383861-7Report Type:Expedited (15-DaCompany Report #001-0945-990900

Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2100	Pulmonary Embolism	Health Professional	Neurontin(Gabapentin)	PS		ORAL

MG(DAILY),

PER ORAL

Celexa(Citalopram Hydrobromide) (Nortriptyline)	C C
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Date:10/29/99ISR Number: 3383833-2Report Type:Direct

Age:52 YR Gender:Male I/FU:I

Company Report #

Outcome	PT
Hospitalization - Initial or Prolonged	Dysarthria Hypotension Lethargy Loss Of Consciousness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Overdose

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Inderal	PS		
			Nefazadone	SS		
			Neurontin	SS		
			Zantac	C		
			Lisinipril	C		

Date:10/29/99ISR Number: 3384938-2Report Type:Direct Company Report #
 Age:64 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State		Neurontin	PS		
		Lethargy		Ultram	SS		

Date:10/29/99ISR Number: 3393339-2Report Type:Periodic Company Report #8-98345-067A
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Consumer	Ativan Tablets	PS		ORAL
Other		Gait Disturbance					
1 MG ONCE							
DAILY ORAL		Paraesthesia		Elavil			
		Sedation		(Amitriptyline)			
		Speech Disorder		Tablets	SS		ORAL
100 MG DAILY							
ORAL				Flomax (Tamsulosin)			
				Capsules	SS		ORAL
0.4 MG DAILY							
ORAL				Neurontin			
				(Gabapentin)			
				Capsules	SS		ORAL
DOSE							

INCREASED UP

TO 1200 MG

DAILY ORAL

Flomax (Tamsulosin)
Capsules C
Elavil
(Amitriptyline)
Tablets C
Neurontin
(Gabapentin)
Capsules C

Date:11/01/99ISR Number: 3386804-5Report Type:Periodic
Age:35 YR Gender:Male I/FU:I

Company Report #M088774

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Amnesia Emotional Disorder	Health Professional	Buspar Tabs (Buspirone Hcl)	PS		ORAL
ORAL		Hostility Psychotic Disorder		Neurontin (Gabapentin)	SS		ORAL
ORAL	1 WK			Claritin	C		
				Iodine	C		
				Inderal	C		
				Remeron	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/01/99ISR Number: 3386879-3Report Type:Expedited (15-DaCompany Report #046-0945-990009

Age: Gender:Unknown I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG EVERY SECOND DAY	Blister Blood Creatine Increased Infection Oedema Peripheral Renal Failure Acute	Foreign Health Professional	Neurontin (Gabapentin) (Insuliin)	PS C		ORAL

Date:11/02/99ISR Number: 3388577-9Report Type:Expedited (15-DaCompany Report #001-0945-990735

Age:70 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 900MG (300MG Q8H)	Aplastic Anaemia Autoimmune Disorder Drug Interaction Haemorrhagic Stroke Platelet Count Decreased Staphylococcal Infection Thrombocytopenia White Blood Cell Count Decreased	Health Professional	Neurontin (Gabapentin) Carbamazepine (Carbamazepine)	PS SS		

Date:11/03/99ISR Number: 3389013-9Report Type:Expedited (15-DaCompany Report #001-0945-990707

Age:71 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (DAILY), UNKNOWN	Blood Catecholamines Increased Epinephrine Increased Norepinephrine Increased	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:11/03/99ISR Number: 3392994-0Report Type:Periodic
Age:52 YR Gender:Male I/FU:I

Company Report #1999070012

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1 TAB Q8H, PO	Drug Toxicity	Health Professional	Carisoprodol + Aspirin	PS		ORAL
DATA NA, PO			Zyprexa (Olanzapine)	SS		ORAL
DATA NA, PO			Neurontin (Gabapentin)	SS		ORAL
DATA NA, PO			Paxil (Paroxetine)	C		ORAL
			Pepcid	C		
			Armour Thyroid	C		

Date:11/04/99ISR Number: 3389255-2Report Type:Expedited (15-DaCompany Report #001-0945-990919
Age:34 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death 3600 MG (1200 MG, TID), PER ORAL	Epilepsy	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/05/99ISR Number: 3390033-9Report Type:Expedited (15-DaCompany Report #001-0945-990363

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged SEE TEXT, PER	Bone Cyst Ecchymosis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL	Increased Tendency To Bruise Platelet Count Decreased Platelet Function Test Abnormal		Prozac (Fluoxetine Hydrochloride) Unspecified Tricyclic Antidepressant	C C		

Date:11/05/99ISR Number: 3390036-4Report Type:Expedited (15-DaCompany Report #001-0945-990786

Age:8 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG (100 MG, 6QD), PER	Anger Asthenia	Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
ORAL	Autoimmune Disorder Balance Disorder Discomfort Epilepsy Fall Fatigue Gingival Bleeding Gingivitis Haematuria Hypotonia Irritability Joint Swelling Oedema Peripheral Personality Disorder Pyrexia Sedation Tooth Abscess	Company Representative	Depakote (Valproate Semisodium) (Zinc) Vitamin E (Tocopherol) (Phenobarbital)	C C C C		

Date:11/05/99ISR Number: 3390039-XReport Type:Expedited (15-DaCompany Report #033-0945-990065
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged SEE IMAGE	Hepatitis	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:11/05/99ISR Number: 3390073-XReport Type:Expedited (15-DaCompany Report #001-0073-990417
Age:48 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600MG (300MG BID) PER ORAL	Angioplasty Aspiration Complex Partial Seizures	Health Professional	Dilantin (Phenytoin Sodium)	PS		ORAL
PER ORAL	Coronary Artery Disease Coronary Artery Surgery		Neurontin (Gabapentin)	SS		ORAL
	Gamma-Glutamyltransferase Increased Myocardial Infarction		Lipitor (Atorvastatin) Aspirin (Acetylsalicylic Acid)	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lescol (Fluvastatin Sodium) C

Date:11/09/99ISR Number: 3391927-0Report Type:Expedited (15-DaCompany Report #044-0945-990139
Age:66 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Abdominal Pain	Foreign	Neurontin	PS		ORAL
Initial or Prolonged	Hepatitis	Health Professional	(Gabapentin) (Lactulose) Losec (Omeprazole)	C C		

Date:11/10/99ISR Number: 3391573-9Report Type:Direct Company Report #
Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Feeling Jittery	Health	Gabapentin	PS		ORAL
Initial or Prolonged	Intentional Misuse Medication Error Nausea	Professional				

Date:11/10/99ISR Number: 3394036-XReport Type:Expedited (15-DaCompany Report #039-0945-990006
Age:44 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion	Foreign	Gabapentin	PS		ORAL
Initial or Prolonged		Health Professional	(Gabapentin)			
160 MG						
(DAILY), PER						

ORAL

Tavor (Lorazepam) C
Axoren (Buspirone Hydrochloride) C

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG		Arteriopathic Disease Coagulopathy	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
(DAILY), PER		Systemic Lupus	Professional				
ORAL		Erythematosis					
300 MG		Thrombosis Vasospasm		Maveral (Fluvoxamine Maleate)	SS		ORAL
(DAILY), PER							
ORAL				Orap (Pimozide)	SS		
				Seroxat (Paroxetine Hydrochloride)	SS		
				Tegretol (Carbamazepine)	SS		
				Surmontil (Trimipramine)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/10/99ISR Number: 3394044-9Report Type:Expedited (15-DaCompany Report #001-0945-991095

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 3200 MG (DAILY), PER ORAL		Hepatic Steatosis Hepatitis Hyperlipidaemia Liver Function Test Abnormal Weight Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Prempro (Medroxyprogesterone , Acetate, Estrogens Conjugated) Prevacid (Lansoprazole) (Amitriptyline) Naprosyn (Naproxen) (Lorazepam) Thorazine (Chlorpromazine Hydrochloride)	C C C C C C		

Date:11/10/99ISR Number: 3394049-8Report Type:Expedited (15-DaCompany Report #001-0945-990508

Age:81 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (AFTER EACH DIALYSIS TREATMENT) 6 MG (QHS)		Activated Partial Thromboplastin Time Prolonged International Normalised Ratio Increased Prothrombin Time Prolonged	Health Professional	Neurontin (Gabapentin)	PS		
				Coumadin (Warfarin Sodium) (Erythropoietin) (Insulin) Cozaar (Losartan Potassium) (Glipizide)	SS C C C C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Abdominal Pain	Consumer	Neurontin Capsules			
Initial or Prolonged	Gastric Ulcer	Health	400 Mg (Gabapentin)	PS		ORAL
1200 MG (QD),						
Other	Gastritis	Professional				
PER ORAL						
	Glycosuria					
	Haematochezia					
	Malaise					
	Nephritis Interstitial					
	Nephrolithiasis					
	Proteinuria					
	Renal Disorder					
	Vomiting					
	Weight Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/12/99ISR Number: 3396961-2Report Type:Expedited (15-DaCompany Report #001-0945-991109

Age:1 DY Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly 300 MG (DAILY), PLACENTAL; IN UTERO EXPOSURE	Duration Complications Of Maternal Exposure To Therapeutic Drugs Polydactyly	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		
750 MG (DAILY), PLACENTAL; IN UTERO EXPOSURE			Depakote (Valproate Semisodium)	SS		
			Unspecified Antibiotics Ritalin (Methylphenidate Hydrochloride) "Clotadin" ...	C C C C		

Date:11/12/99ISR Number: 3397340-4Report Type:Expedited (15-DaCompany Report #001-0945-990505

Age:31 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 3600 MG (2400 Other MG QAM, 1200	Duration Hallucination Logorrhoea Multiple Sclerosis	Health Professional	Neurontin (Gabapentin)	PS		ORAL

MG QPM), PER

ORAL

Many Unspecified Medications C

Date:11/17/99 ISR Number: 3398677-5 Report Type:Direct
Age:65 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 100MG BID PO	Abdominal Pain Pancreatitis		Gabapentin (Neurontin)	PS		ORAL
			Amlodipine (Norvasc)	SS		

Date:11/17/99 ISR Number: 3399557-1 Report Type:Expedited (15-DaCompany Report #001-0945-991130
Age:62 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Blindness Diplopia Optic Nerve Disorder Retinal Disorder Vision Blurred	Consumer	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/18/99ISR Number: 3402695-8Report Type:Expedited (15-DaCompany Report #99USA11289
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Drug Interaction	Health Professional	Carbamazepine Tablet 300 Mg (Carbamazepine)	PS		ORAL
300 MG, TID, ORAL				Gabapentin Tablet 2000 Mg (Gabapentin)	SS		ORAL
2000 MG, TID, ORAL				Celebrex Tablet 200 Mg (Celecoxib)	SS		ORAL
200 MG, DAILY, ORAL							

Date:11/19/99ISR Number: 3403143-4Report Type:Expedited (15-DaCompany Report #033-0945-990064
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (DAILY), PER ORAL		Confusional State Depressed Level Of Consciousness Drug Level Below	Foreign Study Health Professional	Gabapentin (Gabapentin)	PS		ORAL
		Therapeutic Dry Mouth Electroencephalogram Abnormal Epilepsy Faecal Incontinence Grand Mal Convulsion Hyponatraemia Loss Of Consciousness Polydipsia Polyuria		Piascledine (Soya Oil, Avocado Oil) Tegretol (Carbamazepine) Floxyfral (Fluvoxamine Maleate)	C C C		

Status Epilepticus
 Stereotypy
 Thirst
 Urinary Incontinence
 Urine Sodium Decreased

Date:11/22/99ISR Number: 3404156-9Report Type:Direct
 Age:73 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 300MG TID,PO		Body Temperature	Health	Gabapentin	PS		ORAL
Intervention to 10-20 MG HS, Prevent Permanent PO 8 Impairment/Damage	8 WK	Increased Decreased Appetite Dermatitis Hepatitis Hepatocellular Damage Injury Jaundice Liver Function Test Abnormal Nausea Rash Maculo-Papular	Professional	Amitriptylline	SS		ORAL
				Viokase	C		
				Omeprazole	C		
				Timolol Eye Drops	C		
				Xalatan Eye Drops	C		
				Vitamin E	C		
				Vitamin C	C		
				Dyazide	C		
				Zostrix Cream	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/22/99ISR Number: 3405277-7Report Type:Expedited (15-DaCompany Report #9940757

Age:76 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage	5.00 MG	Blood Glucose Increased Diabetes Mellitus Inadequate Control	Health Professional	Glucotrol Xl Extended Release Tablets	PS		ORAL
TOTAL:DAILY:	0	Drug Interaction Epilepsy					
RAL				Neurontin	SS		ORAL
100.00 MG							
TOTAL:DAILY:	0						
RAL				Topiramate	SS		ORAL
50.00 MG							
TOTAL DAILY							
ORAL				Lipitor	C		

Date:11/22/99ISR Number: 3405794-XReport Type:Periodic

Company Report #8-99188-139A

Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other SEE IMAGE		Grand Mal Convulsion	Health Professional	Effexor Xr	PS		ORAL
400 MG FOUR				Neurontin	SS		ORAL
TIMES DAILY,							
ORAL				Remeron (Mirtazapine)	C		
				Neurontin (Gabapentin)	C		
				Xanax (Alprazolam)	C		

Date:11/24/99ISR Number: 3406789-2Report Type:Direct
Age:52 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	1800 MG DAILY	Ejaculation Failure		Neurontin	PS		
AS 600MG 3		Erectile Dysfunction					
X/DAY							

Date:11/30/99ISR Number: 3410392-8Report Type:Direct
Age:51 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	1,800MG /DAY	Arthralgia		Neurontin	PS		
		Sexual Dysfunction					

Date:12/01/99ISR Number: 3412381-6Report Type:Expedited (15-DaCompany Report #001-0945-990799
Age:27 YR Gender:Male I/FU:F

Outcome	PT
Death	Drug Ineffective
Other	Dysphagia
	Dysphonia
	Facial Palsy
	Inflammation
	Laryngeal Disorder
	Laryngeal Oedema
	Neoplasm

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Obstructive Airways Disorder Vocal Cord Paralysis	Report Source	Product	Role	Manufacturer	Route
1200 MG			Health Professional	Neurontin (Gabapentin)	PS		ORAL
(400 MG, TID)							
PER							
ORAL							
				Cyclosporine (Ciclosporin)	C		
				Morphine (Morphine)	C		
				Nystatin (Nystatin)	C		
				Fentanyl Patch (Fentanyl)	C		
				Lorazepam (Lorazepam)	C		
				Scopolamine Patch (Hyoscine)	C		
				Lansoprazole (Lansoprazole)	C		
				Sulfamethoxazole-Tri methoprim (Sulfamethoxazole, Trimethoprim)	C		

Date:12/01/99ISR Number: 3412385-3Report Type:Expedited (15-DaCompany Report #045-0945-990004
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1200 MG	Toxic Epidermal Necrolysis	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
(DAILY)							
ORAL				Eltroxin (Levothyroxine Sodium)	C		

Pamol (Paracetamol) C
 Losec (Omeprazole) C
 Noritren
 (Nortriptyline
 Hydrochloride) C

Date:12/02/99ISR Number: 3413007-8Report Type:Expedited (15-DaCompany Report #049-0945-980013
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 600 MG (DAILY), PER ORAL		Coma Drug Toxicity	Foreign Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
INTRAVENOUS THE EVENING), INTRAVENOUS	680 MG (IN			Carboplatin (Carboplatin)	SS		
				M-Dolor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/03/99ISR Number: 3413575-6Report Type:Expedited (15-DaCompany Report #046-0945-990013

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 300 MG (EVERY Initial or Prolonged OTHER DAY), PER ORAL	Blister Dialysis Leg Amputation Renal Failure	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Achapid-Insulatard (Insulin Human Injection, Isophane)	C		
			Levoxin (Levothyroxine Sodium)	C		
			Rhinocort (Budesonide)	C		
			(Acetylcysteine)	C		
			Nitromex (Glyceryl Trinitrate)	C		
			Glucophage (Metformin Hydrochloride)	C		
			Zantac (Ranitidine Hydrochloride)	C		
			Renitec (Enalapril Maleate)	C		
			Trimgyl	C		
			Lasix (Furosemide)	C		
			Distalgesic (Paracetamol, Dextropropoxyphene Hydrochloride)	C		
			Lopid (Gemfibrozil)	C		
			Zocord (Simvastatin)	C		
			Behepan (Cyanocobalamin)	C		
			Seloken Zoc (Metoprolol Succinate)	C		

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Creatine Phosphokinase Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
400 MG (200 MG BID) PER ORAL		Blood Creatine Phosphokinase Mb Increased					

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged		Alopecia Amnesia Convulsion Tremor	Consumer	Neurontin (Gabapentin) Micro K (Potassium Chloride) Lasix (Furosemide) Oxycontin (Oxycodone)	PS C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C
 Pamelor
 (Nortriptyline
 Hydrochloride) C
 Lodine (Etodolac) C
 Dilantin (Phenytoin
 Sodium) C
 Ultram (Tramadol
 Hydrochloride) C

Date:12/06/99ISR Number: 3414308-XReport Type:Expedited (15-DaCompany Report #044-0945-990166
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Foreign Health	Neurontin(Gabapentin)	PS		ORAL
Other			Professional				
1800 MG							
(DAILY), PER							
ORAL							

Date:12/07/99ISR Number: 3415350-5Report Type:Expedited (15-DaCompany Report #001-0945-991179
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Drug Interaction	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
400 MG		Haemorrhage					
(DAILY), PER		International Normalised					
ORAL/800 MG		Ratio Increased					
(400 MG,		Loss Of Consciousness					
BID), PER		Platelet Count Decreased		Coumadin (Warfarin Sodium)	SS		ORAL
3.75 MG							
(DAILY), PER							

ORAL

(Insulin)

C

Date:12/07/99ISR Number: 3415388-8Report Type:Expedited (15-DaCompany Report #001-0945-991172

Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG, TID), PER		Blood Bilirubin Increased Decreased Appetite Dermatitis Hepatitis	Health Professional	Gabapentin (Gabapentin)	PS		ORAL
ORAL 10 - 20 MG (HS), PER		Jaundice Liver Function Test Abnormal		(Amitriptyline)	SS		ORAL
ORAL	8 WK	Nausea Pyrexia Rash Maculo-Papular		Viokase (Pancrelipase) (Omeprazole) Timolol Eye Drops (Timolol) Xalatan Eye Drops (Latanoprost) Vitamin E (Tocopherol) Vitamin C (Ascorbic Acid) Dyazide (Hydrochlorothiazide)	C C C C C C C		

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Freedom Of Information (FOI) Report

, Triamterene)	C
Zostrix Cream	
(Capsaicin)	C
Nerve Block	C
Antiviral Therapy	C
Steroid Nerve Root	
Therapy	C

Date:12/07/99ISR Number: 3415391-8Report Type:Expedited (15-DaCompany Report #001-0945-990707
 Age:71 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Blood Catecholamines	Health	Neurontin			
Initial or Prolonged	Increased	Professional	(Gabapentin)	PS		
1200 MG		Company				
(DAILY),		Representative				

Date:12/08/99ISR Number: 3416552-4Report Type:Expedited (15-DaCompany Report #033-0945-990076
 Age:48 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Deafness	Foreign	Neurontin			
Initial or Prolonged		Health	(Gabapentin)	PS		ORAL
1200 MG (400		Professional				
MG, TID), PER						
ORAL						

Date:12/08/99ISR Number: 3416555-XReport Type:Expedited (15-DaCompany Report #001-0945-991191
 Age:36 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Convulsion	Health	Neurontin Capsules			
500 MG (100		Professional	100 Mg (Gabapentin)	PS		ORAL

MG, AM, 200

MG NOON, 200

MG PM), PER

ORAL

Date:12/09/99ISR Number: 3417947-5Report Type:Direct
Age:44 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	4 DAY	Death	Health	Neurontin 300 Mg	PS		
ONE TID			Professional	Oxycontin	C		

Date:12/09/99ISR Number: 3417961-XReport Type:Expedited (15-DaCompany Report #001-0073-990417
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300		Aspiration Convulsion	Health Professional	Dilantin (Phenytoin Sodium)	PS		ORAL
MG BID), PER		Coronary Artery Disease					
ORAL		Gamma-Glutamyltransferase					
PER ORAL		Increased Grand Mal Convulsion		Neurontin (Gabapentin)	SS		ORAL
		Myocardial Infarction		Lipitor	C		
				Aspirin	C		
				Lescol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/09/99ISR Number: 3417963-3Report Type:Expedited (15-DaCompany Report #033-0945-990069

Age:83 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER ORAL	Balance Disorder Tremor	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Tegretol	C		
			Alepsal	C		
			Digoxin	C		
			Lasilix	C		

Date:12/10/99ISR Number: 3417908-6Report Type:Direct

Company Report #

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 100MG QAM 700MG HS ORAL 800MG TID ORAL	Dermatitis		Quetiapine 100mg Tablets	PS		ORAL
			Gabapentin 400mg Capsules	SS		ORAL
			Fluphenazine	C		
			Fluvastatin	C		

Date:12/10/99ISR Number: 3417918-9Report Type:Direct

Company Report #

Age:69 YR Gender:Female I/FU:I

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

Date:12/10/99ISR Number: 3418341-3Report Type:Direct
Age:52 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 1800 MG DAILY		Erectile Dysfunction	Health Professional	Neurontin	PS		
AS 600MG 3X/DAY							

Date:12/13/99ISR Number: 3421603-7Report Type:Expedited (15-DaCompany Report #001-0945-991226
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (200 Required MG, TID), PER Intervention to ORAL Prevent Permanent Impairment/Damage		Back Pain Confusional State Coordination Abnormal Diplopia Diverticulitis Dry Throat Fatigue Peritonitis Skin Exfoliation Tremor Vision Blurred	Consumer	Neurontin Capsules 100 Mg (Gabapentin) Premarin (Estrogens Conjugated) Vitamin B	PS C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/14/99ISR Number: 3421213-1Report Type:Expedited (15-DaCompany Report #001-0945-991228

Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health	Neurontin			
		Ear Infection	Professional	(Gabapentin)	PS		
1000 MG DAILY				Topamax (Topiramate)	SS		
75 MG DAILY				Unspecified			
				Antibiotic	C		
				Prenatal Vitamins			
				(Ergocalciferol,			
				Ascorbic Acid,			
				Pyridoxine			
				Hydrochloride,	C		
				Folic Acid	C		

Date:12/14/99ISR Number: 3421380-XReport Type:Expedited (15-DaCompany Report #1999COU1453

Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cerebrovascular Accident	Health	Coumadin			
Initial or Prolonged		Convulsion	Professional	(Crystalline			
		Drug Interaction		Warfarin Sodium)	PS		ORAL
3.0-3.5 MG		International Normalised					
DLY PO		Ratio Decreased		Neurontin			
		International Normalised		(Gabapentin)	SS		
		Ratio Increased		Prempro (Conjugated			
		Liver Function Test		Estrogen			
		Abnormal		Medroxyprogesterone			
				Acetate)	C		
				Lanoxin (Digoxin)	C		
				Zoloft (Sertraline			
				Hydrochloride)	C		
				Lipitor			
				(Atorvastatin			
				Calcium)	C		
				Pen-Vee-K			
				(Phenoxymethylpenici			
				llin Potassium)	C		

Darvocet-N
(Di-Gesic) C
Ni (Centrum) C

Date:12/15/99ISR Number: 3422827-5Report Type:Expedited (15-DaCompany Report #032-0945-990023
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthropathy	Foreign	Gabapentin			
PER ORAL		Difficulty In Walking	Health	(Gabapentin)	PS		ORAL
		Movement Disorder	Professional	(Flunitrazepam)	C		
				(Codeine Phosphate, Paracetamol)	C		

Date:12/16/99ISR Number: 3422780-4Report Type:Direct Company Report #
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Rash Erythematous		Neurontin 100mg			
100MG PO HS				Capsule	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Risperdal C
 Zyprexa C
 Docusate C
 Benztropine C
 Ibuprofen C

Date:12/16/99ISR Number: 3423915-XReport Type:Periodic
 Age:48 YR Gender:Female I/FU:I

Company Report #001-0981-997129

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 20 MG (DAILY), PER ORAL		Confusional State Jaundice Liver Function Test Abnormal	Consumer	Lipitor Tablets 20 Mg (Atorvastatin)	PS		ORAL
300 MG QAM, 600 MG QPM), PER ORAL		Nausea Urinary Tract Infection Vomiting Weight Decreased		Neurontin Capsules 300 Mg (Gabapentin)	SS		ORAL
				Lortab (Paracetamol, Hydrocodone Bitartrate) Clonidine Xanax (Alprazolam) Imitrex (Sumatriptan Succinate) Folic Acid Magnesium	C C C C C C		

Date:12/21/99ISR Number: 3427109-3Report Type:Expedited (15-DaCompany Report #033-0945-990080
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNK, UNKNOWN		Cardiac Disorder	Foreign Study	Gabapentin (Gabapentin)	PS		

Health
Professional

Date:12/22/99ISR Number: 3428503-7Report Type:Expedited (15-DaCompany Report #001-0945-991232
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Health	Neurontin Capsules			
200 MG		Drug Level Below	Professional	100 Mg (Gabapentin)	PS		ORAL
(DAILY), PER		Therapeutic					
ORAL				Cyclosporin A (Ciclosporin)	SS		

Date:12/22/99ISR Number: 3428967-9Report Type:Expedited (15-DaCompany Report #001-0945-990986
Age:42 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Akinesia
Initial or Prolonged	Convulsion
Other	Crying
	Depression
	Difficulty In Walking

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, BID), PER ORAL		Dysarthria Feeling Abnormal Mood Swings Nervousness Pain Palpitations Salivary Hypersecretion Screaming Suicidal Ideation Tremor Vision Blurred	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

Date:12/23/99ISR Number: 3429041-8Report Type:Direct
Age:21 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300MG BID		Pruritus		Neurontin 300mg Synthroid	PS C		

Date:12/23/99ISR Number: 3429829-3Report Type:Expedited (15-DaCompany Report #991217-SK111
Age:91 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - UNKNOWN PO Initial or Prolonged 300.000 MG OD Other PO / YEARS		Bronchitis Convulsion Drug Interaction	Health Professional	Celebrex Neurontin	PS SS		ORAL ORAL

Date:12/23/99ISR Number: 3430269-1Report Type:Expedited (15-DaCompany Report #033-0945-990082
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Genital Disorder Female	Foreign Study Health Professional	Gabapentin (Gabapentin)	PS		

Date:12/23/99ISR Number: 3430270-8Report Type:Expedited (15-DaCompany Report #032-0945-990019
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL			Fall Psychotic Disorder	Foreign Health Professional Company Representative	Gabapentin (Gabapentin) (Vigabatrin)	PS C		ORAL

Date:12/23/99ISR Number: 3430271-XReport Type:Expedited (15-DaCompany Report #001-0945-991048
Age: Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 400 MG (DAILY)		1 MON	Abortion Spontaneous Benign Hydatidiform Mole	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/27/99ISR Number: 3431501-0Report Type:Expedited (15-DaCompany Report #001-0945-991087
 Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Ventricular Fibrillation	Health Professional	Neurontin (Gabapentin)	PS		
300 MG (100 MG, TID),			Company Representative				
UNKNOWN	2 DAY			Betapace (Sotalol Hydrochloride)	C		

Date:12/29/99ISR Number: 3433678-XReport Type:Direct Company Report #
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Deafness		Neurontin	PS		
900MG DAILY							
SEE ITEM B5		Dialysis		Procardia	C		
				Clonidine	C		
				Nephrocaps	C		
				Lipitor	C		

Date:12/29/99ISR Number: 3435167-5Report Type:Expedited (15-DaCompany Report #001-0945-991279
 Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Grand Mal Convulsion Intentional Misuse	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1)PER ORAL;2)							
Other		Lethargy					
3000 MG (AS ONE DOSE),		Loss Of Consciousness					
PER ORAL; 3)		Suicide Attempt					

PER ORAL

Wellbutrin
(Amfebutamone
Hydrochloride) SS ORAL

1) PER ORAL;

2) 4500 MG

(AS ONE

DOSE), PER

ORAL; 3) PER

Date:12/29/99ISR Number: 3435201-2Report Type:Expedited (15-DaCompany Report #001-0945-991179
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1) 400 MG (DAILY), PER ORAL; 2) 800 MG (400 MG, BID), PER 3.75 MG (DAILY), PER ORAL		Abdominal Pain Drug Interaction Gastrointestinal Haemorrhage International Normalised Ratio Increased Loss Of Consciousness Platelet Count Decreased	Health Professional	Neurontin Capsules 400 Mg (Gabapentin) Coumadin (Warfarin Sodium) (Insulin)	PS SS C		ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/30/99ISR Number: 3434744-5Report Type:Expedited (15-DaCompany Report #001-0945-990919
 Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Health Professional	Neurontin Capsule 300 Mg (Gabapentin)	PS		ORAL
3600 (1200 MG, TID), PER ORAL							

Date:01/05/00ISR Number: 3436705-9Report Type:Direct Company Report #
 Age:89 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Posture Abnormal		Neurontin 600mgm	PS		ORAL
600MGM BID PO							
		Tremor		Fv Natural Tears Drops	C		
				Synthroid	C		
				Furosemide	C		
				Zoloft	C		
				Prinivil	C		
				Multivitamins	C		
				K-Dur	C		
				Detrol	C		
				Docusate Sod	C		
				Pentoxifylline	C		
				Metamucil Powder	C		
				Baclofen	C		
				Ranitidine	C		
				Premarin	C		
				Xylocaine	C		
				Zoloft	C		

Date:01/05/00ISR Number: 3437848-6Report Type:Expedited (15-DaCompany Report #001-0945-990197
 Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged 800 MG (200 MG, QID), UNKNOWN	Chest Pain Flushing Insomnia Oesophageal Spasm Pain Peripheral Coldness	Health Professional	Neurontin (Gabapentin)	PS
			Glyburide (Glibenclamide) Paxil (Paroxetine Hydrochloride) Zyprexa (Olanzapine) Levoxine (Levothyroxine Sodium) Trazodone Ambien (Zolpidem Tartrate)	C C C C

Date:01/06/00ISR Number: 3439277-8Report Type:Expedited (15-DaCompany Report #033-0945-990085
Age:67 YR Gender:Female I/FU:I

Outcome Hospitalization - Initial or Prolonged	PT Agranulocytosis Leukopenia Neutropenia	Report Source Foreign 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
1200 MG			Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER						
ORAL			Phenytoin Depakine (Valproate Sodium)	SS SS		

Date:01/07/00ISR Number: 3440278-4Report Type:Expedited (15-DaCompany Report #033-0945-990088
Age:87 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE IMAGE	10 DAY	Respiratory Distress Sedation	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Sleep Apnoea Syndrome	Professional	Lovenox (Heparin-F-Raction, Sodium Salt)	C		
				Bricanyl (Terbutal-Ine Sulfate)	C		
				Lasilix (Furosemid)	C		
				Atrovent (Ipratropium Bromide)	C		

Date:01/11/00ISR Number: 3442691-8Report Type:Expedited (15-DaCompany Report #001-0073-990417
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300		Aspiration Convulsion	Health Professional	Dilantin (Phenytoin Sodium)	PS		ORAL

MG BID)

Coronary Artery Disease

Gamma-Glutamyltransferase
Increased
Grand Mal Convulsion
Myocardial Infarction
Quadruple Vessel Bypass
Graft

Neurontin
(Gabapentin) SS
Lipitor
(Atorvastatin) C
Aspirin
(Acetylsalicylic
Acid) C
Lescol (Fluvastatin
Sodium) C

ORAL

Date:01/11/00ISR Number: 3442693-1Report Type:Expedited (15-DaCompany Report #033-0945-990089

Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (DAILY)		Balance Disorder Electroencephalogram Abnormal Encephalopathy Memory Impairment Mental Disorder	Foreign Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Hept-A-Myl (Heptamino l Hydrochloride) Seropram (Citalopram Hydrobromide)	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/11/00ISR Number: 3442696-7Report Type:Expedited (15-DaCompany Report #001-0945-M000001

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 MG, TID)	Coma Depressed Level Of Consciousness	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
75 MG (Q AM)			Effexor (Venlafaxine Hydrochloride)	SS		ORAL
			Imdur (Isosorbide Mononitrate)	C		
			Prilosec (Omeprazole)	C		
			Pulmicort Inhaler (Budesonide)	C		
			Permax (Pergolide Mesilate)	C		
			Serzone (Nefazodone Hydrochloride)	C		
			Oramorph (Morphine Sulfate)	C		
			Estrace (Estradiol)	C		
			Synthroid (Levothyroxine Sodium)	C		
			Skelaxin (Metaxalone)	C		
			Vicodin (Paracetamol, Hydrocodone Bitartrate)	C		

Date:01/12/00ISR Number: 3443156-XReport Type:Direct

Age:44 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG PO TID 3 MON	Depression Libido Decreased Sedation		Neurontin	PS		ORAL
			Compazine	C		
			Imipramine	C		

Tremor
Weight Increased

Levoxy1

C

Date:01/12/00ISR Number: 3443414-9Report Type:Expedited (15-DaCompany Report #001-0945-990475
Age:8 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Gastric Ulcer
Other	Gastritis
	Glycosuria
	Haematochezia
	Malaise
	Nephritis Interstitial
	Nephrolithiasis
	Proteinuria
	Red Blood Cell
	Sedimentation Rate
	Increased
	Uveitis
	Vomiting

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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
		Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		

Date:01/12/00ISR Number: 3443416-2Report Type:Expedited (15-DaCompany Report #044-0945-M000001
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 80 CAPSULE			Foreign Health Professional	Neurontin (Gabapentin)	PS		
		Loss Of Consciousness Suicidal Ideation	Professional	Carbamazepine (Carbamazepine)	SS		

Date:01/13/00ISR Number: 3443614-8Report Type:Expedited (15-DaCompany Report #001-0945-991308
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged DAILY (AT Other BEDTIME) UNKNOWN		Condition Aggravated Dysmenorrhoea	Health Professional	Neurontin (Gabapentin)	PS		
		Endometriosis Menorrhagia					

Date:01/18/00ISR Number: 3444902-1Report Type:Expedited (15-DaCompany Report #033-0945-990085
Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (DAILY), PER		Agranulocytosis Leukopenia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Neutropenia	Professional				

ORAL

(Phenytoin) SS
Depakine (Valproate Sodium) SS

Date:01/18/00ISR Number: 3445011-8Report Type:Expedited (15-DaCompany Report #001-0945-M000009
Age:91 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (DAILY), PER		Bronchitis Convulsion	Study Health Professional	Neurontin Capsules 300 Mg(Gabapentin)	PS		ORAL

ORAL/ YEARS YR

Celebrex (Celecoxib) SS ORAL

PER ORAL

Date:01/19/00ISR Number: 3446467-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000032
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 1500 MG (300 MG, FIVE TIMES A DAY),		Myocardial Infarction	Health Professional Company Representative	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

PER ORAL

Rocephin (Lidocaine

Freedom Of Information (FOI) Report

INTRAMUSCULAR (ONE DOSE),
 INTRAMUSCULAR

Hydrochloride,
 Ceftriaxone Sodium) SS

Biaxin
 (Clarithromycin) SS
 Soma (Carisoprodol) C
 Ultram (Tramadol
 Hydrochloride) C

Date:01/20/00ISR Number: 3445368-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000022
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Consumer	Neurontin			
Other		Suicidal Ideation		(Gabapentin)	PS		

Date:01/21/00ISR Number: 3445655-3Report Type:Expedited (15-DaCompany Report #A000496
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	150.00 MG	Alopecia	Health	Diflucan	PS		ORAL
Required	TOTAL:DAILY:0	Asthenia	Professional				
Intervention to	RAL	Dry Skin					
Prevent Permanent	ORAL	Fatigue		Neurontin	SS		ORAL
Impairment/Damage		Feeling Cold		Lipitor-"Started			
		Formication		Several Years Ago"	C		
		Hyperinsulinism		Insulin	C		
		Paraesthesia					
		Sedation					

Date:01/21/00ISR Number: 3446297-6Report Type:Expedited (15-DaCompany Report #032-0945-990023
 Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthropathy	Foreign	Gabapentin			
Other		Difficulty In Walking	Health	(Gabapentin)	PS		ORAL
300 MG		Haematoma	Professional				
(DAILY), PER		Insomnia					
ORAL		Movement Disorder		(Flunitrazepam)	C		
				(Codeine Phosphate, Paracetamol)	C		

Date:01/24/00ISR Number: 3446735-9Report Type:Expedited (15-DaCompany Report #001-0945-970529
Age:10 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Aggression	Consumer	Neurontin			
Initial or Prolonged		Convulsion		(Gabapentin)	PS		ORAL
300 MG		Irritability					
Other		Mania					
(DAILY), PER		Panic Attack		Depakote (Valproate			
ORAL		Psychomotor Hyperactivity		Semisodium)	C		
		Suicide Attempt		Wellbutrin			
				(Amfebutamone			
				Hydrochloride)	C		
				Zinc	C		
				Multivatimn			
				(Ergocalciferol,			

Freedom Of Information (FOI) Report

Ascorbic Acid, Folic
Acid, Thiamine
Hydrochloride, C

Date:01/24/00ISR Number: 3446771-2Report Type:Expedited (15-DaCompany Report #001-0945-990846
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:01/24/00ISR Number: 3446901-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000038
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Corneal Oedema Visual Acuity Reduced	Consumer Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
900 MG (300 MG TID) PER ORAL							

Date:01/24/00ISR Number: 3528453-1Report Type:Periodic Company Report #990924010015
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
RESPIRATORY (INHALATION) INH MG	300 MG, BID,	Asthenia Diarrhoea Hyperhidrosis Peripheral Vascular Disorder Vasodilatation	Consumer Health Professional	Tobi Gabapentin (Gabapentin) Nelfinavir	PS SS C	Pathogenesis Corp	

Alprazolam
 (Alprazolam) C
 Doxepin (Doxepin) C
 Oxycontin (Oxycodone
 Hydrochloride) C

Date:01/27/00ISR Number: 3447340-0Report Type:Expedited (15-DaCompany Report #001-0945-991232
 Age:71 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Health	Neurontin Capsules			
200 MG (100		Drug Level Below	Professional	100 Mg (Gabapentin)	PS		ORAL
MG, BID), PER		Therapeutic					
ORAL		Fluid Overload					
		Renal Impairment		Cyclosporin A (Ciclosporin)	SS		

Date:01/27/00ISR Number: 3447378-3Report Type:Expedited (15-DaCompany Report #033-0945-990065
 Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Bilirubin Increased	Foreign	Neurontin			
Initial or Prolonged		Hepatitis	Health	(Gabapentin)	PS		ORAL
1200 MG DAILY		Liver Function Test	Professional				
PER ORAL		Abnormal		Rivotril			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Clonazepam) C

Date:01/27/00ISR Number: 3447391-6Report Type:Expedited (15-DaCompany Report #225690
 Age:35 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL	Confusional State Hallucination, Visual	Foreign Other	Rivotril (Clonazepam)	PS		ORAL
ORAL	Medication Error		Tardyferon (Ferrous Sulfate)	SS		ORAL
100 MG 1 PER 1 DAY ORAL			Tranxene (Clorazepate Dipotassium)	SS		ORAL
600 MG 1 PER 1 DAY ORAL			Leponex (Clozapine)	SS		ORAL
ORAL			Mopral (Omeprazole)	SS		ORAL
200 MG 1 PER 1 DAY ORAL			Neurontin (Gabapentin)	SS		ORAL

Date:01/27/00ISR Number: 3447413-2Report Type:Expedited (15-DaCompany Report #033-0945-990080
 Age:70 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (DAILY),	Cardiac Disorder	Foreign Study	Gabapentin (Gabapentin)	PS		
		Health				
		Professional	Vasten (Pravastatin Sodium) Aspegic	C		

(Acetylsalicylate
 Lysine) C
 Mopral (Omeprazole) C
 Atarax (Hydroxyzine
 Hydrochloride) C
 Temesta (Lorazepam) C
 Lescol (Fluvastatin
 Sodium) C

Date:01/28/00ISR Number: 3448298-0Report Type:Expedited (15-DaCompany Report #001-0945-990219
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Optic Neuropathy	Health	Neurontin Capsules			
		Visual Acuity Reduced	Professional	300 Mg (Gabapentin)	PS		ORAL
900 MG (300							
MG, TID), PER							
ORAL							

Date:01/31/00ISR Number: 3448775-2Report Type:Expedited (15-DaCompany Report #991217-SK111
 Age:91 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Bronchitis	Health	Celebrex	PS		ORAL
PO							
Initial or Prolonged		Convulsion	Professional	Neurontin	SS		ORAL
300.000 MG QD							
Other		Drug Interaction					
PO	YR						

Other	Cognitive Disorder	Consumer	Neurontin Capsules		
1200 MG (300	Cystitis		300 Mg (Gabapentin)	PS	ORAL
MG, QID), PER	Pain In Extremity				
ORAL	Renal Failure				
	Rhabdomyolysis		Unspecified Medication	C	

Date:02/07/00ISR Number: 3454025-3Report Type:Expedited (15-DaCompany Report #032-0945-M0000002
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Mental Disorder	Foreign Study	Gabapentin (Gabapentin)	PS		ORAL
			Health Professional				

Date:02/07/00ISR Number: 3454086-1Report Type:Expedited (15-DaCompany Report #001-0945-991025
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (DAILY)		Dizziness Dysarthria Urinary Incontinence	Consumer	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Percocet
 (Paracetamol,
 Oxycodone
 Hydrochloride,
 Oxycodone C
 Xanax (Alprazolam) C

Date:02/07/00ISR Number: 3454386-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000074
 Age:74 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG (QHS), Other PER ORAL	Blood Creatine Increased Blood Urea Increased Coma Lethargy Toxic Epidermal Necrolysis	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
			Synthroid (Levothyroxine Sodium) Ecotrin (Acetylsalicyclic Acid) Trental (Pentoxifylline) Diovan (Valsartan) Trandate (Labetalol Hydrochloride)	C C C C		

Date:02/07/00ISR Number: 3454396-8Report Type:Expedited (15-DaCompany Report #001-0945-991063
 Age:56 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER ORAL	Cerebrovascular Accident Chills Costochondritis Dysarthria Eye Pain	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
			Lotrel (Benazepril,			

1 TABLET (S)	Face Oedema	Amlodipine)	SS	ORAL
(DAILY), PER	Headache			
ORAL	Hyperhidrosis			
	Hypersensitivity	Glyburide(Glibenclam		
	Hypertension	ide)	C	
	Palpitations	Asa (Acetylsalicylic		
	Vision Blurred	Acid)	C	

Date:02/08/00ISR Number: 3454305-1Report Type:Direct Company Report #
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Congenital Anomaly	Calcinosis		Klonopin	PS		
	Congenital Anomaly		Paxil	SS		
	Nervous System Disorder		Depakote	SS		
			Gabapentin	SS		
			Wellbutrin	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/08/00ISR Number: 3454394-4Report Type:Direct
 Age:54 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Leukopenia Neutropenia		Gabapentin 300mg (Parke-Davis)	PS	Parke-Davis	ORAL
300MG 1T PO							
TID				Bactrim Ds	SS		ORAL
1T PO				Zantac	SS		ORAL
PO TID				Megace	C		
				Roxicodone	C		
				Pogy	C		

Date:02/09/00ISR Number: 3455426-XReport Type:Expedited (15-DaCompany Report #032-0945-990023
 Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthralgia	Foreign	Gabapentin	PS		ORAL
Other		Arthropathy	Health	(Gabapentin)			
300MG							
(DAILY), PER		Depression	Professional				
ORAL		Difficulty In Walking					
		Haematoma		Flunitrazepam	C		
		Insomnia		Codeine Phosphate,			
		Muscle Injury		Paracetamol	C		
		Oedema Peripheral					

Date:02/10/00ISR Number: 3456148-1Report Type:Expedited (15-DaCompany Report #00P-056-0086712-00(0)
 Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Confusional State Hallucination, Visual	Foreign Health	Tranxene (Tranxene) (Clorazepate			

100 MG, 1 IN	Professional	Dipotassium)	PS	ORAL
1 D, PER ORAL				
PER ORAL		Clonazepam (Clonazepam)	SS	ORAL
PER ORAL		Ferrous Sulfate (Ferrous Sulfate)	SS	ORAL
600 MG, 1 IN		Clozapine (Clozapine)	SS	ORAL
1 D, PER ORAL				
PER ORAL		Omeprazole (Omeprazole)	SS	ORAL
200 MG, 1 IN		Gabapentin (Gabapentin)	SS	ORAL
1 D, PER ORAL				

Date:02/11/00ISR Number: 3457020-3Report Type:Expedited (15-DaCompany Report #033-0945-M0000015
Age:34 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged PER ORAL	Agranulocytosis Pyrexia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
PER ORAL			Tegretol (Carbamazepine)	SS		ORAL
PER ORAL			Zocor (Simvastatin)	SS		ORAL
PER ORAL			Bi-Profenid (Ketoprofen)	SS		ORAL
PER ORAL			Lexomil (Bromazepam)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

PER ORAL
 Zoltum (Omeprazole) SS ORAL
 Chemotherapy SS

Date:02/14/00ISR Number: 3457402-XReport Type:Direct Company Report #
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma	Health	Zyprexa 15 Mg/Day	PS		
15MG/DAY		Muscle Rigidity	Professional	Lithobid 600 Mg Bid	SS		
600 MG BID		Pyrexia		Neurontil 600mg Tid	SS		
600MG TID		Tremor		Symmetrel 900mg/Day	SS		
900MG/DAY							

Date:02/14/00ISR Number: 3459086-3Report Type:Periodic Company Report #001-0945-990364
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Deafness	Health	Neurontin			
Other		Tinnitus	Professional	(Gabapentin)	PS		ORAL
1200 MG (QHS)		Vertigo	Company				
, PER ORAL			Representative				

Date:02/14/00ISR Number: 3459088-7Report Type:Periodic Company Report #001-0945-990384
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Bronchospasm	Health	Neurontin			
1800 MG		Dyspnoea	Professional	(Gabapentin)	PS		ORAL
(DAILY), PER		Pulmonary Embolism					

ORAL
Pulmonary Oedema
Pulmonary Thrombosis
Snoring

Date:02/14/00ISR Number: 3459089-9Report Type:Periodic Company Report #001-0945-990404
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Blindness	Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
400 MG (200 MG, BID) , PER ORAL				Lotensin (Benazepril Hydrochloride) (Digoxin) Nph Insulin (Insulin Injection, Isophane) (Acetylsalicylic Acid) Vitamin B12 (Cyanocobalamin)	C C C C C		

Date:02/14/00ISR Number: 3459090-5Report Type:Periodic Company Report #001-0945-990405
Age:40 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Blood Thyroid Stimulating Hormone Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Cardiac Failure Congestive	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
400 MG	(DAILY), PER						
ORAL				(Lithium)	SS		
				Depakote (Valproate Semisodium)	C		
				Zyprexa (Olanzapine)	C		
				Luvox (Fluvoxamine Maleate)	C		

Date:02/14/00ISR Number: 3459091-7Report Type:Periodic Company Report #001-0945-990411
 Age: Gender:Male I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Health Professional	Neurontin (Gabapentin)	PS		
Dose		Confusional State Depression					
Hospitalization - Initial or Prolonged 1500 MG		Drug Interaction					
(DAILY) , /		Hallucination					
MONTHS AGO		Mood Swings		Haldol (Haloperidol)	SS		
5 MG ((DAILY)							

Date:02/14/00ISR Number: 3459092-9Report Type:Periodic Company Report #001-0945-990441
 Age:67 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
Dose		Insomnia Pollakiuria					
Other		Weight Decreased					
100 MG (QHS),							
PER ORAL							

(Amitriptyline)

C

Date:02/14/00ISR Number: 3459093-0Report Type:Periodic
Age:50 YR Gender:Male I/FU:I

Company Report #001-0945-990472

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1600 MG (400 MG, QID), PER ORAL		Confusional State Drug Toxicity Encephalopathy Sedation	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL

(Baclofen)	C
(Atenolol)	C
(Cimetidine)	C
(Ranitidine)	C
Oxybutynin)	C
(Ferrous Sulfate)	C

Date:02/14/00ISR Number: 3459094-2Report Type:Periodic
Age:66 YR Gender:Female I/FU:I

Company Report #001-0945-990511

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (300 MG, BID) , PER ORAL		Breast Cancer Female	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

Premarin (Estrogens)

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Conjugated) C
 Advil (Ibuprofen) C

Date:02/14/00ISR Number: 3459095-4Report Type:Periodic
 Age:77 YR Gender:Female I/FU:I

Company Report #001-0945-990591

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300		Thrombocytopenia	Health Professional	Neurontin Capsules 300 Mg(Gabapentin)	PS		ORAL

MG, TID), PER

ORAL

Cardizem (Diltiazem
 Hydrochloride) C
 Prevacid
 (Lansoprazole) C
 Toprol Xl
 (Metoprolol
 Succinate) C
 Senokot (Senna
 Fruit) C
 Accupril (Quinapril
 Hydrochloride) C

Date:02/14/00ISR Number: 3459096-6Report Type:Periodic
 Age:56 YR Gender:Female I/FU:I

Company Report #001-0945-990641

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1200 MG (300		Cerebral Atrophy	Health Professional	Neurontin Capsules 300 Mg(Gabapentin)	PS		ORAL

MG, 2 QAM,

10PMHS), PER

ORAL

Celebrex (Celecoxib) C
 Xanax (Alprazolam) C
 Premarin (Estrogens)

Conjugated) C
Parafon Forte
(Chlorzoxazone,
Paracetamol) C
Armour Thyroid
(Thyroid) C

Date:02/14/00ISR Number: 3459099-1Report Type:Periodic Company Report #001-0945-990679
Age:36 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2400 MG (800 MG, QID) , PER ORAL	Acute Respiratory Distress Syndrome	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Antibiotic	C		

Date:02/14/00ISR Number: 3459101-7Report Type:Periodic Company Report #001-0945-990744
Age:47 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

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4800 MG (1200 MG, QID), PER ORAL / 3200 MG (400 MG, QID), PER		Abdominal Pain Accidental Overdose Overdose Palpitations Road Traffic Accident Syncope Vision Blurred Vomiting	Health Professional	Neurontin Capsules 400 Mg(Gabapentin)	PS		ORAL

Date:02/14/00ISR Number: 3459108-XReport Type:Periodic
Age:56 YR Gender:Female I/FU:I

Company Report #001-0945-990762

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 MG, BID)		Pancreatitis Urinary Tract Infection	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		
				Coumadin (Warfarin Sodium)	C		
				Ativan (Lorazepam)	C		
				Levoxyl (Levothyroxine Sodium)	C		
				(Propranolol)	C		
				Cortef (Hydrocortisone Acetate)	C		
				Azmacort Inhaler (Triamcinolone Acetonide)	C		
				Albuterol Inhaler (Salbutamol)	C		
				Premarin (Estrogens Conjugated)	C		

Colace (Docusate Sodium) C
Tenormin (Atenolol) C

Date:02/14/00ISR Number: 3459110-8Report Type:Periodic Company Report #001-0945-990770
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Accidental Overdose	Consumer	Neurontin Capsules			
SEE IMAGE ,		Amnesia		300 Mg (Gabapentin)	PS		ORAL
PER ORAL		Constipation					
		Disturbance In Attention		(Trazodone)	C		
		Oedema		Valium (Diazepam)	C		
		Skin Exfoliation		Percocet (Paracetamol, Oxycodone Hydrochloride, Oxycodone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/00ISR Number: 3459111-XReport Type:Periodic
Age:32 YR Gender:Female I/FU:I

Company Report #001-0945-990771

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cataract	Health	Neurontin			
1600 MG (800		Condition Aggravated	Professional	(Gabapentin)	PS		ORAL
MG BID) (800		Sedation					
MG), PER ORAL				(Lithium)	C		
				(Clozapine)	C		
				Colace (Docusate			
				Sodium)	C		

Date:02/14/00ISR Number: 3459112-1Report Type:Periodic
Age:84 YR Gender:Female I/FU:I

Company Report #001-0945-990794

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Amnesia	Consumer	Neurontin Capsules			
Initial or Prolonged		Confusional State		300 Mg (Gabapentin)	PS		ORAL
300 MG (QHS)		Pneumonia					
, PER ORAL				Zocor (Simvastatin)	C		
				Synthroid			
				(Levothyroxine			
				Sodium)	C		
				(Trazodone)	C		
				Colace (Docusate			
				Sodium)	C		
				Paxil (Paroxetine			
				Hydrochloride)	C		

Date:02/14/00ISR Number: 3459113-3Report Type:Periodic
Age:75 YR Gender:Female I/FU:I

Company Report #001-0945-990839

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Confusional State	Consumer	Neurontin Capsules			

Initial or Prolonged 2700 MG (900 MG, TID), PER ORAL	Mental Retardation Severity Unspecified Pneumonia Sedation	300 Mg (Gabapentin)	PS	ORAL
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Tegretol (Carbamazepine)	C
Unspecified Blood Pressure Medication	C
Vicodin (Paracetamol, Hydrocodone Bitartrate)	C

Date:02/14/00ISR Number: 3459115-7Report Type:Periodic Company Report #001-0945-990888
Age:80 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged PER ORAL	Abdominal Pain Blood Bilirubin Increased Hepatic Enzyme Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/00ISR Number: 3459117-0Report Type:Periodic
Age:40 YR Gender:Female I/FU:I

Company Report #001-0945-990903

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Pain	Health	Neurontin			
SEE IMAGE		Vision Blurred	Professional	(Gabapentin)	PS		ORAL
PER ORAL		Visual Acuity Reduced					
				Premarin (Estrogens Conjugated)	C		
				Tylenol (Paracetamol)	C		

Date:02/14/00ISR Number: 3459119-4Report Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-990909

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Convulsion Hypersensitivity	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/14/00ISR Number: 3459500-3Report Type:Periodic
Age:44 YR Gender:Female I/FU:I

Company Report #001-0945-991218

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged TITRATED UP		Asthenia Back Pain	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
TO A DAILY (DOSE OF 2700MG), PER ORAL; SEE		Cardiac Failure Coordination Abnormal Dizziness Fall Headache Nausea		Lasix (Furosemide) Oxycontin (Oxycodone)	C		

Hydrochloride) C
 Wellbutrin
 (Amfebutamone) C
 Soma (Carisoprodol) C
 Pamelor
 (Nortriptylne
 Hydrochloride) C
 Micro-K (Potassium
 Chloride) C

Date:02/14/00ISR Number: 3459504-0Report Type:Periodic Company Report #001-0945-991219
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Deafness	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG, TID), PER							ORAL

Darvocet
 (Paracetamol,
 Dextropropoxyphene) C
 Prilosec
 (Omeprazole) C
 Promethazine C
 Lactulose C
 Levothyroid

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Levothyroxine) C
 Temazepam C
 Prevacid
 (Lansoprazole) C
 K-Dur (Potassium
 Chloride) C
 Neomycin C

Date:02/14/00ISR Number: 3459507-6Report Type:Periodic Company Report #001-0945-991268
 Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Neurontin (Gabapentin)	PS		ORAL
2400-2700 MG							
(DAILY), PER							
ORAL				Depakote (Valproate Semisodium)	C		

Date:02/14/00ISR Number: 3459511-8Report Type:Periodic Company Report #001-0945-991308
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dysmenorrhoea Menorrhagia	Health Professional	Neurontin (Gabapentin)	PS		
3800 MG							
(DAILY)		Overdose					

Date:02/14/00ISR Number: 3459514-3Report Type:Periodic Company Report #001-0945-991310
 Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hallucination Insomnia	Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG							

Logorrhoea
 (DAILY), PER
 Restlessness
 ORAL
 Vitamins C
 Unspecified Water
 Pill C

Date:02/14/00ISR Number: 3459516-7Report Type:Periodic Company Report #001-0945-980487
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE TEXT, PER		Arthralgia Convulsion	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
ORAL		Dermatitis					
250 MG		Dyspnoea		Peganone (Ehtotoin)	SS		ORAL
(DAILY), PER		Mental Disorder					
ORAL		Overdose					
		Stupor Syncope		Zoloft (Sertraline Hydrochloride) Tranxene (Clorazepam Dipotassium) Haldol (Haloperidol) Felbatol (Felbamate) Depakote (Valproate Sodium) Gabitril (Tiagabine)	C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Topamax (Topiramate) C
 Lamictal
 (Lamotrigine) C

Date:02/14/00ISR Number: 3459524-6Report Type:Periodic
 Age:28 YR Gender:Female I/FU:F

Company Report #001-0945-980710

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER		Convulsion Drug Withdrawal Convulsions	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL				Depakote (Valproate Sodium)	SS		ORAL
250 MG (TOOK ONE MG, TID) PER ORAL				Buspar (Buspirone Hydrochloride) Zoloft (Sertraline Hydrochloride) Dilantin (Phenytoin Sodium) Advil (Ibuprofen)	C C C C		

Date:02/14/00ISR Number: 3459550-7Report Type:Periodic
 Age:58 YR Gender:Female I/FU:I

Company Report #001-0945-980831

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 MG, TID), PER		Pancreatitis Post Procedural Complication	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
ORAL							

Vicodin
(Paracetamol,
Hydrocodone
Bitartrate) C
Prilosec
(Omeprazole) C

Date:02/14/00ISR Number: 3459553-2Report Type:Periodic Company Report #001-0945-980872
Age:40 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Suicidal Ideation	Health Professional	Neurontin (Gabapentin)	PS		

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
300 MG (150 MG,BID), PER			Zyban (Amfebutamone)	SS		ORAL

ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/00ISR Number: 3459570-2Report Type:Periodic
Age:35 YR Gender:Male I/FU:F

Company Report #001-0945-990003

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG (300 MG, BID), PER ORAL	Epistaxis Haemoglobin Decreased Headache Hypersensitivity Photophobia Thrombocytopenia	Health Professional Company Representative	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
			Aciclovir Prilosec (Omeprazole) Tylenol (Paracetamol) Imitrex (Sumatriptan Succinate) St. John'S Wort (Hypericum Extract) Vitamin C (Ascorbic Acid) Papain	C C C C C C C		

Date:02/14/00ISR Number: 3459580-5Report Type:Periodic
Age:70 YR Gender:Male I/FU:I

Company Report #001-0945-990011

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (DAILY), PER ORAL	Dyspnoea Liver Function Test Abnormal	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
			Coumadin (Warfarin Sodium) (Insulin) Zoloft (Sertraline Hydrochloride) K-Dur (Potassium Chloride) Lanoxin (Digoxin)	C C C C C		

Lotrel (Benazepril,
Amlodipine) C
Demadex (Torasemide) C

Date:02/14/00ISR Number: 3459583-0Report Type:Periodic
Age:42 YR Gender:Female I/FU:I

Company Report #001-0945-990059

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Lipids Abnormal	Health	Neurontin Capsules			
3200 MG		Nephrotic Syndrome	Professional	400 Mg (Gabapentin)	PS		ORAL

(DAILY), PER

ORAL

Synthroid
(Levothyroxine
Sodium) C
Norvasc (Amlodipine
Besilate) C
Tegretol
(Carbamazepine) C
Wellbutrin
(Amfebutamone
Hydrochloride) C
Motrin (Ibuprofen) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/00ISR Number: 3459587-8Report Type:Periodic
Age:23 YR Gender:Female I/FU:I

Company Report #001-0945-990063

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER ORAL		Bipolar Disorder Condition Aggravated	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				(Lithium)	SS		
				Depakote (Valproate Semisodium)	SS		
				Vistaril (Hydroxyzine Embonate)	SS		
				Trazodone)	SS		
				Synthroid (Levothyroxine Sodium)	C		
				Clonidine	C		

Date:02/14/00ISR Number: 3459588-XReport Type:Periodic
Age:48 YR Gender:Female I/FU:I

Company Report #001-0945-990067

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (DAILY), PER ORAL		Asthenia Coordination Abnormal Dizziness Hypersensitivity Tremor Vomiting	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Synthroid (Levothyroxine Sodium)	C		
				Voltaren (Diclofen-Ac Sodium)	C		
				Nortriptyline	C		
				(Echinacea Extract)	C		
				L-Lysine (Lysine)	C		
				Vitamin E			

(Tocopherol) C
(Calcium) C

Date:02/14/00ISR Number: 3459594-5Report Type:Periodic Company Report #001-0945-990068
Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	White Blood Cell Count	Health	Neurontin Capsules			
Initial or Prolonged	Decreased	Professional	100 Mg (Gabapentin)	PS		ORAL
200 MG (100						
MG, BID), PER						
ORAL						

Motrin (Ibuprofen) C

Date:02/14/00ISR Number: 3459609-4Report Type:Periodic Company Report #001-0945-990073
Age:49 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion	Health	Neurontin			
Initial or Prolonged	Drug Interaction	Professional	(Gabapentin)	PS		
300 MG (100						
MG, TID),						
			Ginkgo Biloba	SS		
			Serzone (Nefazodone			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C

Date:02/14/00ISR Number: 3459611-2Report Type:Periodic
Age:44 YR Gender:Female I/FU:I

Company Report #001-0945-990106

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Agitation Drug Level Above Therapeutic Feeling Drunk Nausea	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (DAILY), PER ORAL				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:02/14/00ISR Number: 3459615-XReport Type:Periodic
Age:18 YR Gender:Female I/FU:I

Company Report #001-0945-990118

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other	Convulsion	Consumer	Neurontin		
900-1100 MG	Delusion	Health	(Gabapentin)	PS	ORAL
(DAILY), PER	Depression	Professional			
ORAL	Suicide Attempt				

Marijuana (Cannabis)	SS
Unspecified Birth	
Control Pills	C
Prozac (Fluoxetine	
Hydrochloride)	C

Date:02/14/00ISR Number: 3459618-5Report Type:Periodic Company Report #001-0945-990129
 Age:86 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Hallucination	Consumer	Neurontin Capsules			
Initial or Prolonged			300 Mg (Gabapentin)	PS		ORAL
2400 MG (600						

MG, QID), PER

ORAL

Cardizem (Diltiazem	
Hydrochloride)	C
Lanoxin (Digoxin)	C
(Acetylsalicylic	
Acid)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/00ISR Number: 3459620-3Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-990159

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pancreatitis	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/14/00ISR Number: 3459624-0Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-990166

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pericardial Effusion	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/14/00ISR Number: 3459627-6Report Type:Periodic
Age:42 YR Gender:Female I/FU:I

Company Report #001-0945-990215

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Hot Hypersensitivity	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		
300 MG		Sinus Headache		Methocarbamol Naprelan (Naproxen Sodium)	C C		

Date:02/14/00ISR Number: 3459631-8Report Type:Periodic
Age:80 YR Gender:Female I/FU:I

Company Report #001-0945-990265

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 300 MG (100		Dysphagia Pneumonia Aspiration	Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

MG, TID), PER

ORAL

Date:02/14/00ISR Number: 3459634-3Report Type:Periodic Company Report #001-0945-990267
Age:63 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (400 MG,TID), PER ORAL	Convulsion	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
			Topamax (Topiramate)	SS		

Date:02/14/00ISR Number: 3459638-0Report Type:Periodic Company Report #001-0945-990268
Age:23 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged YR	Cerebrovascular Accident	Consumer	Neurontin (Gabapentin)	PS		

Date:02/14/00ISR Number: 3459641-0Report Type:Periodic Company Report #001-0945-990269
Age:3 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG (300	Stevens-Johnson Syndrome	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

MG, BID) PER

ORAL

Date:02/14/00ISR Number: 3459646-XReport Type:Periodic Company Report #001-0945-990271
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (DAILY)		Mental Disorder Oedema Peripheral	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		

Date:02/14/00ISR Number: 3459648-3Report Type:Periodic Company Report #001-0945-990284
 Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (300		Visual Disturbance	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

MG,BID), PER

ORAL

Depakote (Valproate
Semisodium) C
(Ibuprofen) C

Date:02/14/00ISR Number: 3459650-1Report Type:Periodic Company Report #001-0945-990319
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Bronchitis Gingival Bleeding Influenza Like Illness	Consumer Health Professional	Neurontin (Gabapentin) Sporanox	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

37.5 MG
 (DAILY), PER
 ORAL; 75 MG
 (DAILY), PER
 ORAL; 150 MG

Hydrochloride) SS ORAL

Remeron
 (Mirtazapine) C
 Xanax (Alprazolam) C

Date:02/14/00ISR Number: 3459740-3Report Type:Periodic Company Report #001-0945-990945
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
Required		Facial Pain		(Gabapentin)	PS		
4100 MG							
Intervention to		Headache		Effexor (Venlafaxine			
Prevent Permanent		Neck Pain		Hydrochloride)	C		
Impairment/Damage		Overdose		Motrin (Ibuprofen)	C		
				Tylenol			
				(Paracetamol)	C		

Date:02/14/00ISR Number: 3459742-7Report Type:Periodic Company Report #001-0945-990967
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Drug Toxicity	Health	Neurontin			
Initial or Prolonged		Mental Impairment	Professional	(Gabapentin)	PS		ORAL
1800 MG (600							
MG, TID), PER		Respiratory Distress	Company				
			Representative				
ORAL; 300 MG							
(100 MG,							
TID), PER							

Date:02/14/00ISR Number: 3459745-2Report Type:Periodic Company Report #001-0945-991015
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Renal Failure Acute	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/14/00ISR Number: 3459748-8Report Type:Periodic Company Report #001-0945-991016
Age:55 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Pneumonia Pyrexia Urinary Tract Infection	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/14/00ISR Number: 3459777-4Report Type:Periodic Company Report #001-0945-991017
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Stevens-Johnson Syndrome	Health Professional	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/00ISR Number: 3459781-6Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #001-0945-991019

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Stevens-Johnson Syndrome	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/14/00ISR Number: 3459784-1Report Type:Periodic
 Age:47 YR Gender:Male I/FU:I

Company Report #001-0945-991034

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG (QHS)	Dermatitis	Health Professional	Neurontin (Gabapentin)	PS		
			Asa (Acetylsalicylic Acid)	C		
			(Citalopram)	C		
			Cardizem Cd (Diltiazem Hydrochloride)	C		
			(Metoprolol)	C		
			(Valproic Acid)	C		

Date:02/14/00ISR Number: 3459786-5Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #001-0945-991037

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1500 MG	Disorientation Drug Interaction		Neurontin Capsules 100 Mg (Gabapentin)	PS		
1750 MG	Drug Level Above Therapeutic		Depakote (Valproate Semisodium)	SS		
	Pyrexia					

Date:02/14/00ISR Number: 3459793-2Report Type:Periodic
 Age:52 YR Gender:Female I/FU:I

Company Report #001-0945-991041

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (SEE TEXT), PER ORAL; 300 MG (DAILY), PER ORAL		Back Injury Drug Interaction Syncope	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ONE (DAILY), PER ORAL 20 MG (DAILY), PER ORAL				Ziac (Hydrochlorothiazide , Bisoprolol Fumarate)	SS		ORAL
ONE (DAILY), PER ORAL				Paxil (Paroxetine Hydrochloride)	SS		ORAL
ONE (DAILY), PER ORAL				Lasix (Furosemide)	SS		ORAL
				Estratab (Estrogens Esterified) Indomethacin (Indometacin)	SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/00ISR Number: 3459797-XReport Type:Periodic
Age:18 YR Gender:Male I/FU:I

Company Report #001-0945-991042

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Intentional Misuse Suicide Attempt	Health Professional	Neurontin (Gabapentin)	PS		
				Luvox (Fluvoxamine Maleate)	SS		
				Zyprexa (Olanzapine)	C		
				Tylenol (Paracetamol)	C		

Date:02/14/00ISR Number: 3459801-9Report Type:Periodic
Age:34 YR Gender:Female I/FU:I

Company Report #001-0945-991067

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1600 MG (400 MG, QID), PER ORAL		Drug Withdrawal Syndrome Feeling Abnormal Tachycardia	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
				(Baclofen)	C		
				(Nortriptyline)	C		

Date:02/14/00ISR Number: 3459807-XReport Type:Periodic
Age:72 YR Gender:Male I/FU:I

Company Report #001-0945-991088

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2700 MG (300 MG, TID), PER ORAL		Urinary Retention	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blindness	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		
1200 MG (300 MG, QID)				Flexeril (Cyclobenzaprine Hydrochloride)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Agitation Dermatitis	Health Professional	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL
1200 MG (400 MG, TID), PER ORAL; 400 MG (DAILY), PER ORAL		Difficulty In Walking Fall Hypotension		Celexa (Citalopram Hydrobromide) Aerocef (Cefixime Trihydrate)	SS C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/00ISR Number: 3459820-2Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-991161

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Other	Neurontin (Gabapentin)	PS		

Date:02/14/00ISR Number: 3459825-1Report Type:Periodic
Age:65 YR Gender:Female I/FU:I

Company Report #001-0945-991164

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (100		Pancreatitis	Health Professional	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL

MG, BID), PER

ORAL

Norvasc (Amlodipine
Besilate) C

Date:02/14/00ISR Number: 3459829-9Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-991171

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Neurontin (Gabapentin)	PS		

900 MG TO

3600 MG

(DAILY)

Company
Representative

Date:02/14/00ISR Number: 3459832-9Report Type:Periodic
Age:51 YR Gender:Female I/FU:I

Company Report #011-0945-991173

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Hepatic Enzyme Increased	Consumer	Neurontin Capsules		
300 MG	Weight Increased		300 Mg (Gabapentin)	PS	ORAL
(DAILY), PER					
ORAL			(Morphine)	C	
			Zestril (Lisinopril)	C	
			Prilosec		
			(Omeprazole)	C	
			Synthroid		
			(Levothyroxine		
			Sodium)	C	

Date:02/14/00ISR Number: 3459838-XReport Type:Periodic Company Report #011-0945-991197
 Age:48 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Drug Interaction	Consumer	Neurontin Capsules			
Initial or Prolonged	Eating Disorder		400 Mg (Gabapentin)	PS		ORAL
1600 MG (400	Fatigue					
MG, QID), PER	Hypotension					
ORAL	Mental Impairment		Tylenol Pm			
	Sedation		(Diphenhydramine,			
			Paracetamol)	SS		
			Propulsid			
			(Cisapride)	C		
			Prevacid			
			(Lansoprazole)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Calcium) C
 Estratest
 (Methyltestosterone,
 Estrogens
 Esterified) C
 (Baclofen) C
 Celebrex (Celecoxib) C
 Multivitamins C

Date:02/16/00ISR Number: 3536823-0Report Type:Periodic Company Report #1999-003129
 Age:82 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
0.4 MG/QD/PO		Gynaecomastia	Health Professional	Flomax	PS	Boehringer Ingelheim Pharmaceuticals Inc	ORAL
				Neurontin	SS		
				Lorazepam	C		
				Vitamins	C		

Date:02/17/00ISR Number: 3459377-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000148
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage ORAL		Fracture Impaired Healing Injury	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Zoloft	C		
				Motrin	C		

Date:02/18/00ISR Number: 3459733-6Report Type:Expedited (15-DaCompany Report #033-0945-M0000012
 Age:89 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Death Cardiac Disorder Foreign Gabapentin PS ORAL
1200 MG Study (Gabapentin)
(DAILY), PER Health
ORAL Professional

Adancor (Nicorandil) C
Tildiem (Diltiazem
Hydrochloride) C
Renitec (Enalapril
Maleate) C
Kardegic
(Acetylsalicylate
Lysine) C
Digoxine Nativelle
(Digoxin) C
Lasilix (Furosemide) C

Date:02/22/00ISR Number: 3460749-4Report Type:Expedited (15-DaCompany Report #032-0945-M0000002
Age:32 YR Gender:Female I/FU:F

Outcome PT Report Source
Hospitalization - Depression Foreign
Initial or Prolonged Mental Disorder Study
Health

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
1200 MG		Gabapentin (Gabapentin)	PS		ORAL
(DAILY), PER					
ORAL		Valproate (Valproate Bismuth)	C		
		Clonazepam	C		
		Venlafaxine	C		
		Dicalium			
		Chlorazepate	C		

Date:02/22/00ISR Number: 3460889-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000141

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous Convulsion	Health Professional	Neurontin (Gabapentin)	PS		
1200 MG							
(DAILY)							
1200 MG				Tegretol (Carbamzepine)	SS		
(DAILY)							
600 MG				Lamictal (Lamotrigine)	SS		
(DAILY)							
25 MG (DAILY)				Topamax (Topiramate)	SS		
				Allegra (Fexofenadine Hydrochloride)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 300 MG Initial or Prolonged (DAILY) PER ORAL	Blister Dermatitis Bullous Dialysis Gangrene Leg Amputation Oedema Peripheral Renal Failure Skin Infection	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Actrapid (Insulin Human)	C		
			Levoxin (Levothyroxine Sodium)	C		
			Rhinocort (Budesone)	C		
			(Acetylcysteine)	C		
			Nitromex (Glyceryl Trinitrate)	C		
			Glucophage (Metformin Hydrochloride)	C		
			Zantac (Ranitidine Hydrochloride)	C		
			Renitec (Enalapril Maleate)	C		
			Trombyl (Acetylsalicylic Acid)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lasix (Furosemide) C
 Distalgesic
 (Paracetamol,
 Dextropropoxyphene) C
 Lipid (Gemfibrozil) C
 Zocord (Simvastatin) C
 Behepan
 (Cyanocobalamin) C
 Seloken Zoc
 (Metoprolol
 Succinate) C

Date:02/22/00ISR Number: 3461092-XReport Type:Expedited (15-DaCompany Report #WAES 00021101
 Age:66 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening PO	Biopsy Liver Abnormal	Health	Tab Vioxx Unk	PS		ORAL
Hospitalization - Initial or Prolonged	Blood Bilirubin Increased Hepatic Cirrhosis Hepatic Failure Hepatitis Cholestatic Jaundice Liver Function Test Abnormal Prothrombin Time Prolonged	Professional	Neurontin Celebrex Ambien Cozaar Norvasc Reglan	SS SS C C C C		

Date:02/22/00ISR Number: 3462547-4Report Type:Periodic Company Report #S99-USA-02204-01
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Bradycardia Hypotension Sedation	Health Professional Company Representative	Celexa Neurontin	PS SS		

Date:02/23/00ISR Number: 3462050-1Report Type:Periodic Company Report #9941953
 Age:22 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination	Health	Zoloft Tablets	PS		
		Hostility	Professional	Effexor	SS		
		Nervousness		Unspecified			
		Neurosis		Medication	SS		
		Pain		Neurontin	SS		
		Suicidal Ideation					

Date:02/23/00ISR Number: 3462475-4Report Type:Periodic Company Report #9937754
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorgasmia	Health	Zoloft Tablets	PS		ORAL
ORAL		Libido Increased	Professional	Neurontin	SS		ORAL

1500.00 MG

TOTAL: TID :

ORAL

Levoxyl	C
Zantac	C

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Freedom Of Information (FOI) Report

Date:02/23/00ISR Number: 3468455-7Report Type:Periodic
 Age:33 YR Gender:Female I/FU:I

Company Report #9948357

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Consumer	Zoloft Tablets	PS		ORAL
50.00 MG		Insomnia					
TOTAL:DAILY:0							
RAL							
				Neurontin	SS		
				Amitriptyline	SS		

Date:02/23/00ISR Number: 3469143-3Report Type:Periodic
 Age: Gender:Female I/FU:F

Company Report #9828199

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea	Health	Zoloft Tablets	PS		ORAL
ORAL		Drug Interaction	Professional	Neurontin	SS		
				Ambiem	C		
				Hydrocodone/ Acetaminophen	C		
				Unknown Estrogen	C		

Date:02/24/00ISR Number: 3462590-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000139
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Coombs Direct Test Positive	Health Professional	Neurontin (Gabapentin)	PS		
600 MG		Haematocrit Decreased Haemoglobin Decreased		Seroquel (Quetiapine)	SS		
		Haemolytic Anaemia Psychotic Disorder					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (DAILY)	Atrioventricular Block Cardiac Disorder Dyspnoea	Foreign Study Health Professional	Gabapentin (Gabapentin) Vasten (Pravastatin Sodium) Aspegic (Acetylsalicylate Lysine) Mopral (Omeprazole) Atarax (Hydroxyzine Hydrochloride) Temesta (Lorazepam) Lescol (Fluvastatin Sodium)	PS C C C C C C		

Outcome	PT
Hospitalization - Initial or Prolonged Other	Blood Creatinine Increased Blood Urea Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Coma Headache Lethargy	Report Source	Product	Role	Manufacturer	Route
300 MG (QHS), PER ORAL		Nasal Congestion Sneezing Toxic Epidermal Necrolysis Vomiting	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Avandia (Rosiglitazone)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Ecotrin (Acetylsalicylic Acid)	C		
				Trental (Pentoxifylline)	C		
				Diovan (Valsartan)	C		
				Trandate (Labetalol Hydrochloride)	C		
				Insulin N	C		
				Normodyne (Labetalol Hydrochloride)	C		

Date:02/28/00ISR Number: 3464506-4Report Type:Expedited (15-DaCompany Report #032-0945-M0000005
Age:39 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Agitation Delirium Hallucination Mental Impairment	Report Source Foreign Health Professional Company Representative	Product Gabapentin (Gabapentin)	Role PS	Manufacturer	Route ORAL
1200 MG (DAILY), PER ORAL				(Valporate Sodium)	SS		
INTRAVENOUS (DAILY), INTRAVENOUS	1600 MG			Phenytoin (Pheytoin)	C		

(Vigabatrin) C
(Carbamazepine) C

Date:02/28/00ISR Number: 3464509-XReport Type:Expedited (15-DaCompany Report #055-0945-M0000003
Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Lymphopenia	Foreign	Gabapentin	PS		ORAL
800 MG			Health				
(DAILY), PER			Professional				
ORAL				Captopril	C		
				Nifedipine	C		
				Hydrochlorothiazide	C		
				Glibenclamide	C		

Date:02/28/00ISR Number: 3464512-XReport Type:Expedited (15-DaCompany Report #033-0945-990068
Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Foreign	Neurontin			
6 TABLET(S)		Rash Pustular	Health	(Gabapentin)	PS		ORAL
(DAILY), PER			Professional				
ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Daonil
 (Glibenclamide) C
 Glucophage
 (Metformin
 Hydrochloride) C
 Creon (Pancreatin) C
 Atarax (Hydroxyzine
 Hydrochloride) C

Date:02/28/00ISR Number: 3464660-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000167
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1500 MG (DAILY), PER ORAL		Astrocytoma Feeling Drunk	Consumer	Neurontin	PS		ORAL

Tegretol
 (Carbamazepine) C

Date:02/29/00ISR Number: 3465569-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000175
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG 400 MG TID PER ORAL		Deafness Neurosensory	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL

(Trazodone) C
 Tricor (Fenofibrate) C
 Vicodin
 (Paracetamol,
 Hydrocodone
 Bitartrate) C
 Oxycontin (Oxycodone
 Hydrochloride) C

Date:03/02/00ISR Number: 3468824-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000168
Age:79 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG BID, 600 MG QHS, PER ORAL	Abnormal Dreams Confusional State Coordination Abnormal Dehydration Disturbance In Attention Dysphemia International Normalised Ratio Increased Parkinsonian Gait Tremor	Health Professional	Neurontin Capsules 300 Mg (Gapapentin) Coumadin (Warfarin Sodium)	PS C		ORAL

Date:03/02/00ISR Number: 3468827-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000174
Age:79 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Bronchitis Fall

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hallucination Vision Blurred	Report Source	Product	Role	Manufacturer	Route
200 MG			Consumer	Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER				Unspecified Anti-Hypertensive	C		

Date:03/02/00ISR Number: 3468836-1Report Type:Expedited (15-DaCompany Report #039-0945-M0000001
Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Purpura Rash Erythematous	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
15 DROPS			Professional	Laroxyl (Amitriptyline Hydrochloride)	SS		ORAL
(DAILY) , PER							

Date:03/02/00ISR Number: 3469856-3Report Type:Expedited (15-DaCompany Report #USA/00/00481/MES
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aspiration Convulsion	Health Professional	Mesantoin (Mephenytoin)	PS		ORAL
100 MG, THREE							
TIMES A DAY,							
ORAL							

Dilantin (Phenytoin Sodium) SS
 Neurontin (Gabapentin) SS

Date:03/06/00ISR Number: 3471053-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000172
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG (300	Asthenia Burning Sensation	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
MG, TID), PER		Condition Aggravated					
ORAL		Micturition Urgency					
		Oliguria					
		Restless Legs Syndrome					
		Spinal Column Stenosis					

Date:03/06/00ISR Number: 3552811-2Report Type:Periodic Company Report #1999UW04406
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG HS PO;		Movement Disorder	Consumer	Seroquel	PS	Astrazeneca Uk Ltd	ORAL
50 MG QAM PO;		Muscle Rigidity					
400 MG HS PO		Peripheral Sensory					
1500 MG DAILY		Neuropathy		Neurontin	SS		
		Speech Disorder		Ambien	C		
				Lamictal	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Effexor C
Zoloft C

Date:03/09/00ISR Number: 3472211-3Report Type:Direct
Age:56 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Sexual Dysfunction		Gabapentin	PS		
				Oxycodone Hcl	C		
				Docusate Na	C		
				Fosinopril Na	C		

Date:03/09/00ISR Number: 3472317-9Report Type:Direct
Age:54 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Sexual Dysfunction		Gabapentin	PS		
				Warfarin (Coumadin)			
				Na	C		
				Isosorbide Dinitrate	C		
				Captopril	C		
				Metoprolol Tartrate	C		
				Pravastatin	C		
				Ranitidine Hcl	C		
				Bepiridil Hcl	C		
				Nitroglycerin	C		
				Gabapentin	C		

Date:03/09/00ISR Number: 3472319-2Report Type:Direct
Age:49 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Sexual Dysfunction		Gabapentin	PS		
				Fentanyl	C		
				Beclomethasone	C		
				Morphine S04	C		

Date:03/09/00ISR Number: 3472320-9Report Type:Direct
Age:32 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Gabapentin	PS		
Other		Sexual Dysfunction		Venlafaxine Hcl	C		
				Lansoprazole Sa	C		
				Tolmetin Sodium	C		
				Fentanyl	C		

Date:03/09/00ISR Number: 3472445-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000188
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin			
Life-Threatening		Respiratory Arrest	Health Professional Company Representative	(Gabapentin)	PS		
				(Methadone)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/09/00ISR Number: 3472446-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000196
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged TWO (TID), PER ORAL		Ovarian Cyst Vomiting	Consumer	Neurontin (Gabapentin)	PS		ORAL
				(Baclofen)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Ms Contin (Morphine Sulfate)	C		

Date:03/10/00ISR Number: 3473306-0Report Type:Direct Company Report #
 Age:86 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100MG PO TID		Movement Disorder	Health Professional	Neurontin	PS		ORAL
				Asa	C		
				Calcium	C		
				Cardizem	C		
				Premarin	C		
				Fe-So4	C		
				Lasix	C		
				Prevacid	C		
				Synthroid	C		
				Mag Ox	C		
				Peri-Colace	C		
				K-Dur	C		
				Prednisone	C		
				Vit E	C		

Date:03/10/00ISR Number: 3473911-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000200
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization - Initial or Prolonged 1200 MG (300 MG,DAILY)	Drug Interaction Respiratory Arrest	Health Professional	Neutrontin (Gabapentin)	PS
			Ms Contin (Morphine Sulfate)	SS
			Synthroid (Levothy-Roxine Sodium)	C
			Norvasc (Amlodipine Besilate)	C
			Enteric Coated Aspirin (Acetylsalicylic Acid)	C
			Plavix (Clopidogrel)	C
			Diovan (Valsartan)	C
			Duragesic (Fentanyl)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/13/00ISR Number: 3475095-2Report Type:Expedited (15-DaCompany Report #002-0945-990030

Age:10 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Disorder	Foreign	Neurontin			
Other		Cardio-Respiratory Arrest	Health	(Gabapentin)	PS		ORAL
1400 MG		Drug Interaction	Professional				
(DAILY), PER		Liver Disorder					
ORAL		Viral Infection		Morphine (Morphine)	SS		
INTRAVENOUS	INTRAVENOUS			Carbamazepine			
				(Carbamazepine)	SS		ORAL
PER ORAL				Amitriptyline			
				(Amitriptyline)	SS		ORAL
PER ORAL							

Date:03/14/00ISR Number: 3475239-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000022

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Neurontin			
Other		Suicidal Ideation		(Gabapentin)	PS		

Date:03/14/00ISR Number: 3475240-9Report Type:Expedited (15-DaCompany Report #001-0945-991191

Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Health	Neurontin Capsules			
Other			Professional	100 Mg (Gabapentin)	PS		ORAL
400 MG (200MG							
AT 7AM, 200MG							
AT 12 NOON),							
PER ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Blood Pressure Decreased	Consumer	Neurontin			
		Confusional State		(Gabapentin)	PS		
		Disturbance In Attention		Thorazine			
		Dizziness		(Chlorpromazine			
		Drug Level Above		Hydrochloride)	SS		ORAL
200 MG (AT		Therapeutic					
BEDTIME), PER		Dry Mouth					
ORAL		Dysarthria		Valium (Diazepam)	C		
		Dysphagia		Xanax (Alprazolam)	C		
		Dyspnoea		Prozac (Fluoxetine			
		Hyperhidrosis		Hydrochloride)	C		
		Hypoaesthesia		Buspar (Buspirone			
		Hypotension		Hydrochloride)	C		
		Hypotonia		Adipex (Phentermine			
		Muscle Disorder		Hydrochloride)	C		
		Muscle Rigidity		Topamax (Topiramate)	C		
		Muscular Weakness		Vioxx	C		
		Nausea		Skelatin			
		Pharyngeal Oedema		(Metaxalone)	C		
		Tremor		Ultram (Tramadol			
		Urinary Incontinence		Hydrochloride)	C		
		Vertigo		Maxalt	C		
				Risperdal			
				(Risperidone)	C		

Freedom Of Information (FOI) Report

Ceroquel C

Date:03/16/00ISR Number: 3477633-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000022
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		

Date:03/17/00ISR Number: 3478331-1Report Type:Expedited (15-DaCompany Report #001-0945-991153
 Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Amyotrophic Lateral	Health	Neurontin	PS		
3600 MG (DAILY), (1200MG TID)		Sclerosis Condition Aggravated	Professional				
		Decreased Appetite		...	C		
		Drug Withdrawal Syndrome		...	C		
		Flatulence		Aspirin			
		Influenza Like Illness		(Acetylsalicylic Acid)	C		
		Insomnia		Multivitamins			
		Malaise		(Ergocalciferol, Ascorbid Acid, Folic Acid, Thiamine Hydrochloride,	C		
		Sedation					
		Weight Decreased					

Date:03/20/00ISR Number: 3477886-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000232
 Age:1 DY Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Complications Of Maternal Exposure To Therapeutic	Health Professional	Neurontin (Gabapentin)	PS		

SEE TEXT,

PLACENTAL
 (IN UTERO
 EXPOSURE)
 Drugs
 Stillbirth
 Umbilical Cord Around
 Neck

Levoxyl
 (Levothyroxine
 Sodium) C
 Antibiotics Nos C
 Haldol (Haloperidol) C
 Celexa (Citalopram
 Hydrobromide) C
 Benadryl
 (Diphenhydramine
 Hydrochloride) C

Date:03/20/00ISR Number: 3477887-2Report Type:Expedited (15-DaCompany Report #034-0945-M0000027
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (DAILY), PER ORAL		Microcytic Anaemia Pulmonary Embolism	Foreign Study Health Professional	Gabapentin Or Placebo Placebo Levodopa	PS SS C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

L-Dopa Retard C
 Donperidona C
 Bromazepam C
 Amantadine C
 Entacapone C
 Pramipexole C

Date:03/20/00ISR Number: 3477889-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000229
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, DAILY) PER ORAL		Inflammation Oedema Peripheral	Consumer	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Xanax (Alprazolam) Zoloft (Sertraline Hydrochloride) Trazodone)	C C C		

Date:03/20/00ISR Number: 3477970-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000224
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion Drug Interaction	Health Professional Company Representative	Neurontin (Gabapentin) Ultram (Tramadol Hydrochloride)	PS SS		

Date:03/21/00ISR Number: 3478591-7Report Type:Expedited (15-DaCompany Report #225690
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Confusional State	Foreign	Rivotril			

Initial or Prolonged ORAL	Hallucination	Other	(Clonazepam)	PS	ORAL
ORAL			Tardyferon (Ferrous Sulfate)	SS	ORAL
100 MG 1 PER			Tranxene (Clorazepate Dipotassium)	SS	ORAL
1 DAY ORAL					
600 MG 1 PER			Leponex (Clozapine)	SS	ORAL
1 DAY ORAL					
ORAL			Mopral (Omeprazole)	SS	ORAL
200 MG 1 PER			Neurontin (Gabapentin)	SS	ORAL
1 DAY ORAL					

Date:03/22/00ISR Number: 3478695-9Report Type:Expedited (15-DaCompany Report #033-0945-M0000021
Age:87 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Alanine Aminotransferase
Initial or Prolonged	Increased
	Diarrhoea
	Hepatitis Cholestatic
	Pyrexia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100 MG		Foreign Health Professional	Gabapentin (Gabapentin)	PS		
INCREASE TO 600 MG DAILY			Renitec (Enalapril Maleate)	C		
			Lasilix (Furosemide)	C		
			Nitriderm(Glyceryl Trinitrate)	C		
			Loxen (Nicardipine)	C		
			Digoxine (Digoxin)	C		
			Asasantine (Acetylsalicylic Acid, Dipyridamole)	C		

Date:03/22/00ISR Number: 3478699-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000168
Age:79 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG BID,	Abnormal Dreams Confusional State	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
600 MG QHS,		Coordination Abnormal					
PER ORAL		Dehydration					
		Disturbance In Attention Dysphemia International Normalised Ratio Increased Parkinsonian Gait Tremor		Coumadin (Warfarin Sodium)	C		

Date:03/22/00ISR Number: 3478702-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000250
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (100 MG, QID) PER ORAL		Ascites Pyrexia Sepsis Urine Analysis Abnormal	Consumer	Neurontin Capsules 100 Mg (Gabapentin)	PS		ORAL
				Lasix (Furosemide) Aldactone (Spironolactone) Lactulose Noroxin (Norfloxacin Magnesium Tylenol Pm (Diphenhydramine, Paracetamol)	C C C C C C		

Date:03/22/00ISR Number: 3479004-1Report Type:Direct
Age:44 YR Gender:Female I/FU:I

Company Report #

Outcome	PT
Death Life-Threatening	Culture Urine Positive Depressed Level Of Consciousness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Discomfort Escherichia Infection Escherichia Sepsis				
NEURONTIN 300		Fluid Overload Leg Amputation	Neurontin Capsules 300 Mg	PS		ORAL
MG PO TID		Multi-Organ Failure				
25-50 MG		Nausea Oedema	Sandimmune Capsules 25,50 Mg	SS		
QD-BID		Oesophageal Disorder				
		Polyuria Renal Failure Acute Sepsis Vomiting	Morphine Vancomycin Mag Ox Imuran Bumex Prednisone Coumadin	C C C C C C C		

Date:03/23/00ISR Number: 3479097-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000260
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE IMAGE		Arrhythmia Chest Pain Pain In Extremity	Consumer	Neurontin (Gabapentin) Synthroid (Levothyroxine Sodium) Estrace (Estradiol) Sinequan (Doxepin Hydrochloride)	PS C C C		ORAL

Date:03/23/00ISR Number: 3479098-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000264
Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Confusional State	Consumer	Neurontin Tablets			

Initial or Prolonged Disorientation Health 600 Mg (Gabapentin) PS ORAL
1800 MG (600
Renal Cell Carcinoma Professional
MG, TID), PER
Stage Unspecified
ORAL

Zestril (Lisinopril) C
Levoxyl
(Levothyroxine
Sodium) C
Celebrex (Celecoxib) C
Baby Aspirin
(Acetylsalicylic
Acid) C

Date:03/23/00ISR Number: 3479103-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000243
Age:86 YR Gender:Female I/FU:I

Outcome PT
Disability Aphasia
Other Cerebrovascular Accident
Confusional State
Decreased Activity
Depressed Level Of
Consciousness
Fall

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Outcome	PT
Hospitalization -	Abnormal Behaviour
Initial or Prolonged	Aphasia
Other	Balance Disorder
	Clonic Convulsion
	Convulsion
	Coordination Abnormal
	Decreased Appetite
	Drooling
	Drug Level Above
	Therapeutic
	Drug Level Below
	Therapeutic
	Ear Disorder
	Eye Rolling
	Gingival Bleeding
	Gingival Pain
	Grand Mal Convulsion
	Insomnia
	Intentional Self-Injury
	Nervous System Disorder
	Neurosis
	Nystagmus

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Petit Mal Epilepsy Psychotic Disorder Pyrexia	Report Source	Product	Role	Manufacturer	Route
200 MG(100 MG, BID), PER ORAL		Status Epilepticus Tremor Vomiting	Consumer	Dilantin Kapseals 100 Mg (Phenytoin Sodium)	PS		ORAL
	INTRA VENOUS (ONE DOSE), INTRA VENOUS			Dilantin (Phenytoin Sodium)	SS		
2 MILLILITERS (BID), PER ORAL				Dilantin Suspension 125 Mg/5 ML (Phenytoin Sodium)	SS		ORAL
200 MG (100 MG, BID), PER ORAL				Dilantin Infatabs 50 Mg (Phenytoin Sodium)	SS		ORAL
300 MG (DAILY), PER ORAL				Neurontin (Gabapentin)	SS		ORAL
				(Phenobarbital)	SS		
				Lamictal (Lamotrigine)	SS		
				Tegretol (Carbamazepine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/00ISR Number: 3482018-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000257
 Age:54 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG (600 MG, TID), PER ORAL	Back Pain Blood Pressure Fluctuation Blood Pressure Systolic Increased Condition Aggravated Cyst Drug Withdrawal Syndrome Spinal Disorder Tachycardia	Health Professional Company Representative	Neurontin (Gabapentin) Percocet Unspecified Anti Depressant	PS C C		ORAL

Date:03/30/00ISR Number: 3482026-8Report Type:Expedited (15-DaCompany Report #034-0945-M0000027
 Age:72 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG Required (DAILY), PER Intervention to ORAL Prevent Permanent Impairment/Damage	Dyspnoea Epistaxis Fatigue Influenza Iron Deficiency Anaemia Microcytic Anaemia Parkinson'S Disease Pulmonary Embolism	Foreign Study Health Professional	Gabapentin Or Placebo (Levodopa) L-Dopa Retard Donperidona (Bromazepam) (Amantadine) Entacapon E Pramipexole	PS C C C C C C		ORAL

Date:03/31/00ISR Number: 3482626-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000288
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abortion Spontaneous	Health Professional	Neurontin (Gabapentin)	PS		
1200 MG							
(DAILY)	7 WK			Prenatal Vitamins	C		

Date:03/31/00ISR Number: 3482703-9Report Type:Periodic Company Report #9940167
 Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Confusional State	Consumer	Cardura	PS		ORAL
4.00 MG TOTAL		Diarrhoea					
DAILY ORAL		Headache		Neurontin	SS		ORAL
100.00 MG		Nervousness					
TOTAL DAILY		Oedema Peripheral					
ORAL		Sedation		Stool Softner Risperidone	C C		

Date:04/03/00ISR Number: 3483336-0Report Type:Expedited (15-DaCompany Report #032-0945-M0000010
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source
Death	Completed Suicide Intentional Misuse	Foreign Study

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Health Professional	Product	Role	Manufacturer	Route
PER ORAL			Gabapentin (Gabapentin)	PS		ORAL
			(Carbamazepine)	SS		
			(Loprazolam)	SS		

Date:04/06/00ISR Number: 3484701-8Report Type:Direct
Age:31 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2 DAY		Asthenia	Health Professional	Neurontin	PS		
Initial or Prolonged		Dizziness Nausea					

Date:04/07/00ISR Number: 3486543-6Report Type:Expedited (15-DaCompany Report #034-0945-M0000028
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 600 MG		Drug Interaction	Foreign	Gabapentin	PS		ORAL
(DAILY), PER		Hepatic Function Abnormal	Study				
ORAL		Sedation	Health Professional	(Morphine Sulfate)	C		
				(Prednisone)	C		
				(Metamizole)	C		
				(Clebopride Hydrogen Maleate)	C		
				(Dimeticone)	C		
				(Ranitidine)	C		
				(Megestrol Acetate)	C		
				(Alprazolam)	C		
				(Bisacodyl)	C		
				(Furosemide)	C		
				(Lormetazepam)	C		

Date:04/11/00ISR Number: 3487404-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000314
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Lung Infiltration Pneumonia Respiratory Failure	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:04/12/00ISR Number: 3487347-0Report Type:Direct Company Report #
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG PO QNS		Urinary Incontinence		Neurontin 300mg	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/12/00ISR Number: 3487365-2Report Type:Expedited (15-DaCompany Report #033-0945-990082

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Genital Disorder Female Vaginal Laceration	Foreign Study Health Professional	Gabapentin (Gabapentin)	PS		

Date:04/12/00ISR Number: 3487585-7Report Type:Expedited (15-DaCompany Report #001-0945-990641

Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Atrophy Nervous System Disorder	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL

1200 MG (300

MG, 2QAM,

1QPMHS), PER

ORAL

Celebrex (Celecoxib)	C
Xanax (Alprazolam)	C
Premarin (Estrogens Conjugated)	C
Parafon Forte (Chlorzoxazone, Paracetamol)	C
Armour Thyroid (Thyroid)	C

Date:04/13/00ISR Number: 3487954-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000329

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blister Dermatitis Joint Swelling Neuralgia	Consumer	Neurontin (Gabapentin)	PS		

Date:04/17/00ISR Number: 3488983-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000328
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1200 MG (600 Initial or Prolonged MG QAM, 600 MG QPM) PER ORAL		Amnesia Convulsion	Consumer	Neurontin	PS		ORAL

Date:04/18/00ISR Number: 3490109-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000347
 Age:10 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL; 900 MG (300 MG, TID)		Convulsion Status Epilepticus	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
				Methylphenidate (Methylphenidate) Gabitril (Tiagabine)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/19/00ISR Number: 3490454-XReport Type:Expedited (15-DaCompany Report #001-0945-990404

Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blindness	Health	Neurontin Capsules			
Disability		Cerebrovascular Accident	Professional	100 Mg (Gabapentin)	PS		ORAL

400 MG (200

MG, BID), PER

ORAL

Lotensin (Benazepril Hydrochloride) C
 (Digoxin) C
 Nph Insulin (Insulin Injection, Isophane) C
 (Acetylsalicylic Acid) C
 Vitamin B12 (Cyanocobalamin) C

Date:04/19/00ISR Number: 3490455-1Report Type:Expedited (15-DaCompany Report #032-0945-M0000011

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dyskinesia	Foreign	Gabapentin			
SEE IMAGE		Middle Insomnia	Health	(Gabapentin)	PS		
			Professional	(Sertraline)	C		

Date:04/20/00ISR Number: 3490311-9Report Type:Direct

Company Report #U-026700

Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation Mucosal Medication Error		Neurontin Capsules			
		Skin Lesion		Generic Garbapentin	PS	Parke Davis Div Warner Lambert Co	ORAL

Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Foreign	Claritin			
Required		Drug Interaction	Health	(Loratadine) Tablets	PS		ORAL
10MG ONCE							
Intervention to			Professional				
ORAL							
Prevent Permanent				Epival Tablets	SS		ORAL
1500 MG DAILY							
Impairment/Damage							
ORAL							
				Lamotrigine Tablets	SS		ORAL
150 MG BID							
ORAL							
				Gabapentin Capsules	SS		ORAL
2000 MG DAILY							
ORAL							

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Amputation	Consumer	Glucotrol Tablets	PS		
				Nifedipine	SS		
				Glucophage	SS		
				Neurontin	SS		
				Imdur	C		
				Lasix	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Plavix	C
Prevacid	C
Hydralazine	C
Multi Vitamin	C
Iron	C
Lopid	C

Date:04/24/00ISR Number: 3492908-9Report Type:Periodic Company Report #A007326
 Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chromatopsia	Consumer	Viagra Tablets	PS		ORAL
100.00 MG		Drug Ineffective					
TOTAL PRN							
ORAL				Neurontin	SS		ORAL
ORAL							

Date:04/26/00ISR Number: 3493066-7Report Type:Expedited (15-DaCompany Report #001-0073-M0000173
 Age:22 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Chills	Consumer	Dilantin Kapseals			
Initial or Prolonged		Colonic Polyp		100 Mg (Phenytoin			
Required		Constipation		Sodium)	PS		ORAL
200 MG QAM,		Dizziness					
Intervention to		Electroencephalogram					
100 MG QPM,		Abnormal					
Prevent Permanent		Gastrooesophageal Reflux					
PER ORAL; 200		Disease		Neurontin			
Impairment/Damage		Hiatus Hernia		(Gabapentin)	SS		ORAL
MG QAM, 200		Nausea					
MG QPM, PER							
SEE TEXT, PER							
ORAL							

Nervousness
Paranoia

Centrum (Vitamins
Nos, Minerals Nos) C

Date:04/26/00ISR Number: 3493518-XReport Type:Expedited (15-DaCompany Report #000418-SK917

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Celebrex	PS		ORAL
100.000 MG QD		Drug Interaction	Professional				
PO				Neurontin	SS		ORAL
400.000 MG							
TID PO							
500.000 MG QD				Tegretol	SS		ORAL
PO							

Date:04/26/00ISR Number: 3493555-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000022

Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Agitation	Health	Neurontin			
		Anger	Professional	(Gabapentin)	PS		ORAL
1800 MG (600		Convulsion					
MG, TID), PER		Suicidal Ideation					
ORAL				Propulsid			
				(Cisapride)	C		
				Nasalcrom			
				(Cromoglicate			
				Sodium)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Celebrex (Celecoxib) C
 Copaxone (Glatiramer
 Acetate) C
 Alphagan
 (Brimonidine
 Tartrate) C
 Maxair (Pirbuterol
 Acetate) C

Date:04/26/00ISR Number: 3493556-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000375
 Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (400 MG, TID), PER ORAL (SEVERAL YEARS)		Bronchitis Eye Disorder Pruritus Scab	Consumer	Neurontin Capsules 400 Mg (Gabapentin)	PS		ORAL

Insulin (Insulin) C
 Glucophage
 (Metformin
 Hydrochloride) C
 Actos (Pioglitazone) C

Date:04/26/00ISR Number: 3493557-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000022
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, TID), PER ORAL		Agitation Anger Condition Aggravated Convulsion Suicidal Ideation	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Propulsid (Cisapride)	C		

Nasalcrom
 (Cromoglicate
 Sodium) C
 Celebrex (Celecoxib) C
 Copaxone (Glatirame
 Acetate) C
 Alphagan
 (Brimonidine
 Tartrate) C
 Maxair (Pirbuterol
 Acetate) C

Date:04/26/00ISR Number: 3559476-4Report Type:Periodic Company Report #HQ1012110FEB2000
 Age:49 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT Drug Interaction	Report Source Consumer	Product Alesse	Role PS	Manufacturer Wyeth Ayerst Laboratories Inc	Route ORAL
1 TABLET 1 X		Neuropathy Peripheral					
PER 1 DAY,							
ORAL				Celecoxib	SS		
				Neurontin	SS		
				Pamelor	SS		
				Synthroid	SS		

Freedom Of Information (FOI) Report

Date:04/27/00ISR Number: 3493649-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000301
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 MG, TID), PER ORAL; 2400 MG (600 MG BID, 1200 MG QHS),		Abnormal Dreams Amblyopia Amnesia Back Pain Confusional State Coordination Abnormal Cyst Delusional Disorder, Persecutory Type Depression Dizziness Drug Withdrawal Syndrome Dry Mouth Dry Throat Dysarthria Euphoric Mood Fatigue Feeling Abnormal Headache Hostility Meningeal Disorder Muscle Twitching Oedema Thinking Abnormal Tremor Vomiting Weight Decreased	Consumer	Neurontin Tablets 600 Mg (Gabapentin)	PS		ORAL
				Pain Medication Unspecified Celexa	C C		

Date:04/27/00ISR Number: 3493651-2Report Type:Expedited (15-DaCompany Report #033-0945-M0000036
 Age:11.5 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Effect Decreased	Foreign	Neurontin			

1200 MG Mydriasis Study (Gabapentin) PS ORAL
 (DAILY), PER Health
 ORAL; 2000 MG Professional
 (DAILY), PER
 ORAL

Date:04/28/00ISR Number: 3494391-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000387
 Age:1 DY Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Neurontin			
Other		Complications Of Maternal Exposure To Therapeutic	Professional	(Gabapentin)	PS		
TRANSPLACENTAL	SEE TEXT ,	Drugs					
PLACENTAL							
		Growth Retardation		Tegretol			
		Premature Baby		(Carbamazepine)	SS		
TRANSPLACENTAL	SEE TEXT,						
PLACENTAL							
				Trazodone	C		
				Prenatal Vitamins	C		
				Folic Acid	C		

Prothrombin Time
Prolonged

Date:05/03/00ISR Number: 3496676-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000397
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE IMAGE		Abdominal Pain Blepharospasm Diarrhoea Eye Rolling Hyperhidrosis Intervertebral Disc Protrusion Mood Swings Muscle Twitching Pyrexia Tongue Disorder	Consumer	Neurontin Capsules 300 Mg (Gabapentin) Oxycontin (Oxycodone Hydrochloride)	PS C		ORAL

Date:05/03/00ISR Number: 3496680-8Report Type:Expedited (15-DaCompany Report #033-0945-M0000037
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Abdominal Pain Upper Liver Function Test Abnormal	Foreign Health Professional	Neurontin (Gabapentin) Unspecified Hormonal	PS		ORAL

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Freedom Of Information (FOI) Report

Replacement C

Date:05/03/00ISR Number: 3496913-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000403
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 4000 MG (800 MG TID, 1600 MG QHS), PER ORAL		Pulmonary Fibrosis Respiratory Distress	Health Professional	Neurontin Tablets 800 Mg (Gabapentin)	PS		ORAL
				Zoloft (Sertraline Hydrochloride)	C		
				Depakote (Valproate Semisodium)	C		

Date:05/08/00ISR Number: 3498035-9Report Type:Expedited (15-DaCompany Report #A014180
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Pain Road Traffic Accident	Consumer	Zoloft Tablets	PS		
				Neurontin	SS		
				Xanax	SS		
				Lorcet	SS		
				Acetaminophen/Propox yphene Napsylate	C		
				Hctz/Triamterene	C		
				Omeprazole	C		
				Gemfibrozil	C		
				Famotidine	C		
				Other Medication	C		

Date:05/09/00ISR Number: 3498444-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000421
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE IMAGE, Other PER ORAL		Arterial Occlusive Disease	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
		Cerebrovascular Accident					
		Dermatitis		Aspirin			
		Drug Interaction		(Acetylsalicylic Acid)	SS		ORAL
325 MG		Headache					
(DAILY), PER		Loss Of Consciousness					
ORAL		Rosacea					
				Premarin (Estrogens Conjugated)	C		
				Aldactone (Spironolactone)	C		
				Prilosec (Omeprazole)	C		
				(Allopurinol)	C		
				Wellbutrin (Amfebutamone Hydrochloride)	C		
				Diamox (Acetazolamide)	C		
				(Oxybutynin)	C		
				Synthroid (Levothyroxine			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sodium) C
 Kaon Cl 10
 (Potassium Chloride) C

Date:05/10/00ISR Number: 3497915-8Report Type:Direct
 Age:30 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Neurontin (Gabapeptin)	PS	Parke-Davis	
Other				Rezulin (Troglitazine)	SS	Parke Davis	

Date:05/11/00ISR Number: 3499450-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000434
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Other				Topamax (Topiramate)	C		
				Valium (Diazepam)	C		
				Prozac (Fluoxetine Hydrochloride)	C		
				Desyrel (Trazodone Hydrochloride)	C		
				Zanaflex (Tizanidine)	C		
				Lasix (Furosemide)	C		
				(Potassium)	C		
				(Glucosamine)	C		
				(Calcium)	C		
				Multi Vitamin (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride,	C		

2400 MG

(DAILY)

Date:05/12/00ISR Number: 3500049-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000424
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Bilirubin Increased Cholelithiasis Jaundice Liver Function Test Abnormal	Health Professional Company Representative	Neurontin (Prednisone) Antivert (Nicotinic Acid, Meclozine Hydrochloride)	PS C C	Parke Davis Pharmaceuticals Ltd	

Date:05/12/00ISR Number: 3500051-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000192
Age:8 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness Otitis Media	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1200 MG (400 MG, TID), PER ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/12/00ISR Number: 3500052-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000074

Age:74 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG (QHS), Other PER ORAL	Atrial Flutter Coma Depressed Level Of Consciousness Dermatitis Diabetes Mellitus Inadequate Control Headache Hyperglycaemia Laceration Lethargy Nasal Congestion Pyrexia Renal Failure Acute Skin Exfoliation Sneezing Toxic Epidermal Necrolysis Vasculitis Vomiting	Health Professional	Neurontin Avandia (Rosiglitazone) Synthroid (Levothyroxine Sodium) Ecotrin (Acetylsalicylic Acid) Trental (Pentoxifylline) Diovan (Valsartan) Trandate (Labetalol Hydrochloride) Insulin N (Insulin) Normodyne (Labetalol Hydrochloride)	 C C C C C C C C C C	Parke Davis Pharmaceuticals Ltd	ORAL

Date:05/12/00ISR Number: 3500082-5Report Type:Expedited (15-DaCompany Report #009-056-0089670-00 (0)

Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged PER ORAL 3600 MG, 1 IN 1D, PER ORAL	Ileus Paralytic Mesenteric Occlusion	Foreign Health Professional	Depakene Gabapentin (Gabapentin)	 SS	Abbott Laboratories Pharmaceutical Products Div	ORAL ORAL

Date:05/15/00ISR Number: 3499784-9Report Type:Direct
Age:53 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - TID	Duration Fall		Neurontin 300mg Tid	PS		ORAL
Initial or Prolonged 500 MG PO IN AM; 1000 MG PO AT NIGHT	Lethargy Loss Of Consciousness Syncope		Depakote 500 In Am & 1000mg At Night	SS		ORAL
			Oxybutynin	C		
			Celexa	C		
			Lorazepam	C		
			Clozapine	C		
			...	C		

Date:05/17/00ISR Number: 3501974-3Report Type:Periodic
Age: Gender:Male I/FU:F

Company Report #S99-USA-02204-01

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Bradycardia Hypotension Sedation	Health Professional Company Representative	Celexa Neurontin	PS SS	Forest Laboratories Inc	

Other 300 MG HS Dermatitis Health Neurontin PS

Pruritus Professional

Date:05/22/00ISR Number: 3504442-8Report Type:Periodic Company Report #99USA11129
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Photosensitivity Reaction Systemic Lupus Erythematosus	Consumer	Tegretol-Xr	PS	Novartis Pharmaceuticals Corp	ORAL
	400 MG, BID, ORAL						
	600 MG, DAILY,ORAL	White Blood Cell Count Decreased		Neurontin Tablet 300 Mg (Gabapentin)	SS		ORAL
				Imitrex	C		
				Inderal	C		
				Prednisone	C		

Date:05/24/00ISR Number: 3504448-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000347
Age:10 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	600 MG (300	Convulsion Status Epilepticus	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

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Freedom Of Information (FOI) Report

MG, BID), PER

ORAL

SEE IMAGE

Gabitril (Tiagabine)	SS	ORAL
Methylphenidate (Methylphenidate)	C	
Risperdal (Risperidone)	C	

Date:05/24/00ISR Number: 3504450-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000260
Age:48 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG (QHS), PER	Arrhythmia Bipolar Disorder Chest Pain Condition Aggravated	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
ORAL	Faecaloma Memory Impairment Pain In Extremity Sedation Weight Increased		Synthroid (Levothyroxine Sodium) Estrace (Estradiol) Sinequan (Doxepin Hydrochloride) Depakote (Valproate Semisodium) Zestril (Lisinopril)	C C C C		

Date:05/24/00ISR Number: 3504470-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000446
Age:36 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 600 MG (300 MG, BID), PER	Colour Blindness Ear Pain Eye Movement Disorder	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

ORAL

Eye Pain

Headache

Hyperaesthesia

Hyperthyroidism

Muscle Disorder

Photosensitivity Reaction

Visual Acuity Reduced

Visual Disturbance

Mestinon

(Pyridostigmine

Bromide)

C

Paxil (Paroxetine

Hydrochloride)

C

Tapazole

(Thiamazole)

C

(Trazodone)

C

(Prednisone)

C

Date:05/25/00ISR Number: 3504988-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000488

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Pulmonary Oedema	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Unspecified Medications	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/26/00ISR Number: 3565099-3Report Type:Periodic
Age:48 YR Gender:Female I/FU:I

Company Report #JRFUSA2000002822

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain Chills Depersonalisation	Health Professional	Propulsid	PS	Janssen Research Fdn Div Johnson And Johnson	ORAL
20 MG, 3 IN 1		Headache					
DAY (S), ORAL		Myalgia Nausea		Neurontin (Gabapentin)	SS		ORAL
ORAL		Visual Disturbance Vomiting		Lasix Trilisate Hydrocodone Imodium Cystospaz Trimpex Miacalcin Vasotec Hytrin Evista Zantac	C C C C C C C C C C C		

Date:05/31/00ISR Number: 3506570-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000496
Age:1 DY Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiac Aneurysm Complications Of Maternal	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Congenital Anomaly		Exposure To Therapeutic Drugs Congenital Anomaly		Unspecified Anti-Nausea Medication Prenatal Vitamins	C C C		
PLACENTAL							

Date:05/31/00ISR Number: 3506574-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000493
Age:1 DY Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
SEE TEXT,		Drugs					
PLACENTAL		Congenital Anomaly Microcephaly		Prenatal Vitamins	C		

Date:05/31/00 ISR Number: 3506578-4 Report Type:Expedited (15-DaCompany Report #001-0945-M0000504
Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Bone Disorder Bone Infection	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1800 MG (300		Infection					
MG , EVERY 4		Jaw Disorder					
HOURS), PER		Oral Soft Tissue Disorder					
ORAL				Hytrin	C		
				Catapres	C		
				(Baclofen)	C		
				Prevacid	C		
				Effexor	C		
				Remeron	C		
				Septra	C		

FDA - Adverse Event Reporting System (AERS)

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(Nifedipine) C
 Ms Contin C
 Dalmane C
 Percocet C
 (Prednisone) C
 (Nortriptyline) C
 Cipro (Ciprofloxacin Hydrochloride) C

Date:06/02/00ISR Number: 3507300-8Report Type:Direct
 Age:70 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Required 1/2 OF 8MG Intervention to TAB QD PO Prevent Permanent Impairment/Damage OPHTHALMIC		Hypoglycaemia		Avandia 8 Mg Tab -1/2 Tab Qd	PS		ORAL
		1T CAP PO QD		Neurontin 300mg Cap Qd	SS		
				...	C		

Date:06/02/00ISR Number: 3507547-0Report Type:Expedited (15-DaCompany Report #A014180
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to UNKNOWN Prevent Permanent UNKNOWN Impairment/Damage UNKNOWN UNKNOWN		Decreased Activity Pain UNKNOWN Road Traffic Accident UNKNOWN UNKNOWN	Consumer Health Professional	Zoloft Neurontin Xanax Lorcet Acetaminophen/Propox yphene Napsylate Hctz/Triamterene Omeprazole Gemfibrozil	PS SS SS SS C C C C	Pfizer Pharmaceuticals Inc	

Famotidine C
Other Medication C

Date:06/05/00ISR Number: 3508848-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000487
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ecchymosis Retinal Haemorrhage	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:06/06/00ISR Number: 3508774-9Report Type:Direct Company Report #
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening SEE IMAGE		Amnesia		Neurotin	PS		
SEE IMAGE		Cognitive Disorder		Klonopin	SS		
SEE IMAGE		Road Traffic Accident		Lamictal	SS		
SEE IMAGE		Speech Disorder		Lithium	SS		
		Visual Acuity Reduced					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/06/00ISR Number: 3509324-3Report Type:Expedited (15-DaCompany Report #033-0945-M0000022

Age:74 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 1200 MG Initial or Prolonged (DAILY), PER ORAL	Aphasia Electroencephalogram Abnormal Epilepsy Fall Haemorrhagic Stroke Head Injury Hemiplegia	Foreign Study Health Professional	Neurontin Gardenal (Phenobarbital) Lioresal (Baclofen) Tildiem (Diltiazem Hydrochloride) Nitriderm (Glyceryl Trinitrate) Ecazide (Hydrochlorothiazide , Captopril)	 PS C C C C C C	Parke Davis Pharmaceuticals Ltd	ORAL

Date:06/06/00ISR Number: 3509327-9Report Type:Expedited (15-DaCompany Report #033-0945-M0000012

Age:89 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 1200 MG (DAILY), PER ORAL	Atrial Fibrillation Cardiac Disorder Cardiac Enzymes Increased Electrocardiogram Abnormal Myocardial Infarction Pulmonary Oedema Ventricular Fibrillation	Foreign Study Health Professional	Neurontin Adancor (Nicorandil) Renitec (Enalapril Maleate) Kardegic (Acetylsalicylate Lysine) Digoxine Nativelle (Digoxin) Lasilix (Furosemide)	 PS C C C C C C	Parke Davis Pharmaceuticals Ltd	ORAL

Date:06/07/00ISR Number: 3508630-6Report Type:Direct
Age: Gender: I/FU:I

Company Report #UPC 081291

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Noroxin 400mg Neurontin (Gabapentin)	PS SS	Roberts Pharm Parke-Davis	

Date:06/07/00ISR Number: 3509637-5Report Type:Expedited (15-DaCompany Report #049-0945-M0000012
Age:56 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Required	Acetabulum Fracture Blood Iron Abnormal Colitis
Intervention to Prevent Permanent Impairment/Damage	Difficulty In Walking Escherichia Infection Fall Gastric Ulcer Haematocrit Decreased Haemoglobin Decreased Haemorrhoids

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
		Hepatic Steatosis Microcytic Anaemia Oesophageal Ulcer					
		Pain Renal Cyst Rib Fracture Spondylolisthesis Acquired Urinary Tract Infection	Foreign Health Professional	Neurontin Nonsteroidal Antiinflammatory Medication	PS SS	Parke Davis Pharmaceuticals Ltd	

Date:06/07/00ISR Number: 3509913-6Report Type:Expedited (15-DaCompany Report #033-0945-M0000045
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other PER ORAL		Optic Neuritis Retrobulbar	Foreign Health Professional	Neurontin Dafalgan	PS C	Parke Davis Pharmaceuticals Ltd	ORAL

Date:06/08/00ISR Number: 3510771-4Report Type:Expedited (15-DaCompany Report #044-0945-M0000083
Age:6 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 8 MON		Rhinorrhoea Sarcoma	Foreign Health Professional	Neurontin (Carbamazepine)	PS C	Parke Davis Pharmaceuticals Ltd	

Date:06/14/00ISR Number: 3514932-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000421
Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG, TID), PER		Cerebral Artery Occlusion Cerebrovascular Accident Dermatitis	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

ORAL (SEE	Drug Interaction				
IMAGE)/ YEARS	Headache				
325 MG	Loss Of Consciousness	Aspirin	SS		ORAL
(DAILY), PER	Malaise				
ORAL	Rosacea				
		Premarin	C		
		Aldactone	C		
		Prilosec	C		
		Allopurinol	C		
		Wellbutrin	C		
		Diamox	C		
		Oxybutynin	C		
		Synthroid	C		
		Kaon Cl 10	C		
		Aspirin	C		

Date:06/15/00ISR Number: 3514232-8Report Type:Expedited (15-DaCompany Report #033-0945-M0000037
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNK, PER ORAL		Abdominal Pain Upper Cholelithiasis	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
		Liver Function Test Abnormal	Professional	Utrogestan (Progesterone)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/15/00ISR Number: 3514245-6Report Type:Expedited (15-DaCompany Report #033-0945-M0000022

Age:74 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Aphasia	Foreign	Neurontin	PS	Parke Davis	
Hospitalization - 1200 MG	Brain Contusion	Study			Pharmaceuticals Ltd	ORAL
Initial or Prolonged (DAILY), PER	Electroencephalogram	Health				
ORAL	Abnormal	Professional				
	Epilepsy		Gardenal			
	Fall		(Phenobarbital)	C		
	Haemorrhagic Stroke		Lioresal (Baclofen)	C		
	Hemiplegia		Tildiem (Diltazem Hydrochloride)	C		
			Nitriderm (Glyceryl Trinitrate)	C		
			Ecazide (Hydrochlorothiazide , Captopril)	C		

Date:06/16/00ISR Number: 3514712-5Report Type:Direct

Age:73 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 300MG TID	Syncope		Gabapentin	PS		
Initial or Prolonged			Percocet	C		
			Ca Acetate	C		
			Diltiazem	C		
			Prevacid	C		
			Combivent	C		
			Flexeril	C		
			Flezeril	C		
			Zoloft	C		
			Cipro	C		

Date:06/16/00ISR Number: 3514718-6Report Type:Direct

Age:50 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 900MG TID Initial or Prolonged		Syncope		Gabapentin	PS		
				Furosemide	C		
				Elavil	C		
				Baclofen	C		

Date:06/16/00ISR Number: 3515102-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000556
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other SEE IMAGE		Blindness Unilateral Blood Pressure Increased	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
SEE IMAGE		Convulsion Dizziness		Lamictal (Lamotrigine)	SS		
		Dysarthria Dysgraphia Ecchymosis Haemorrhage Hepatic Function Abnormal Hepatitis C Retinal Vein Thrombosis Speech Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/19/00ISR Number: 3515159-8Report Type:Direct
Age:64 YR Gender:Male I/FU:I

Company Report #USP 51901

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Neurontin	PS	Parke-Davis	
				Noroxin (Norfloxacin)	SS	Roberts Pharmaceutical	

Date:06/19/00ISR Number: 3516052-7Report Type:Periodic
Age:56 YR Gender:Female I/FU:F

Company Report #049-0945-M0000012

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Required		Colitis Fall	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Intervention to Prevent Permanent Impairment/Damage		Femur Fracture Gastric Ulcer Haemorrhoids Iron Deficiency Anaemia Obesity Urinary Tract Infection		Nonsteroidal Antiinrheumatic Medication Unspecified	SS		

Date:06/20/00ISR Number: 3515844-8Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Dermatitis	Health Professional	Dilantin 300mg			
Intervention to 3 PO QD				Capseal	PS		ORAL
Prevent Permanent 1 PO TID Impairment/Damage				Neurontin 100mg	SS		ORAL

Date:06/20/00ISR Number: 3515928-4Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Hepatitis Health Gabapentin PS
400 MGM QID 3 MON
Initial or Prolonged Liver Function Test Professional
Abnormal

Date:06/20/00ISR Number: 3516398-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000461
Age:63 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Back Pain	Consumer	Neurontin	PS	Parke Davis	
Initial or Prolonged	Dysuria				Pharmaceuticals Ltd	ORAL
300 MG						
Other	Nephrolithiasis					
(DAILY), PER						
	Pain					
ORAL	Pyrexia					

Date:06/20/00ISR Number: 3516404-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000446
Age:36 YR Gender:Female I/FU:F

Outcome	PT
Other	Colour Blindness
	Ear Pain
	Eye Movement Disorder
	Eye Pain

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, BID), PER ORAL		Headache Hyperthyroidism Myopathy Optic Neuropathy Photophobia Tenderness Vision Blurred Visual Acuity Reduced	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Mestinon (Pyridostigmine Bromide)	C		
				Paxil (Paroxetine Hydrochloride)	C		
				Tapazole (Thiamazole)	C		
				(Trazodone)	C		
				(Prednisone)	C		

Date:06/20/00ISR Number: 3516800-6Report Type:Expedited (15-DaCompany Report #049-0945-M0000012
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Acetabulum Fracture Arthropathy Bladder Disorder Blood Iron Decreased Bone Disorder Colitis Cystitis Escherichia Difficulty In Walking Fall Haematocrit Decreased Haemoglobin Decreased Haemorrhoids Hepatic Steatosis Microcytic Anaemia Renal Cyst Rib Fracture Spondylolisthesis Acquired Ulcer	Foreign Health Professional	Neurontin Nonsteroidal Antiinrheumatic Medication	PS SS	Parke Davis Pharmaceuticals Ltd	

Urinary Tract Infection

Date:06/21/00ISR Number: 3517187-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000585

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Gastrointestinal Carcinoma	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
200 MG DAILY				Glyburide (Glibenclamide)			
				(Glibenclamide)	C		
				Procardia (Nifedipine)	C		
				Calcium	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/21/00ISR Number: 3517277-7Report Type:Expedited (15-DaCompany Report #A013075

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Amputation	Consumer	Glucotrol	PS	Pfizer Inc	
			Health	Nifedipine	SS		
			Professional	Glucophage	SS		
				Neurontin	SS		
				Imdur	C		
				Lasix	C		
				Plavix	C		
				Prevacid	C		
				Hydralazine	C		
				Multi Vitamin	C		
				Iron	C		
				Lopid	C		

Date:06/23/00ISR Number: 3518913-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000564

Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anxiety	Consumer	Neurontin	PS	Parke Davis	
		Asthenia				Pharmaceuticals Ltd	ORAL
1500 MG		Chronic Fatigue Syndrome					
(DAILY), PER		Condition Aggravated					
ORAL 1300		Dizziness					
MG (DAILY), O		Drug Dependence		Duragesic Patch			
		Euphoric Mood		(Fentanyl)	C		
		Viral Infection		Robaxin			
				(Methocarbamol)	C		

Date:06/26/00ISR Number: 3519080-0Report Type:Direct

Company Report #

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 100MG PO QD	Nightmare	Health	Neurontin 100mg	PS	ORAL
	Vomiting	Professional			

Date:06/26/00ISR Number: 3519483-4Report Type:Expedited (15-DaCompany Report #200166
 Age:59 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 40 MG BID PO	Dehydration	Health	Oxycontin	PS	Purdue Pharma Lp	ORAL
Initial or Prolonged PO	Pneumonia Renal Failure	Professional Company Representative	Neurontin (Gabapentin)	SS		ORAL
			Medrol (Methylprednisolone)	C		
			Naprosyn (Naproxen)	C		
			Inderal (Propranolol)	C		
			Prilosec (Omeprazole)	C		
			Lotensin (Benazepril)	C		
			Indocin (Indomethacin)	C		
			Ambien (Zolpidem Tartrate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/27/00ISR Number: 3520577-8Report Type:Expedited (15-DaCompany Report #001-0945-990475

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 400 MG (200 Other MG, BID), PER ORAL	Abdominal Pain Gastric Ulcer Gastritis Glycosuria Haematochezia Malaise Nephritis Interstitial Nephrolithiasis Proteinuria Red Blood Cell Sedimentation Rate Increased Uveitis Vomiting Weight Decreased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:06/28/00ISR Number: 3521445-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000456

Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1500 MG (DAILY); 2400 MG (800 MG, TID) 1000 MG (AT BEDTIME)	Mental Disorder White Blood Cell Count Decreased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
			Depakote (Valproate Semisodium)	SS		
			Ambien (Zolpidem Tartrate)	C		

Wellbutrin
 (Amfebutamone
 Hydrochloride) C
 Lorazepam C

Date:06/30/00ISR Number: 3522928-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000384
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (100 MG BID AND 200 MG 400 MG (100 MG BID AND 200 MG HS), PER ORAL		Paraesthesia	Health Professional Other	Neurontin Neurontin Capsules 100 Mg (Gabapentin)	PS SS	Parke Davis Pharmaceuticals Ltd	ORAL
				Diphenhydramine Divalproex (Valproate Semisodium) Hydroxyzine Citalopram Clonazepam	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/30/00ISR Number: 3522929-9Report Type:Expedited (15-DaCompany Report #044-0945-M0000100

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Failure Congestive	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
300 MG (100 MG, TID) SEE IMAGE		Coronary Artery Occlusion	Professional				
600 MG (200, TID), 900 MG (300MG TID)600 MG (200 MG, TID)				Gabapentin (Gabapentin)	SS		

Date:06/30/00ISR Number: 3522931-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000616

Age:2 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Speech Disorder	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
300 MG (100 MG, TID), PER ORAL				Hydroxyzine Hydrochloride)	C		

Date:06/30/00ISR Number: 3522932-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000612

Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Coordination Abnormal Prostate Cancer	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
600 MG (300 MG, TWO TIMES				Neurontin Capsules 300 Mg (Gabapentin)	SS		ORAL
600 MG (300 MG, TWO TIMES DAILY), PER ORAL							

Glucophage (Metformin Hydrochloride)	C
Glucotrol (Glipizide)	C
Lopid (Gemfibrozil)	C
Vitamin E (Tocopherol)	C
Flomax (Morniflumate)	C

Date:06/30/00ISR Number: 3523184-6Report Type:Expedited (15-DaCompany Report #034-0945-M0000032
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG DAILY PER ORAL		Dizziness Mouth Ulceration Vomiting	Foreign Health Professional	Neurontin (Hydrochlorothiazide , Amiloride)	PS C	Parke Davis Pharmaceuticals Ltd	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/30/00ISR Number: 3583462-1Report Type:Periodic Company Report #2000032
 Age:15 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
500 MG 1/D PO		Anorexia	Consumer	Keppra	PS	Ucb Pharma Inc	ORAL
250 MG 2/D PO		Depression		Keppra	SS		ORAL
300 MG DAILY PO		Hallucination		Neurontin	SS		ORAL
		Insomnia					
		Nausea					
		Sedation					
		Vertigo					

Date:07/03/00ISR Number: 3523547-9Report Type:Expedited (15-DaCompany Report #M0456-2000
 Age:57 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 225 MG PO		Coma	Health Professional	Remeron	PS	Organon Inc Sub Akzona Inc	ORAL
Initial or Prolonged Required		Suicide Attempt		Benadryl	SS		
Intervention to Prevent Permanent Impairment/Damage				Neurontin	SS		
				Prozac	SS		
				Unisom	SS		
				Lipitor	C		
				Tricor	C		

Date:07/03/00ISR Number: 3524069-1Report Type:Expedited (15-DaCompany Report #049-0945-M0000012
 Age:56 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required		Acetabulum Fracture Bladder Disorder	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Intervention to Prevent Permanent Impairment/Damage		Blood Iron Decreased Colitis Congenital Spondylolysis Escherichia Infection		Nonsteroidal Antiinrheumatic Medication	SS		

Fall
Gastric Ulcer
Haematocrit Decreased
Haemorrhoids
Hepatic Steatosis
Microcytic Anaemia
Musculoskeletal Disorder
Oesophageal Ulcer
Pain
Renal Cyst
Urinary Tract Infection
Urine Analysis Abnormal

Date:07/03/00ISR Number: 3524082-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000421
Age:72 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Cerebral Artery Occlusion
Initial or Prolonged	Drug Interaction
Other	Fatigue
	Headache
	Loss Of Consciousness
	Malaise
	Rosacea

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Freedom Of Information (FOI) Report

Dose	Duration	Transient Ischaemic Attack Vertigo	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		Weight Decreased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
325 MG (DAILY), PER ORAL				Aspirin	SS		ORAL
				Premarin	C		
				Aldactone	C		
				Prilosec	C		
				Allopurinol	C		
				Wellbutrin	C		
				Diamox	C		
				Oxybutynin	C		
				Synthroid	C		
				Kaon Cl 10	C		
				Aspirin	C		
				Mylosine	C		

Date:07/03/00ISR Number: 3524605-5Report Type:Expedited (15-DaCompany Report #A020973
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent TOTAL:DAILY Impairment/Damage 100.00 MG		Anxiety Chest X-Ray Abnormal	Consumer	Zoloft	PS	Pfizer Pharmaceuticals Inc	
200.00 MG TOTAL: BID		Confusional State					
		Fear		Celebrex	SS		
		Influenza Like Illness					
900.00 MG		Insomnia		Neurontin	SS		

TOTAL:DAILY	Interstitial Lung Disease			
PRN	Lung Disorder	Trazodone	SS	
	Pain	Nicoderm (Subject Drug)	C	
	Photophobia	Tylenol (Subject Drug)	C	
	Scar			

Date:07/05/00ISR Number: 3524156-8Report Type:Direct Company Report #

Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Agitation		Neurontin 100mg	PS		ORAL
100MG 3 X DAY	Convulsion					
ORAL	Drug Interaction					
	Dysarthria					
	Injury					
	Jaw Disorder					
	Loss Of Employment					
	Therapeutic Agent					
	Toxicity					
	Tremor					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Brain Neoplasm	Consumer	Dilantin	PS	Parke Davis Div	
Initial or Prolonged	Brain Oedema				Warner Lambert Co	ORAL
300-600 MG						
Other	Convulsion					
DAILY, PER						
Required	Drug Level Above					
ORAL						
Intervention to	Therapeutic		Neurontin			
Prevent Permanent	Haematemesis		(Gabapentin	SS		
Impairment/Damage	Meningioma		Slow Mag Magnesium	C		
	Nervous System Disorder		Detropan	C		
	Sedation					
	Weight Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/07/00ISR Number: 3526414-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000603

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Concussion	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
200 MG (100 MG, BID), PER ORAL		Confusional State Convulsion					
3600 MG (1200 MG TID), PER ORAL		Dental Caries Ecchymosis Enamel Anomaly Fall		Neurontin Tablets 600 Mg (Gabapentin)	SS		ORAL
		Haemorrhage Head Injury Laceration Migraine Tooth Disorder Tooth Resorption		Klonopin (Clonazepam) Paxil (Paroxetine Hydrochloride)	C C		

Date:07/07/00ISR Number: 3526415-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000653

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaphylactoid Reaction	Health Professional Other	Neurontin Vibramycin (Doxycycline Hyclate)	PS SS	Parke Davis Pharmaceuticals Ltd	

Date:07/07/00ISR Number: 3526416-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000652

Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haematocrit Decreased	Health	Neurontin	PS	Parke Davis	

1200 MG (400

MG, TID), PER

ORAL

Paxil (Paroxetine Hydrochloride)	C
Glucophage (Metformin Hydrochloride)	C
Klonopin (Clonazepam)	C
Prilosec (Omeprazole)	C
Diovan (Valsartan)	C
Cardura (Doxazosin Mesilate)	C
Ditropan (Oxybutynin)	C
Vioxx (Rofecoxib)	C
Furosemide	C

Drugs

Depakote (Valproate Semisodium) SS

1500 MG

(DAILY)

Folic Acid C
Prenatal Vitamins (Ergocalciferol, Ascoric Acid, Pyridoxine Hydrochloride, Baby Aspirin (Acetylsalicylic Acid) C

Date:07/11/00ISR Number: 3527372-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000661

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Complications Of Maternal Exposure To Therapeutic	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
PLACENTAL; IN UTERO EXPOSURE		Drugs Intra-Uterine Death		Depakote (Valproate			

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Freedom Of Information (FOI) Report

Semisodium) SS

PLACENTAL IN

UTERO

EXPOSURE

Date:07/11/00ISR Number: 3527375-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000659
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Depakote (Valproate Semisodium)	SS		
				...	C		
				...	C		

Date:07/11/00ISR Number: 3527788-6Report Type:Expedited (15-DaCompany Report #JRFUSA2000004632
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Health	Duragesic	PS	Alza Corp	
TRANSDERMAL	75, TRANSD	Overdose	Professional	Paroxetine (Paroxetine)	SS		
				Carisoprodol (Carisoprodol)	SS		
				Methadone (Methadone)	SS		
				Alprazolam (Alprazolam)	SS		
				Propoxyphene (Dextropropoxyphene)	SS		
				Gabapentin (Gabapentin)	SS		
				Meperidine Hydrochloride (Pethidine Hydrochloride)	SS		

Date:07/12/00ISR Number: 3527368-2Report Type:Expedited (15-DaCompany Report #033-0945-M0000036
Age:11 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Mydriasis	Foreign Study	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
SEE IMAGE			Health Professional	Magne-Be (Pyridoxine Hydrchloride, Magnesium Lactate)	C		

Date:07/14/00ISR Number: 3529931-1Report Type:Expedited (15-DaCompany Report #033-0945-M0000060
Age:32 YR Gender:Female I/FU:I

Outcome	PT
Other	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Condition Aggravated Intra-Uterine Death Mediastinum Neoplasm

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Respiratory Tract Neoplasm Teratoma	Report Source	Product	Role	Manufacturer	Route
1600 MG	(DAILY), PER		Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
600 MG	(DAILY), PER			Carbamazepine	SS		ORAL

Date:07/14/00ISR Number: 3529933-5Report Type:Expedited (15-DaCompany Report #033-0945-M0000061
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Life-Threatening TRANSPLACENTAL Congenital Anomaly PLACENTAL IN Other UTERO EXPOSURE	SEE TEXT,	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Multiple Congenital Abnormalities	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
TRANSPLACENTAL PLACENTAL IN UTERO EXPOSURE	SEE TEXT,	Neuroblastoma Teratoma		Carbapazepine	SS		

Date:07/17/00ISR Number: 3530654-3Report Type:Expedited (15-DaCompany Report #048-0945-M0000001
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Complications Of Maternal Exposure To Therapeutic	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
TRANSPLACENTAL	SEE TEXT,	Drugs	Professional				
PLACENTAL		Stillbirth		(Carbamazepine(C		

Date:07/17/00ISR Number: 3530655-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000684
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Clonic Convulsion Grand Mal Convulsion	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
		Loss Of Consciousness					
		Salivary Hypersecretion					
		Urinary Incontinence		(Oxycodone)	C		
				(Ibuprofen)	C		
				Remeron			
				(Mirtazapine)	C		

Date:07/17/00ISR Number: 3530680-4Report Type:Expedited (15-DaCompany Report #00F--10588
Age:32 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Other	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Intra-Uterine Death Neoplasm

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pregnancy Teratoma	Report Source	Product	Role	Manufacturer	Route
800 MG DAILY			Foreign Health Professional	Tegretol-Xr	PS	Novartis Pharmaceuticals Corp	ORAL
1600 MG DAILY			Other	Neurontin Capsule (Gabapentin)	SS		ORAL

Date:07/20/00ISR Number: 3532098-7Report Type:Direct
Age:20 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dyspnoea Feeling Jittery		Neurontin 300mg (Parke-Davis)	PS	Parke Davis	ORAL
TID PO (300MG)		Hyperventilation					

Date:07/20/00ISR Number: 3532361-XReport Type:Direct
Age:66 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 300 4 ORAL		Gingival Bleeding		Neurontin 300mg	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage		Medication Error Retinal Haemorrhage		Meclomen Premarin Claritin	C C C		

Date:07/20/00ISR Number: 3532568-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000603
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Amnesia	Health	Neurontin	PS	Parke Davis	
Other		Bronchitis	Professional			Pharmaceuticals Ltd	ORAL
300-3000 MG		Concussion					
(DAILY), PER		Confusional State					
ORAL;3600 MG		Convulsion					
9DAILY), PER		Dental Caries					
ORAL		Ecchymosis		Klonopin			
		Fall		(Clonazepam)	C		
		Haemorrhage		Paxil (Paroxetine			
		Head Injury		Hydrochloride)	C		
		Injury		Premarin (Estrogens			
		Laceration		Conjugated)	C		
		Loss Of Consciousness					
		Migraine					
		Oedema					
		Respiratory Disorder					
		Tooth Disorder					
		Tooth Extraction					
		Tooth Injury					
		Weight Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/21/00ISR Number: 3533344-6Report Type:Direct
Age:58 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Sexual Dysfunction		Gabapentin	PS		

Date:07/25/00ISR Number: 3534544-1Report Type:Direct
Age:45 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthralgia		Neurontin 300mg Tid			
Other		Depression		X 7	PS		
TID 300MG	7 DAY	Joint Stiffness Joint Swelling Listless					

Date:07/25/00ISR Number: 3535251-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000704
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Joint Dislocation Urinary Incontinence	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
900 MG (300 MG, TID), PER ORAL				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Advil (Ibuprofen)	C C		

Date:07/27/00ISR Number: 3536358-5Report Type:Expedited (15-DaCompany Report #055-0945-M0000003
Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Lymphopenia	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
800 MG, PO			Professional	Captopril	C		
				Nifedipine	C		
				Hydrochlorothiazide	C		
				Glibenclamide	C		

Date:07/27/00ISR Number: 3536593-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000652
Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Haematocrit Decreased Haemoglobin Decreased	Health Professional	Neurontin	PS	Boston Life Sciences Inc	ORAL
1200 MG (400 MG, TID), PER							
ORAL				Paxil (Paroxetine Hydrochloride)	C		
				Glucophage (Metformin Hydrochloride)	C		
				Klonopin (Clonazepam)	C		
				Prilosec			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Omeprazole) C
 Diovan (Valsartan) C
 Cardura (Doxazosin Mesilate) C
 Ditropan (Oxybutynin) C
 Vioxx (Rofecoxib) C
 Furosemide C

Date:07/28/00ISR Number: 3537248-4Report Type:Direct
 Age:70 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Faecal Incontinence		Neurontin	PS		ORAL
PO 2700MG				Trental	C		
				Elavil	C		
				Vasotec	C		
				Lipitor	C		
				Glucophage	C		
				Glipizide	C		
				Tylenol With Codeine	C		
				Arthrotec	C		
				Plavix	C		

Date:07/28/00ISR Number: 3540354-1Report Type:Periodic
 Age:54 YR Gender:Male I/FU:I

Company Report #8-99104-073A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Consumer	Ponderex	PS	Ah Robins Co	ORAL
Other		Convulsion					
20 MG THREE		Lethargy					
TIMES DAILY,		Syncope		Neurontin	SS		
ORAL							
2 TABLETS							
EVERY 4 HOURS							
AS NEEDED							

40 MG DAILY
2 TABLETS
EVERY 4 HOURS
AS NEEDED
50MG AT
BEDTIME

Prozac SS
Ultram SS
Amitriptyline SS
Propulsid C
Tagamet C

Date:07/31/00ISR Number: 3537477-XReport Type:Direct
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Chest Discomfort Dyspnoea		Neurontin 300mg Parke Davis	PS	Parke Davis	ORAL
300MG 1 DOSE		Dystonia					
ORAL		Myalgia Sedation Suffocation Feeling Swelling		Plaquenil	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/01/00ISR Number: 3538330-8Report Type:Expedited (15-DaCompany Report #2000UW02610
Age:89 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 50 MG DAILY	Hypoventilation Respiratory Depression	Foreign Health Professional	Elavil	PS	Astrazeneca Pharmaceuticals Lp	ORAL
PO 2700 MG DAILY		Other	Gabapentin	SS		ORAL
PO 2400 UG TP			Fentanyl	SS		

Date:08/01/00ISR Number: 3539513-3Report Type:Expedited (15-DaCompany Report #044-0945-M0000114
Age:1 DY Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly TRANSPLACENTAL PLACENTAL (IN UTERO EXPOSURE)	Complications Of Maternal Exposure To Therapeutic Drugs Congenital Anomaly	Foreign Health Professional	Neurontin (Valproate Sodium)	PS	Parke Davis Pharmaceuticals Ltd	SS
TRANSPLACENTAL PLACENTAL (IN UTERO EXPOSURE)						

Date:08/02/00ISR Number: 3540047-0Report Type:Expedited (15-DaCompany Report #032-0945-M0000002
Age:32 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	Depression	Foreign	Neurontin	PS	Parke Davis	

Initial or Prolonged
1200 MG

(DAILY), PER

ORAL

Study

Health

Professional

Pharmaceuticals Ltd ORAL

Valproate (Valproate
Bismuth) C
Clonazepam C
Venlafaxine C
Dicalium
Chlorazepate C

Date:08/02/00ISR Number: 3540480-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000603
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Amnesia	Health	Neurontin	PS	Parke Davis	
Other		Bronchitis	Professional			Pharmaceuticals Ltd	ORAL
300-3000 MG		Bronchospasm					
(DAILY), PER		Concussion					
ORAL; DOSE		Confusional State					
INCREASE TO		Convulsion					
3600 MG		Dental Caries		Klonopin	C		
		Dyspnoea		Paxil	C		
		Ecchymosis		Premarin	C		
		Fall					
		Haemorrhage					
		Loss Of Consciousness					
		Migraine					
		Oedema					
		Tooth Disorder					
		Weight Increased					

Bromide)

C

Date:08/09/00ISR Number: 3546777-9Report Type:Expedited (15-DaCompany Report #033-0945-M0000066
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Coma	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
200 MG DAILY		Dyspnoea					
PER ORAL		Laryngeal Disorder	Professional				
		Lung Disorder		Laroxyl			
		Respiratory Depression		(Amitriptyline Hydrochloride)	SS		ORAL
25 MG DAILY							
PER ORAL							
				Lioresal (Baclofen)	SS		ORAL
15 MG DAILY							
PER ORAL							
				Lexomil (Bromazepam)	SS		ORAL
9 MG DAILY							
PER ORAL							
				Levothyrox			
				(Levothyroxine Sodium)	C		
				Diffu-K (Potassium Chloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/00ISR Number: 3546781-0Report Type:Expedited (15-DaCompany Report #032-0945-M0000023

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Myocardial Infarction Vasospasm	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:08/09/00ISR Number: 3546783-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000721

Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 900 MG (300 MG, TID) PER ORAL		Chest Pain Hallucinations, Mixed Nausea	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Vasotec (Enalapril Maleate)	C
Prilosec (Omeprazole)	C
Inderal (Propranolol Hydrochloride)	C
Buspar (Buspirone Hydrochloride)	C
Wellbutrin (Amfebutamone Hydrochloride)	C
Depakote (Valproate Semisodium)	C

Date:08/09/00ISR Number: 3546786-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000653

Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 300 MG (ONE		Anaphylactic Reaction Dyspnoea	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

DOSE ONLY)	Throat Tightness	Other	
PER ORAL	Urticaria		
100 MG (ONE		Vibramycin (Doxycycline Hyclate)	SS
DOSE ONLY)		(Guaifenesin)	C
		Bupropion	
		(Amfebutamone)	C
		(Simvastatin)	C
		(Atenolol)	C
		(Liothyronine)	C
		Enteric Coated Aspirin	
		(Acetylsalicylic Acid)	C

Date:08/09/00ISR Number: 3546788-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000220
Age:84 YR Gender:Female I/FU:I

Outcome	PT
Other	Deafness
	Hearing Impaired

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Macular Degeneration Vision Blurred	Report Source	Product	Role	Manufacturer	Route
100 MG DAILY PER ORAL;			Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1200 MG (400 MG TID) PER ORAL; 900 MG				(Choline Magnesium Trisalicylate)	C		

Date:08/10/00ISR Number: 3548299-8Report Type:Expedited (15-DaCompany Report #00F--10677
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 15 MG DAILY Initial or Prolonged ORAL		Coma Depressed Level Of Consciousness	Foreign Health Professional	Lioresal	PS	Novartis Pharmaceuticals Corp	ORAL
9 MG DAILY ORAL		Drug Interaction Dyspnoea Laryngeal Disorder	Other	Lexomil Tablet (Bromazepam)	SS		ORAL
25 MG DAILYORAL		Lung Disorder Overdose Respiratory Depression Respiratory Disorder		Laroxyl Tablet (Amitriptyline Hydrochloride)	SS		ORAL
200 MG DAILY ORAL				Neurontin Capsule (Gabapentin)	SS		ORAL
				Diffu-K Capsule	C		

Date:08/14/00ISR Number: 3550651-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000684
 Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Clonic Convulsion Grand Mal Convulsion	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1200 MG (400 MG TID) PER ORAL		Loss Of Consciousness Urinary Incontinence					
				(Oxycodone)	C		
				(Ibuprofen)	C		
				Remeron (Mirtazapine)	C		

Date:08/14/00ISR Number: 3550652-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000742
 Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Pain Sensory Loss	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
900 MG (300 MG TID) :		Transient Ischaemic Attack					
2400 MG (600 MG QID)		Vomiting Vulval Disorder					
600 MG (100 MG PRN EVERY 4 HOURS)				Ultram (Tramadol Hydrochloride)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/14/00ISR Number: 3550701-2Report Type:Expedited (15-DaCompany Report #032-0945-M0000017

Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Lung Disorder	Foreign Study	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1) 400 MG, 2) 800 MG, 3) 1200 MG, 4) 1600 MG, 5) 2000 MG, 6)			Health Professional	(Valproate Sodium) (Phenytoin) (Vigabatrin)	C C C		

Date:08/14/00ISR Number: 3550704-8Report Type:Expedited (15-DaCompany Report #044-0945-M0000094

Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Bronchospasm Oedema Peripheral	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1)300 MG (DAILY), PER ORAL; 2) 600 MG (DAILY), PER ORAL; 3)			Professional	Mst (Morphine Sulfate) (Diazepam) Voltarol (Diclofenac Sodium) Prothiaden (Dosulepin)	C C C C		

Date:08/15/00ISR Number: 3550788-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000748
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Failure Congestive	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
900 MG		Condition Aggravated	Company				
(DAILY)	1	WK	Representative				

Date:08/15/00ISR Number: 3550789-9Report Type:Expedited (15-DaCompany Report #033-0945-M0000021
Age:87 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Blood Bilirubin Increased	Foreign	Neurontin	PS	Parke Davis	
Hospitalization -		Cholestasis	Health			Pharmaceuticals Ltd	
100 MG		Diarrhoea	Professional				
Initial or Prolonged INCREASED TO		Hepatitis					
600 MG		Liver Function Test Abnormal		Renitec (Enalapril Maleate)	C		
		Pyrexia		Lasilix (Fursemide)	C		
		Vomiting		Nitriderm (Glyceryl Trinitrate)	C		
				Loxen (Nicardipine)	C		
				Digoxine (Digoxin)	C		
				Asasantine (Acetylsalicylic Acid, Dipyridamole)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/15/00ISR Number: 3551074-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000534

Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1100 MG (DAILY), PER ORAL		Condition Aggravated Confusional State Decreased Appetite Delusion	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
600 MG (DAILY), PER ORAL		Difficulty In Walking Drug Level Above Therapeutic Feeding Disorder		Neurontin (Gabapentin)	SS		ORAL
		Hallucination, Visual Psychotic Disorder Vascular Dementia Visual Disturbance		Zoloft (Sertraline Hydrochloride) Synthroid (Levothyroxine Sodium) Prinivil (Lisinopril) Lipitor (Atorvastatin) Os-Cal (Ergocalciferol, Calcium) Risperdal (Risperidone)	C C C C C C		

Date:08/15/00ISR Number: 3551075-3Report Type:Expedited (15-DaCompany Report #033-0945-M0000065

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (DAILY), PER		Infection Leukopenia Lung Disorder	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Neutropenia

ORAL

Nootropyl
(Piracetam)

C

Date:08/15/00ISR Number: 3551076-5Report Type:Expedited (15-DaCompany Report #032-0945-M0000010
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Foreign Study	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
ORAL			Health Professional	(Carbamazepine)	SS		
				(Loprazolam)	SS		
				Sertralin			
				(Sertraline)	C		
				(Periciazin			
				(Periciazine)	C		
				(Prothipendyl)			
				(Prothipendyl)	C		

Date:08/16/00ISR Number: 3551317-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000488
Age: Gender:Male I/FU:I

Outcome	PT
Death	Medication Error Pulmonary Hypertension

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pulmonary Oedema

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
5600 MG (1800 MG, TID), PER ORAL		Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
			Oxycontin (Oxycodone Hydrochloride)	C		
			Unspecified Medications	C		

Date:08/16/00ISR Number: 3551899-2Report Type:Direct
Age:44 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Hepatic Enzyme Increased		Neurontin (Gabapentin)	PS		
VARIOUS				Baclofen	C		

Date:08/17/00ISR Number: 3552349-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000616
Age:2 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Speech Disorder Vomiting	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
300 MG (100 MG, TID) PER ORAL				Depakote (Valproate Semisodium)	SS		
				Hydroxyzine Hydrochloride	C		

Phenobarbital C
 Lactulose C
 Senokot (Senna Fruit) C

Date:08/18/00ISR Number: 3589082-7Report Type:Periodic Company Report #2000025335US
 Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
INTRAVENOUS	600 MG, BID;	Blood Creatinine Increased	Health Professional	Zyvox	PS	Pharmacia And Upjohn Co	
IV				Gabapentin (Gabapentin)	SS		
2 DAY				Prozac (Fluoxetine Hydrochloride)	C		
				Buspar (Buspirone Hydrochloride)	C		
				Ambien (Zilpidem Tartrate)	C		
				Benadryl (Diphenhydramine Hydrochloride)	C		
				Ferrous Sulfate (Ferrous Sulfate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/21/00ISR Number: 3554563-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000782

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

7200 MG (2400

MG, TID), PER

ORAL

Nortriptyline (Nortriptyline)	SS
Duragesic (Fentanyl)	SS
Oxycontin (Oxycodone Hydrochloride)	C
Paxil (Paroxetine Hydrochloride)	C
Allopurinol (Allopurinol)	C
Sudafed (Pseudoephedrine Hydrochloride)	C
Clonapin (Clonazepam)	C
Benadryl (Diphenhydramine Hydrochloride)	C
Lomotil (Atropine Sulfate, Diphenoxylate Hydrochloride)	C
Vioxx (Rofecoxib)	C

Date:08/21/00ISR Number: 3554633-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000775

Age:90 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Gastrointestinal Haemorrhage Prothrombin Time Prolonged	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Coumadin (Warfarin Sodium)	C
Lasix (Furosemide)	C

Glucotrol	
(Glipizide)	C
Insulin	C
Benadryl	
(Diphenhydramine	
Hydrochloride)	C
Lanoxin (Digoxin)	C

Date:08/21/00ISR Number: 3554634-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000781
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
SEE IMAGE				Nortriptyyline (Nortriptyline)	SS		
				Methadone (Methadone)	C		
				Topamax (Topiramate)	C		
				Xanax (Alprazolam)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Effexor (Venlafaxine Hydrochloride) C

Date:08/23/00ISR Number: 3556467-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000488
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
5600 MG (1800 MG, TID), PER ORAL		Pulmonary Hypertension Pulmonary Oedema					

Oxycontin (Oxycodone Hydrochloride) C
Unspecified Medications C

Date:08/24/00ISR Number: 3557189-6Report Type:Direct Company Report #
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other				Neurontin 600mg Tablets/Parke-Davis	PS	Parke-Davis	ORAL
1T PO TID		Sedation		Oxycontin Nortriptyline Vioxx	C C C		

Date:08/24/00ISR Number: 3557541-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000684
Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1200 MG (400		Grand Mal Convulsion Loss Of Consciousness					

MG, TID), PER
 ORAL
 Movement Disorder
 Salivary Hypersecretion
 Urinary Incontinence (Oxycodone) C
 (Ibuprofen) C
 Remeron
 (Mirtazapine) C

Date:08/24/00ISR Number: 3557542-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000802
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dehydration Drug Interaction Drug Toxicity Vomiting	Health Professional	Neurontin Dilantin (Phenytoin Sodium)	PS SS	Parke Davis Pharmaceuticals Ltd	
3	YR						

Date:08/24/00ISR Number: 3557543-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000812
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200 MG (100 MG, BID)		Blood Creatine Phosphokinase Increased	Health Professional	Neurontin (Promethazine)	PS C	Parke Davis Pharmaceuticals Ltd	

Freedom Of Information (FOI) Report

Tylenol (Paracetamol)	C
Asa (Acetylsalicylic Acid)	C
(Clonidine)	C
Synthroid (Levothyroxine Sodium)	C
Betoptic (Betaxolol Hydrochloride)	C
Xalatan (Latanoprost)	C
Flonase (Fluticasone Propionate)	C
Lanoxin (Digoxin)	C
Nitro-Dur (Glyceryl Trinitrate)	C
Tiazac (Diltiazem Hydrochloride)	C
Coumadin (Warfarin Sodium)	C
Darvocet (Paracetamol, Dextropropoxyphene)	C
Albuterol (Salbutamol)	C
Ultram (Tramadol Hydrochloride)	C
Hyzaar (Hydrochlorothiazide , Losartan Potassium)	C

Date:08/24/00ISR Number: 3557597-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000818
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER		Dyspnoea Heart Rate Irregular	Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
ORAL							

Flomax
 (Morniflumate) C
 Prilosec
 (Omeprazole) C
 Unspecified Pain
 Medications C

Date:08/24/00ISR Number: 3557600-0Report Type:Expedited (15-DaCompany Report #033-0945-M0000069
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 1) 800 MG Initial or Prolonged (DAILY, PER ORAL; 2) 2000 MG (DAILY), PER PER ORAL		Cardio-Respiratory Arrest Convulsion Mydriasis	Foreign Health Professional	Neurontin Diovenor (Diosmin)	PS SS	Parke Davis Pharmaceuticals Ltd	ORAL ORAL

Freedom Of Information (FOI) Report

PER ORAL	Naramig (Naratriptan Hydrochloride)	SS	ORAL
20 MG	Zocor (Simvastatin)	SS	ORAL
(DAILY), PER			
ORAL			

Date:08/24/00ISR Number: 3557604-8Report Type:Expedited (15-DaCompany Report #047-0945-M0000003
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Cleft Lip And Palate Complications Of Maternal	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
TRANSPLACENTAL PLACENTAL (IN UTERO EXPOSURE)	SEE TEXT,	Exposure To Therapeutic Drugs	Professional				
TRANSPLACENTAL PLACENTAL (IN UTERO EXPOSURE)	SEE TEXT,	Congenital Anomaly		Frisium (Clobazam)	SS		
				(Folic Acid)	C		
				Vitamin D (Ergocalciferol)	C		

Date:08/24/00ISR Number: 3557620-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000816
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Health	Neurontin	PS	Parke Davis	

900 MG (300	Drug Withdrawal Syndrome	Professional		Pharmaceuticals Ltd	ORAL
MG, TID), PER	Grand Mal Convulsion				
ORAL	Loss Of Consciousness				
	Medication Error		Ativan (Lorazepam)	SS	
			Wellbutrin		
			(Amfebutamone		
			Hydrochloride)	C	

Date:08/24/00ISR Number: 3557675-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000196
Age:40 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Biliary Colic	Consumer	Neurontin	PS	Parke Davis	
Initial or Prolonged	Hypothyroidism				Pharmaceuticals Ltd	ORAL
900 MG (300	Liver Function Test					
MG, TID), PER	Abnormal					
ORAL;2400 MG	Ovarian Cyst					
(800 MG,	Vomiting					
TID);2100 MG			(Baclofen)	C		
			Ms Contin (Morphine			
			Sulfate)	C		

Date:08/25/00ISR Number: 3559072-9Report Type:Expedited (15-DaCompany Report #2000-06-0135
Age:39 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Amnesia
Initial or Prolonged	Dermatitis
	Drug Toxicity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
		Epilepsy Grand Mal Convulsion Hypoaesthesia					
800 MG QD		Malaise Petit Mal Epilepsy	Health Professional	Rebetol	PS	Schering Plough Research Institute	ORAL
ORAL		Pyrexia					
		Weight Decreased		Inton A (Interferon Alfa-2b Recombinant) Injectable Solution	SS		
SUBCUTANEOUS	3 MU TIW						
SUBCUTANEOUS				Dilantin	SS		
TID				Neurontin	SS		
TID				Tylenol W/Codeine	C		

Date:08/30/00ISR Number: 3562156-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000815
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Mellaril (Thioridazine Hydrochloride)	C		

Date:08/30/00ISR Number: 3562157-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000843
Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Chest Pain Dyspnoea	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1200 MG (400 MG, TID), PER		Muscle Spasms					

Panic Attack

ORAL

(Furosemide)	C
Kdur (Potassium Chloride)	C
Pravachol (Pravastatin Sodium)	C
Tylenol #3 (Codeine Phosphate, Paracetamol)	C

Date:08/30/00ISR Number: 3562158-6Report Type:Expedited (15-DaCompany Report #044-0945-M0000114
 Age:1 DY Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
TRANSPLACENTAL	PLACENTAL	Drugs	Professional				
		Foetal Valproate Syndrome	Other				

Date:08/30/00ISR Number: 3562165-3Report Type:Expedited (15-DaCompany Report #046-0945-980003
 Age:58 YR Gender:Female I/FU:F

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Epilepsy Grand Mal Convulsion	Foreign Health Professional

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Other

Dose	Duration	Product	Role	Manufacturer	Route
1200 MG (400 MG, TID);		Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
2400 MG (800 MG, TID), PER					
ORAL					

Renitec (Enalapril Maleate)	C
Seloken (Metoprolol Tartrate)	C
Trombyl (Acetylsalicylic Acid)	C
Cipramil (Citalopram Hydrobromide)	C
Lamictal (Lamotrigine)	C

Date:08/31/00ISR Number: 3562881-3Report Type:Direct
 Age:72 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 100 @HS PO	Blood Creatine Phosphokinase Increased Pain In Extremity		Neurontin 100 Mg (Parke-Davis)	PS	Parke-Davis	ORAL
THEN 100MG @ 0800						

Tylenol	C
Albuterol	C
Coumadin	C
Transderm Nitro	C
Digoxin	C
Flonase	C

Betoptic	C
Xalatan	C
Synthroid	C
Ec Aspirin	C
Ultram	C
Darvocet N-100	C
Ocean Nasal Spray	C
Clonidine	C
Apap	C
Phenergan	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
Required				Remeron	PS		ORAL
60MG PO Q HS							
Intervention to		Pulse Absent		Wellbutrin Sr	SS		ORAL
150MG PO TID							
Prevent Permanent				Neurontin	SS		ORAL
600MG 1 PO							
Impairment/Damage							
QAM AMD 2 PO							
Q HS							
200MG PO BID				Lamictal	SS		ORAL
25MCG PO QD				Cytomel	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/00ISR Number: 3571451-2Report Type:Expedited (15-DaCompany Report #00070414

Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams	Consumer	Enbrel 25 Mg	PS		
SUBCUTANEOUS	25 MG,	BIW,					
SUBCUTANEOUS		Clavicle Fracture	Health				
		Coma	Professional	Oxybutynin	SS		
		Dizziness		Atenolol/			
		Fall		Chlorthalidone	SS		
		Hallucination		Gabapentin	SS		
		Headache		Estrogen	C		
		Injection Site Pain		Prednisone	C		
		Loss Of Consciousness		Buffered Aspirin	C		
		Meniere'S Disease		Potassium Chloride	C		
		Nausea					
		Precerebral Artery Occlusion					

Date:09/05/00ISR Number: 3565734-XReport Type:Expedited (15-DaCompany Report #B0086640A

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardio-Respiratory Arrest	Foreign	Amerge	PS	Glaxo Wellcome Inc	ORAL
ORAL		Mydriasis		Diosmin (Diosmin)	SS		ORAL
ORAL				Simvastatin Tablet-Controlled Release (Simvastatin)	SS		
				Gabapentin Capsule (Gabapentin)	SS		ORAL
400 MG /	ORAL						

Date:09/07/00ISR Number: 3567398-8Report Type:Expedited (15-DaCompany Report #044-0945-M0000138

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged 300 MG (DAILY), PER ORAL	Dizziness Fall Femur Fracture Hyponatraemia Nausea	Foreign Health Professional	Neurontin (Amlodipine) (Atenolol) (Enalapril)	PS C C C	Parke Davis Pharmaceuticals Ltd	ORAL
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Date:09/07/00ISR Number: 3567399-XReport Type:Expedited (15-DaCompany Report #039-0945-M0000004
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hepatitis Non-A Non-B Non-C	Foreign	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				En (Delorazepam)	SS		
				Fevarin (Fluvoxamine Maleate)	SS		
				Rivotril (Clonazepam)	SS		
				Sereupin (Paroxetine Hydrochloride)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/07/00ISR Number: 3567609-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000881
 Age:57 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Colitis Ischaemic	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
			Dyazide (Hydrochlorothiazide , Triamterene)	C		
			Avapro (Irbesartan)	C		
			Provera (Medroxyprogesterone Acetate)	C		
			Prilosec (Omeprazole)	C		
			Estrace (Estradiol)	C		
			Amitriptyline	C		

Date:09/07/00ISR Number: 3567612-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000639
 Age:46 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER ORAL	Acute Respiratory Distress Syndrome Dermatitis Eosinophilia Pneumonia Pyrexia Respiratory Failure	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
			Dulcolax (Bisacodyl)	C		
			Colace (Docusate Sodium)	C		
			Senna (Senna)	C		
			Zantac (Ranitidine Hydrochloride)	C		
			Reglan (Metoclopramide)	C		
			Cacl	C		
			Heparin	C		
			Atrovent (Irratropium Bromide)	C		
			Albuterol			

(Salbutamol)	C
Diazepam	C
Fentanyl	C
Klonopin	
(Clonazepam)	C
Methadone	C
Decadron	
(Dexamethasone)	C

Date:09/08/00ISR Number: 3568199-7Report Type:Direct
 Age: Gender: I/FU:I

Company Report #USP 53257

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Valproic Acid	PS	Banner Pharmacaps/Udl Labs	
				Neurontin (Gabapentin) Parke-Davis	SS	Parke-Davis	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/08/00ISR Number: 3568485-0Report Type:Expedited (15-DaCompany Report #033-0945-M0000020
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Life-Threatening		Convulsion					
1200 MG			Professional				
Hospitalization - (DAILY), PER Initial or Prolonged ORAL		Limb Discomfort					
20 MG		Mydriasis					
(DAILY), PER		Sudden Death		Zocor (Simvastatin)	SS		ORAL
ORAL							
1200 MG				Diovenor (Diosmin)	SS		ORAL
(DAILY), PER							
ORAL							
PER ORAL				Naratriptan	SS		ORAL
				Tegretol (Carbamazepine)	C		

Date:09/08/00ISR Number: 3568968-3Report Type:Expedited (15-DaCompany Report #001-0184-M0000026
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Coma	Health	Benadryl			
Hospitalization - Initial or Prolonged Required		Intentional Misuse	Professional	(Diphenhydramine Hydrochloride)	PS	Parke Davis Div Warner Lambert Co	
Intervention to Prevent Permanent Impairment/Damage		Suicide Attempt		Neurontin (Gabapentin)	SS		
225 MG , PER		Toxicologic Test Abnormal		Remeron (Mirtazapine)	SS		ORAL
ORAL							

Prozac (Fluoxetine
 Hydrochloride) SS
 Unisom (Doxylamine
 Succinate) SS
 Tricor (Fenofibrate) C
 Lipitor
 (Atorvastatin) C

Date:09/11/00ISR Number: 3568218-8Report Type:Expedited (15-DaCompany Report #243804
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatitis		En	PS		
TAKEN FOR							
YEARS.							
TAKEN FOR				Fevarin	SS		
YEARS.							
TAKEN FOR				Gabapentin	SS		
YEARS.							
TAKEN FOR				Paroxetin	SS		
YEARS.							
TAKEN FOR				Rivotril	SS	Roche	
YEARS.							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/11/00ISR Number: 3569156-7Report Type:Expedited (15-DaCompany Report #061-0945-M0000007
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Apathy Confusional State	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
PER ORAL			Professional	Unspecified Medication	C		

Date:09/11/00ISR Number: 3569221-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000885
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 900 MG (300		Electroencephalogram Abnormal	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Other MG TID) PER		Epilepsy					
ORAL		Fall					
		Mental Impairment Temporal Lobe Epilepsy		Altram (Cimetidine)	C		
				Stadol (Butorphanol Tartrate)	C		
				Plavix (Clopidogrel	C		
				Carafate (Sucralfate)	C		
				Lopid (Gemfibrozil)	C		
				Zyrtec (Cetirizine Hydrochloride)	C		
				Elavil (Amitriptyline Hydrochloride)	C		
				Triavil (Perphenazine, Amitriptyline Hydrochloride)	C		
				Inderal La (Propranolol Hydrochloride)	C		
				Folic Acid	C		
				Prevacid (Lansoprazole)	C		

Propulsid
(Cisapride) C

Date:09/11/00ISR Number: 3569222-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000895
Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Respiratory Disorder	Consumer	Neurontin	PS	Parke Davis	
Other		Snoring				Pharmaceuticals Ltd	
UNKNOWN	UNKNOWN	Ventricular Fibrillation		Lipitor (Atorvastatin)	C		
				Vasotec (Enalapril Maleate)	C		
				Insulin (Insulin)	C		
				Ultram (Tramadol Hydrochloride)	C		
				Sleeping Pill (Unspecified)	C		
				Vitamin B Complex (Pyridoxine Hydrochloride,			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Thiamine
Hydrochloride, C

Date:09/12/00ISR Number: 3570455-3Report Type:Periodic Company Report #001-0945-M0000314
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged PER ORAL	Pneumonia	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:09/12/00ISR Number: 3570619-9Report Type:Expedited (15-DaCompany Report #20000900037
Age:81 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 20 MG QD Initial or Prolonged	Condition Aggravated Epilepsy Loss Of Consciousness Malaise	Foreign Health Professional	Prilosec Neurontin Idarac Praxilene	PS SS SS SS	Astrazeneca Lp	

Date:09/13/00ISR Number: 3571290-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000748
Age:64 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 800 MG (400 Other MG, BID), PER ORAL	Cardiac Failure Congestive Condition Aggravated Weight Increased	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1600 MG (800 MG, BID), PER			Neurontin (Gabapentin)	SS		ORAL

ORAL

Lasix (Furosemide)	SS
Lanoxin (Digoxin)	C
Capotin (Captopril)	C
Glynase (Glibenclamide)	C
Zaroxolyn (Metolazone)	C

Date:09/15/00ISR Number: 3571515-3Report Type:Expedited (15-DaCompany Report #243804

Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hepatitis	Foreign Other	Clonopin	PS	Hoffmann La Roche Inc	ORAL
2 MG 3 PER							
DAY ORAL							
				En (Delorazepam) 0.5 Mg	SS		ORAL
0.5 MG 1 PER							
DAY ORAL							
				Fevarin (Fluvoxamine) 100 Mg	SS		ORAL
150 MG 1 PER							
DAY ORAL							
				Gabapentin (Gabapentin)	SS		
				Paroxetine (Paroxetine)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/15/00ISR Number: 3572552-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000818

Age:73 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER ORAL	Bradycardia Dyspnoea Heart Rate Irregular Tachycardia	Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
			Flomax	C		
			Prilosec	C		
			Unspecified Pain Medications	C		

Date:09/15/00ISR Number: 3572553-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000257

Age:49 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER ORAL	Back Pain Blood Pressure Fluctuation Condition Aggravated	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
	Cyst Drug Withdrawal Syndrome Tachycardia		Lsinopril (Lisinopril) Cyclobenzaprine (Cyclobenzaprine)	SS SS		
30 MG (10 MG, PRN)			Percocet Unspecified Anti Depressant	C C		

Date:09/18/00ISR Number: 3573121-3Report Type:Direct

Age:79 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Lethargy		Gabapentin	PS		

Date:09/18/00ISR Number: 3575012-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000329
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Blister Dermatitis	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
UNKNOWN	UNKNOWN	Joint Swelling Neuralgia Oedema Peripheral Staphylococcal Infection					

Date:09/19/00ISR Number: 3575155-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000742
 Age:69 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged UNK; 1800MG (600MG,TID); 900MG (300MG, TID) 600MG		Fall Hypoaesthesia Transient Ischaemic Attack Vertigo Vomiting	Health Professional	Neurontin Ultram (Tramadol Hydrochloride)	PS SS	Parke Davis Pharmaceuticals Ltd	

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Freedom Of Information (FOI) Report

(100MG;PRN/

EVERY 4

HOURS)

Date:09/19/00ISR Number: 3575156-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000250
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	400 MG (100 MG, QID), PER	Ascites Pyrexia Sepsis	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
ORAL	400 MG (100 MG, QID), PER	Urine Analysis Abnormal		Neurontin Capsules 100 Mg (Gabapentin)	SS		ORAL
				Lasix (Furosemide)	C		
				Aldactone (Spironolactone)	C		
				(Lactulose)	C		
				Noroxin (Norfloxacin)	C		
				Magnesium (Magnesium)	C		
				Tylenol Pm (Diphenhydramine, Paracetamol)	C		

Date:09/20/00ISR Number: 3576782-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000937
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Impaired Gastric Emptying	Health	Neurontin	PS	Parke Davis	

Date:09/22/00ISR Number: 3578483-9Report Type:Expedited (15-DaCompany Report #A0127729A

Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - ORAL	Duration Asthma	Consumer	Imitrex	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged	Carpal Tunnel Syndrome Cerebrospinal Fluid Leakage		Imitrex Aqueous Spray (Sumatriptan Succinate)	SS		NASAL
INTRANASAL	Chronic Fatigue Syndrome Convulsion Dizziness		Imitrex Injection (Sumatriptan Succinate)	SS		
SUBCUTANEOUS	SUBCUTANEOUS 6 MON Drug Ineffective Fibromyalgia Loss Of Consciousness Nausea Pain Tremor Visual Acuity Reduced		Gabapentin (Gabapentin) Semisodium Valproate (Divalproex Sodium) Acetazolamide (Acetazolamide) Loratadine Fluticasone Propionate Montelukast Sodium	SS SS SS C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/22/00ISR Number: 3578806-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000942
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chills Convulsion	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
300MG (DAILY)							
PER ORAL		Cystitis					
		Sedation		Nph Insulin	C		
		White Blood Cells Urine Positive		Humalong (Insulin Lispro)	C		
				Glucopahage (Metformin Hydrochloride)	C		
				Prinivil (Lisinopril)	C		
				Aciphex (Rabeprazole Sodium)	C		
				Lipitor (Atorvastatin)	C		
				Soy Isoflavones (Calcium)	C		
				Flaxseed Oil	C		

Date:09/22/00ISR Number: 3578808-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000946
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ecchymosis Hepatocellular Damage	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

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
Dose							
Hospitalization - 150 MG /		Amnesia	Consumer	Zyban	PS	Glaxo Wellcome Inc	ORAL

Initial or Prolonged
TWICE DAY /

Hypotension

Pneumonia

ORAL

Sedation
Speech Disorder
Urine Analysis Abnormal

Nifedipine
(Formulation
Unknown) SS
(Nifedipine)
Atenolol
(Formulation
Unknown) (Atenolol) SS
Gabapentin
(Formulation
Unknown)
(Gabapentin) SS

2 YR

Date:09/26/00ISR Number: 3580593-7Report Type:Periodic Company Report #001-0073-M0000285
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Withdrawal Convulsions	Consumer	Dilantin	PS	Parke Davis Div Warner Lambert Co	ORAL
400 MG (200 MG QAM AND QPM) PER ORAL		Drug Withdrawal Syndrome					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

600 MG (300 MG, QAM AND QPM), PER ORAL			Neurontin Capsules 300 Mg(Gabapentin)	SS		ORAL
			..	C		
			..	C		

Date:09/26/00ISR Number: 3580611-6Report Type:Periodic Company Report #001-0073-990364
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (100 MG, 3QD), PER ORAL		Hypersensitivity	Consumer	Dilantin	PS	Parke Davis Div Warner Lambert Co	ORAL

900 MG (300 MG, TID), PER ORAL				Neurontin Capsules 300 Mg (Gabapentin)	SS		ORAL
				Norvasc	C		
				Azmacort	C		
				Proventil	C		
				Prilosec	C		
				Tenormin	C		
				Celebrex	C		

Date:09/26/00ISR Number: 3580645-1Report Type:Periodic Company Report #001-0073-M0000022
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Condition Aggravated	Consumer	Dilantin	PS	Parke Davis Div	

Initial or Prolonged SEE TEXT, PER	Confusional State			Warner Lambert Co	ORAL
	Convulsion				
ORAL	Nasal Congestion				
50 MG (QOD),			Dilantin Infatabs 50mg (Phenytoin Sodium)	SS	ORAL
PER ORAL					
300 MG (100			Neurontin Capsules 100 Mg (Gabapentin)	SS	ORAL
MG, TID), PER					
ORAL					
			Donnatal	SS	
			Verapamil	C	
			Prempro	C	

Date:09/27/00ISR Number: 3581564-7Report Type:Expedited (15-DaCompany Report #001-0184-M0000026
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent 225 MG (UNK), Impairment/Damage PER ORAL		Coma Laboratory Test Abnormal Suicide Attempt	Health Professional	Benadryl Hcl Neurontin (Gabapentin) Remeron (Mirtazapine)	PS SS SS	Parke Davis Div Warner Lambert Co	ORAL
				Prozac (Fluoxetine Hydrochloride)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Unisom (Doxylamine Succinate) SS
 Tricor (Fenofibrate) C
 Lipitor (Atorvastatin) C

Date:09/27/00ISR Number: 3581715-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000949
 Age:1 DY Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly Required		Arrhythmia Neonatal Cardiac Disorder	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
TRANSPLACENTAL, IN	SEE TEXT,	Complications Of Maternal					
Prevent Permanent UTERO Impairment/Damage EXPOSURE		Exposure To Therapeutic Drugs					
TRANSPLACENTAL, IN	SEE TEXT,	Heart Block Congenital Neonatal Disorder		Keppra (Levetiracetam)	SS		
UTERO EXPOSURE							

Date:09/28/00ISR Number: 3582746-0Report Type:Direct Company Report #
 Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 400MG PO		Asthenia		Gabapentin Cap Oral	PS		ORAL
Initial or Prolonged		Mental Impairment Tremor		Acetaminophen Supp,Rtl	C		
				Human Nph Insulin Inj	C		
				Sulfamethox 400/Trimethop 80mg	C		

Metoprolol C
 Lansoprazole C
 Digoxin C
 Piperacillin/Tazobactam C
 Human Regular Insulin U-100 C

Date:09/28/00ISR Number: 3582912-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000885

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG,TID); ORAL		Clonic Convulsion Complex Partial Seizures	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
200 MG (50 MG, QID); ORAL		Electroencephalogram Abnormal Epilepsy Fall Mental Impairment		Ultram (Tramadol Hydrochloride)	SS		ORAL
50 MG (QHS)		Simple Partial Seizures		Elavil(Amitriptyline Hydrochloride)	SS		
1700 MG (425 MG,QID)				Triavil(Perphenazine , Amitriptyline Hydrochloride)	SS		
				Lopid (Gemfibrozil)	C		

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Freedom Of Information (FOI) Report

Plavix (Clopidogrel) C
 Carafate
 (Sucralfate) C
 Stadol (Butorphanol
 Tartrate) C
 Zyrtec (Cetirizine
 Hydrochloride) C
 Inderal La
 (Propranolol
 Hydrochloride) C
 (Folic Acid) C
 Prevacid
 (Lansoprazole) C
 Propulsid
 (Cisapride) C

Date:09/28/00ISR Number: 3582950-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000038
 Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Corneal Oedema Visual Acuity Reduced	Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
900 MG (300 MG, TID), PER ORAL							

Contact Lens SS
 Alphagan
 (Brimonidine
 Tartrate) C
 Timoptic Xe (Timolol
 Maleate) C
 Prinivil
 (Lisinopril) C
 Calan (Verapamil
 Hydrochloride) C
 Excedrin
 (Acetylsalicylic
 Acid, Cafeine,
 Salicylamide,
 Paracetamol) C
 Vancenase
 (Beclometasone

Dipropionate) C
Prilosec
(Omeprazole) C

Date:09/28/00ISR Number: 3583024-6Report Type:Expedited (15-DaCompany Report #033-0945-M0000020
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia	Foreign	Neurontin	PS	Parke Davis	
Life-Threatening		Cardio-Respiratory Arrest	Health			Pharmaceuticals Ltd	ORAL
SEE IMAGE							
Hospitalization -		Convulsion	Professional	Zocor (Simvastatin)	SS		ORAL
20 MG							
Initial or Prolonged		Mydriasis					
(DAILY), PER							
		Sudden Death					
ORAL							
				Diovenor (Diosmin)	SS		ORAL
1200 MG							
(DAILY), PER							
ORAL							
				Naratriptan	SS		ORAL
PER ORAL							
				Tegretol	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/29/00ISR Number: 3584736-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000188

Age: Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Drug Interaction	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Other	Respiratory Arrest	Company Representative	(Methadone)	SS		

Date:09/29/00ISR Number: 3584739-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000998

Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged (DAILY) PER	Convulsion Fall	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
ORAL	Loss Of Consciousness		Cognex (Tacrine Hydrochloride)	C		
			Detrol (Tolterodine Tartrate)	C		
			Selenium	C		
			Vitamin E (Tocopherol)	C		

Date:09/29/00ISR Number: 3584741-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000966

Age:47 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged 800 MG (DAILY), 1600 MG (DAILY), 3600 MG	Blindness Medication Error	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
	Optic Neuritis					
	Retinal Haemorrhage					
	Vision Blurred					

Visual Disturbance

(DAILY), 4800

Date:10/02/00ISR Number: 3585871-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000942

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chills	Consumer	Neurontin	PS	Parke Davis	
300 MG		Convulsion				Pharmaceuticals Ltd	ORAL
(DAILY), PER		Cystitis					
ORAL		Sedation					
		White Blood Cells Urine		Nph Insulin (Insulin			
		Positive		Injection Isophane)	C		
				Humalog Insulin			
				(Insulin Lispro)	C		
				Glucophage			
				(Metformin			
				Hydrochloride)	C		
				Prinivil			
				(Lisinopril)	C		
				Aciphex (Rabeprazole			
				Sodium)	C		
				Lipitor			
				(Atorvastatin)	C		
				Soy Isoflavones	C		
				Calcium	C		
				Flaxseed Oil	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/02/00ISR Number: 3585877-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000616

Age:2 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Autism	Health	Neurontin	PS	Parke Davis	
300 MG (100		Communication Disorder	Professional			Pharmaceuticals Ltd	ORAL
MG, TID), PER		Encephalopathy					
ORAL		Mental Impairment					
		Psychomotor Hyperactivity		Depakote (Valproate			
		Speech Disorder		Semisodium)	SS		
		Vomiting		Hydroxyzine			
				Hydrchloride	C		
				Phenobarbital	C		
				Lactulose	C		
				Senokot (Senna			
				Fruit)	C		

Date:10/03/00ISR Number: 3586510-8Report Type:Direct

Company Report #

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
PO		Hyperglycaemia		Gabapentin	PS		ORAL
		Neuralgia					

Date:10/04/00ISR Number: 3587595-5Report Type:Expedited (15-DaCompany Report #001-0073-M0000261

Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Brain Neoplasm	Health	Dilantin	PS	Parke Davis Div	
Initial or Prolonged		Brain Oedema	Professional			Warner Lambert Co	ORAL
300-600 MG							
Other		Convulsion					
DAILY, PER							
Required		Haematemesis					
ORAL							
Intervention to		Meningioma		Neurontin			

Prevent Permanent Impairment/Damage	Oligodendroglioma Sedation Weight Decreased	(Gabapentin) Slow Mag (Magnesium Chloride Anhydrous) Detropan	SS C C
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Date:10/04/00ISR Number: 3587597-9Report Type:Expedited (15-DaCompany Report #001-0073-M0000120
Age:2 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abnormal Behaviour
Initial or Prolonged	Aphasia
Other	Balance Disorder Brain Scan Abnormal Clonic Convulsion Constipation Convulsion Coordination Abnormal Decreased Appetite Drooling Drug Toxicity Dystonia Ear Infection Electroencephalogram Abnormal Eye Rolling Fall

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue Gait Disturbance Gastrointestinal Disorder	Report Source				
		Gingival Bleeding Gingival Pain	Consumer	Dilantin	PS	Parke Davis Div Warner Lambert Co	ORAL
SEE IMAGE							
		Grand Mal Convulsion Insomnia Joint Stiffness		Dilantin Injection Ampoules Ml (Phenytoin Sodium)	SS		
INTRAVENOUS	SEE IMAGE						
		Lethargy Motor Dysfunction		Dilantin Suspension (Phenytoin Sodium)	SS		ORAL
2 MILLILITERS							
		Nervous System Disorder					
(BID), PER							
		Neurosis					
ORAL							
		Nystagmus Otorrhoea		Dilantin Infatabs (Phenytoin Sodium)	SS		ORAL
SEE IMAGE							
		Petit Mal Epilepsy Pruritus		Neurontin (Gabapentin)	SS		ORAL
300 MG							
		Psychomotor Hyperactivity					
(DAILY), PER							
		Psychotic Disorder					
ORAL							
		Pyrexia		Phenobarbital	SS		
80 MG (40 MG,							
BID)		Respiratory Arrest					
		Status Epilepticus Tremor Vomiting		Lamictal (Lamotrigine) Tegretol (Carbamazepine)	SS C		

Date:10/05/00ISR Number: 3588846-3Report Type:Direct
Age:76 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mental Impairment		Neurontin 300g	PS		ORAL
300MG PO							

100MG PO

Celebrex 100mg Qd

SS

ORAL

Date:10/06/00ISR Number: 3590629-5Report Type:Expedited (15-DaCompany Report #001-0945-M0001009

Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Alcohol Interaction	Company	Neurontin	PS	Parke Davis	
	Hepatic Enzyme Increased	Representative			Pharmaceuticals Ltd	
UNKNOWN	UNKNOWN					
			Alcohol (Ethanol)	SS		
			Serzone (Nefazodone Hydrochloride)	C		
			Celexa (Citalopram Hydrobromide)	C		
			Seroquel (Quetiapine)	C		
			Unspecified Medications	C		

Date:10/06/00ISR Number: 3590665-9Report Type:Expedited (15-DaCompany Report #001-0945-M0001003

Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged	Drug Interaction	Health Professional	Neurontin	PS	Parke Davis	
	Mental Impairment				Pharmaceuticals Ltd	
	Renal Impairment		Celebrex (Celecoxib)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/06/00ISR Number: 3590674-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000970

Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG TID) PER ORAL : 300 MG (100 MG TID) PER ORAL	Diabetes Mellitus	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:10/06/00ISR Number: 3590675-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000937

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (DAILY) PER ORAL	Impaired Gastric Emptying	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
			Oxycontin (Oxycodone Hydrochloride)	C		
			(Amitriptyline)	C		
			Reglan (Metoclopramide)	C		
			Klonopin (Clonazepam)	C		
			Prilosec (Omeprazole)	C		
			(Iron)	C		
			Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride,	C		

Date:10/06/00ISR Number: 3590677-5Report Type:Expedited (15-DaCompany Report #001-0945-M0001021
Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Drug Interaction	Health	Neurontin	PS	Parke Davis	
Initial or Prolonged	Mental Impairment	Professional			Pharmaceuticals Ltd	
	Renal Impairment		Celebrex (Celecoxib)	SS		

Date:10/06/00ISR Number: 3590679-9Report Type:Expedited (15-DaCompany Report #001-0945-M0001020
Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Drug Interaction	Health	Neurontin	PS	Parke Davis	
Initial or Prolonged	Mental Impairment	Professional			Pharmaceuticals Ltd	
	Renal Impairment		Celebrex (Celecoxib)	SS		

Date:10/06/00ISR Number: 3590680-5Report Type:Expedited (15-DaCompany Report #001-0945-M0001019
Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Drug Interaction	Health	Neurontin	PS	Parke Davis	
Initial or Prolonged	Mental Impairment	Professional			Pharmaceuticals Ltd	
	Renal Impairment		Celebrex (Celecoxib)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/06/00ISR Number: 3590681-7Report Type:Expedited (15-DaCompany Report #001-0945-M0001018

Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Drug Interaction	Health	Neurontin	PS	Parke Davis	
Initial or Prolonged	Mental Impairment	Professional			Pharmaceuticals Ltd	
	Renal Impairment		Celebrex (Celecoxib)	SS		

Date:10/06/00ISR Number: 3590682-9Report Type:Expedited (15-DaCompany Report #001-0945-M0001016

Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Drug Interaction	Health	Neurontin	PS	Parke Davis	
Initial or Prolonged	Mental Impairment	Professional			Pharmaceuticals Ltd	
	Renal Impairment		Celebrex (Celecoxib)	SS		

Date:10/06/00ISR Number: 3601896-3Report Type:Periodic Company Report #USA015094

Age:29 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
	Nonspecific Reaction	Consumer	Meridia	PS	Knoll Pharmaceutical	ORAL
10 MG OD PO		Other			Co Sub Basf Corp	
			Neurontin	SS		ORAL
400 MG TID PO			Motrin	SS		ORAL
MG OD PO			Darvocet	SS		ORAL
2 TAB PRN PO			Percodan	SS		ORAL
1 TAB Q2HR PO			Ambien	SS		ORAL
15 MG PRN PO						

Date:10/11/00ISR Number: 3592825-XReport Type:Direct

Age:73 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Gingival Hyperplasia		Gabapentin 100mg	PS		ORAL
2TT PO TID		Gingivitis					

Date:10/11/00ISR Number: 3593414-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000942
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Consumer	Neurontin	PS	Parke Davis	
300 MG		Asthenia				Pharmaceuticals Ltd	ORAL
(DAILY), PER		Chills					
ORAL		Convulsion					
		Cystitis		Nph Insulin (Insulin			
		Hyperglycaemia		Injection, Isophane)	C		
		Joint Stiffness		Humalog Insulin			
		Sedation		(Insulin Lispro)	C		
		Tremor		Glucophage			
		White Blood Cells Urine		(Metformin			
		Positive		Hydrochloride)	C		
				Prinivil			
				(Lisinopril)	C		
				Aciphex (Rabeprazole			
				Sodium)	C		
				Lipitor			
				(Atorvastatin)	C		
				Soy Isoflavones	C		
				Calcium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Flaxseed Oil C

Date:10/11/00ISR Number: 3597496-4Report Type:Periodic Company Report #000928-SK358
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
UNKNOWN PO		Confusional State	Health	Celebrex	PS	Gd Searle And Co	ORAL
UNKNOWN PO		Drug Interaction	Professional	Neurontin	SS		ORAL

Date:10/11/00ISR Number: 3597497-6Report Type:Periodic Company Report #000928-SK359
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
UNKNOWN PO		Confusional State	Health	Celebrex	PS	Gd Searle And Co	ORAL
UNKNOWN PO		Drug Interaction	Professional	Neurontin	SS		ORAL

Date:10/11/00ISR Number: 3597498-8Report Type:Periodic Company Report #000928-SK360
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
UNKNOWN PO		Confusional State	Health	Celebrex	PS	Gd Searle And Co	ORAL
UNKNOWN PO		Drug Interaction	Professional	Neurontin	SS		ORAL

Date:10/11/00ISR Number: 3597499-XReport Type:Periodic Company Report #000928-SK361
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
UNKNOWN PO		Confusional State	Health	Celebrex	PS	Gd Searle And Co	ORAL

UNKNOWN PO Drug Interaction Professional Neurontin SS ORAL

Date:10/11/00ISR Number: 3597500-3Report Type:Periodic Company Report #000928-SK362
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Health	Celebrex	PS	Gd Searle And Co	ORAL
UNKNOWN PO		Drug Interaction	Professional	Neurontin	SS		ORAL
UNKNOWN PO							

Date:10/11/00ISR Number: 3597501-5Report Type:Periodic Company Report #000928-SK363
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Health	Celebrex	PS	Gd Searle And Co	ORAL
UNKNOWN PO		Drug Interaction	Professional	Neurontin	SS		
OPHTHALMIC	UNKNOWN PO						

Date:10/11/00ISR Number: 3597869-XReport Type:Periodic Company Report #000707-SK453
 Age: Gender:Not SpecifiedI/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Mouth Ulceration	Health	Celebrex	PS	Gd Searle And Co	ORAL
PO			Professional	Neurontin	SS		ORAL
PO							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/11/00ISR Number: 3598630-2Report Type:Periodic
Age:57 YR Gender:Female I/FU:I

Company Report #000821-SK376

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Neuropathy Peripheral	Health	Celebrex	PS	Gd Searle And Co	ORAL
200.000 MG			Professional				
BID PO				Neurontin	SS		ORAL
200.000 MG							
TID PO				Atorvastatin	C		
				Triamterene	C		
				Hydrochlorothiazide	C		
				Potassium	C		
				Zolpidem	C		
				Conjugated Estrogens	C		
				Diltiazem			
				Hydrochloride	C		
				Omeprazole	C		
				Methocarbamol	C		
				Hydrocodone	C		

Date:10/12/00ISR Number: 3595695-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000990
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anuria	Literature	Neurontin	PS	Parke Davis	
Other		Blood Creatinine	Health			Pharmaceuticals Ltd	
400 MG (200		Increased	Professional				
MG, BID), 600		Blood Urea Increased					
MG (200 MG,		Drug Interaction					
BID)		Failure Of Implant		Salsalate(Salsalate)	SS		
		Renal Impairment		Furosemide			
				(Furosemide)	SS		
				Asathioprine			
				(Azathioprine)	C		
				Macrochantin			

(Nitrofurantoin) C
 Isosorbide Dinitrate
 (Isosorbide
 Dinitrate) C
 Nph Insulin (Insulin
 Injection, Isophane) C
 Metoprolol
 (Metoprolol) C
 Lovastatin
 (Lovastatin) C
 L-Thyroxine
 (Levothyroxine) C

Date:10/13/00ISR Number: 3594860-4Report Type:Expedited (15-DaCompany Report #044-0945-M0000100
 Age:73 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Failure Congestive	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
300 MG (100 MG TID) PER ORAL; 600 MG (200 MG TID); 900 MG (300		Condition Aggravated	Professional				
		Coronary Artery Disease					
		Coronary Artery Occlusion					
		Medication Error		Trypitzol (Amitriptyline)			

Freedom Of Information (FOI) Report

Hydrochloride) C

Date:10/13/00ISR Number: 3594937-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000972
 Age:84 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hepatic Neoplasm Malignant	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
400 MG (100 MG, QID) PER ORAL		Metastases To Liver					

Zocor (Simvastatin) C
 Synthroid
 (Levothyroxine
 Sodium) C
 (Iron) C
 Vitamins C

Date:10/16/00ISR Number: 3595964-2Report Type:Expedited (15-DaCompany Report #JRFUSA2000006335
 Age:48 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability	Abdominal Pain
Other	Abnormal Behaviour
Required	Aggression
Intervention to Prevent Permanent Impairment/Damage	Angina Pectoris
	Anhedonia
	Anxiety
	Arachnoiditis
	Asthenia
	Back Pain
	Bronchitis Acute
	Cardiac Disorder
	Cardiomegaly
	Cervical Spinal Stenosis
	Chest Pain
	Condition Aggravated
	Constipation
	Coronary Artery Disease
	Cough
	Decreased Appetite

Depression
Diarrhoea
Dizziness
Dry Mouth
Dyspnoea
Echocardiogram Abnormal
Ejection Fraction
Abnormal
Electrocardiogram
Abnormal
Electrocardiogram Qt
Prolonged
Emotional Disorder
Fall
Fibrosis
Gastric Ulcer
Gastrooesophageal Reflux
Disease
Haematochezia
Hallucination

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Adverse Event	Report Source	Product	Role	Manufacturer	Route
20 MG, 4 IN 1 DAY(S), ORAL		Headache Hyperhidrosis Hypoaesthesia	Health Professional	Propulsid	PS	Janssen Research Fdn Div Johnson And Johnson	ORAL
75 MG, 1 IN 1 DAY (S), ORAL		Joint Swelling Malaise Mental Disorder		Pamelor (Nortriptyline Hydrochloride)	SS		ORAL
50 MG, 4 IN 1 DAILY, ORAL		Mitral Valve Incompetence Mononeuropathy Muscle Spasms		Promethazine	SS		ORAL
600 MG, 2 IN 1 DAILY		Myalgia Myocardial Infarction Nasal Congestion Nausea Neck Pain Nervous System Disorder Nervousness Nuclear Magnetic Resonance Imaging		Mellaril (Thioridazine Hydrochloride) Compazine (Prochlorperazine Edisylate) Neurontin (Gabapentin)	SS SS SS		
500 MG, 2 IN 1 DAILY		Abnormal Pain Pain In Extremity		Biaxin (Clarithromycin)	SS		
		Palpitations Pancreatic Disorder Pancreatitis Pharyngitis Pharyngolaryngeal Pain Polydipsia Polyp Polyuria Pruritus Pyrexia Rash Erythematous		Elavil (Amitriptyline Hydrochloride) Prilosec (Omeprazole) Paxil (Paroxetine Hydrochloride) Prozac (Fluoxetine Hydrochloride) Nitroglycerin (Nitroglycerin Comp.	SS C C C		

Rash Papular	/Net/)	C
Rectal Haemorrhage	Vicodin	C
Renal Colic	Procardia	
Rhinitis	(Nifedipine)	C
Road Traffic Accident	Valium (Diazepam)	C
Sinus Tachycardia	Prevacid	
Spinal Osteoarthritis	(Lansoprazole)	C
Suicidal Ideation	Reglan	
Swelling	(Metoclopramide)	C
Syncope	Baclofen	C
Tachycardia	Vistaril	
Tricuspid Valve	(Hydroxyzine	
Incompetence	Embonate)	C
Ultrasound Scan Abnormal	Vitamins	C
Urinary Retention	Darvocet	C
Urinary Tract Infection	Parafon (Parafon	
Ventricular Extrasystoles	Forte)	C
Ventricular Fibrillation	Flexeril	
Vision Blurred	(Cyclobenzaprine	
Vomiting	Hydrochloride)	C
Weight Decreased	Isocet (Axotal (Old	
	Form))	C
	Trazodone	C
	Amoxil (Amoxicillin)	C
	Metoprolol	C

Freedom Of Information (FOI) Report

Carbamazepine	C
Talacen (Fortagesic)	C
Nifrex-150 Forte	C
Celexa (Citalopram Hydrobromide)	C
Medrol Dose Pack (Methylprednisolone)	C
Lactulose	C
Atarax (Hydroxyzine Hydrochloride)	C
Limbitrol	C
Baclofen	C
Remeron (Mirtazapine)	C
Ms Contin (Morphine Sulfate)	C
Oxycontin (Oxycodone Hydrochloride)	C
Hemorrhoidal Suppository (Unspecified)	C
Midrin (Midrid)	C
Zithromax (Azithromycin)	C
Viokase (Pancrelipase)	C
Lorazepam	C

Date:10/17/00ISR Number: 3597073-5Report Type:Expedited (15-DaCompany Report #001-0945-M0001041
 Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Anxiety Apraxia	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
2400 MG (800 MG TID)		Condition Aggravated		Lamictal (Lamotrigine)	SS		
		Convulsion Dizziness Memory Impairment Mental Impairment Psychomotor Retardation Sedation Tearfulness					

Date:10/17/00ISR Number: 3597076-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000603
Age:54 YR Gender:Female I/FU:F

Outcome	PT
Disability	Accidental Overdose
Other	Amnesia
	Bronchitis
	Bronchospasm
	Confusional State
	Convulsion
	Dental Caries
	Dyspnoea
	Ecchymosis
	Enamel Anomaly
	Fall
	Haemorrhage

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300-3000 MG	(DAILY) PER	Laceration Migraine Oedema Tooth Injury Toxic Epidermal Necrolysis Weight Increased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
ORAL				Klonopin (Clonazepam)	C		
				Paxil (Paroxetine Hydrochloride)	C		
				Premarin (Estrogens Conjugated)	C		

Date:10/17/00ISR Number: 3597083-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000887

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Clonic Convulsion Epilepsy Insomnia	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
2700 MG (900 MG TID) PER				Neurontin (Gabapentin)	SS		ORAL
ORAL							
3600 MG (1200 MG TID) PER				Paxil (Paroxetine Hydrochloride)	C		
ORAL				Ultram (Tramadol Hydrochloride)	C		
				Aspirin (Acetylsalicylic Acid)	C		
				Baclofen (Baclofen)	C		

(Potassium) C
 Lasix (Furosemide) C
 Prilosec
 (Omeprazole) C
 Medrin (Paracetamol
 ,
 Dichloralphenazone,
 Isometheptene) C

Date:10/17/00ISR Number: 3597587-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000895
 Age:76 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Disease	Health	Neurontin	PS	Parke Davis	ORAL
Other		Medication Error	Professional			Pharmaceuticals Ltd	
600 MG (300		Sedation					
MG QPM QAM		Snoring					
PRN) PER ORAL		Ventricular Fibrillation		Lipitor (Atorvastatin)	C		
				Vasotec (Enalapril Maleate)	C		
				Insulin (Insulin)	C		
				Ultram (Tramadol Hydrochloride)	C		
				Sleeping Pill	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vitamin B Complex
 (Pyridoxine
 Hydrochloride,
 Thiamine
 Hydrochloride, C

Date:10/18/00ISR Number: 3597552-0Report Type:Expedited (15-DaCompany Report #001-0945-M0001039
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE IMAGE		Chest Pain Convulsion	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
2400 MG (600 MG, QID), PER ORAL		Dizziness Feeling Jittery		Neurontin Tablets	SS		ORAL
				Bromocriptine	C		

Date:10/20/00ISR Number: 3599374-3Report Type:Expedited (15-DaCompany Report #A030358
 Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - ORAL Initial or Prolonged		Coma Intentional Misuse	Consumer Health	Unisom	PS	Pfizer Laboratories Div Pfizer Inc	ORAL
225.00 MG Required TOTAL:ORAL Intervention to ORAL Prevent Permanent ORAL Impairment/Damage		Suicide Attempt	Professional	Remeron	SS		ORAL
				Benadryl	SS		ORAL
				Neurontin	SS		ORAL
				Lipitor	C		
				Tricor	C		
				Prozac	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (400 Required MG, TID), PER Intervention to ORAL Prevent Permanent Impairment/Damage	Condition Aggravated Fistula Haemorrhage Hypercholesterolaemia Weight Decreased	Consumer	Neurontin Premarin (Estrogens Conjugated) Claritin-D (Pseudoephedrine Sulfate, Loratadine) Robaxin (Methocarbamol) Darvocet (Paracetamol, Dextropropoxyphene) Lortab (Paracetamol, Hydrocodone Bitartrate)	PS C C C C C C	Parke Davis Pharmaceuticals Ltd	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/23/00ISR Number: 3599982-XReport Type:Expedited (15-DaCompany Report #001-0945-M0001092

Age:52 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1) 300 MG (QHS), PER ORAL; 2) 100 MG (DAILY), PO; 3) 200 MG	Bradycardia Chest Pain Dyspnoea Electrolyte Imbalance Feeling Hot	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
			Humulin N (Insulin Human Injection, Isophane)	C		
			Humulin R (Insulin Human)	C		
			Cardizem (Diltiazem Hydrochloride)	C		
			Lanoxin (Digoxin)	C		
			Zocor (Simvastatin)	C		
			Bufferin (Acetylsalicylic Acid, Magnesium Carbonate, Aluminium Glycinate)	C		
			Ntg Slow Cap (Glyceryl Trinitrate)	C		

Date:10/23/00ISR Number: 3599983-1Report Type:Expedited (15-DaCompany Report #001-0945-M0001064

Age:48 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG, TID), PER	Colour Blindness Retinal Oedema Visual Disturbance	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

ORAL

Demadex (Torasemide)	C
Zoloft (Sertraline Hydrochloride)	C
Remeron (Mirtazapine)	C
Premarin (Estrogens Conjugated)	C
Prevacid (Lansoprazole)	C
Imitrex (Sumatriptan Succinate)	C

Date:10/23/00ISR Number: 3600259-4Report Type:Expedited (15-DaCompany Report #001-0073-M0000385
 Age:1 DY Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Complications Of Maternal Exposure To Therapeutic	Health Professional	Dilantin	PS	Parke Davis Div Warner Lambert Co	
SEE TEXT,		Drugs					
PLACENTAL				Neurontin (Gabapentin)	SS		
SEE TEXT,							
PLACENTAL				Proventil	C		
				Theo-Dur	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/23/00ISR Number: 3600264-8Report Type:Expedited (15-DaCompany Report #001-0945-M0001065

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 200 MG (100 MG BID) PER ORAL; 100 MG (DAILY) PER ORAL	Balance Disorder Blood Pressure Increased Dyspnoea Haematuria Headache Hypoaesthesia Muscle Rigidity Pain Paraesthesia Urinary Tract Infection	Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
			Glucotrol (Glipizide)	C		
			Actos (Pioglitazone)	C		
			Zestril (Lisinopril)	C		
			Celebrex (Celecoxib)	C		
			Lipitor (Atorvastatin)	C		

Date:10/23/00ISR Number: 3600485-4Report Type:Expedited (15-DaCompany Report #002-0945-M0000031

Age:50 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1600 MG (DAILY), PER ORAL	Myopia Presbyopia	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:10/23/00ISR Number: 3600690-7Report Type:Expedited (15-DaCompany Report #PHRM2000FR01473

Age:25 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization -	Convulsion	Foreign	Tegretol-Xr	PS	Novartis	

Initial or Prolonged	Drug Interaction	Health	Pharmaceuticals Corp	ORAL
1200 MG/DAY,		Professional		
ORAL		Other	Neurontin(Gabapentin) Capsule	SS ORAL
2400 MG/DAY,				
ORAL			Nutrition Supplements(Nutrition Supplements)	SS ORAL
ORAL			Urbanyl	C

Date:10/24/00ISR Number: 3600703-2Report Type:Expedited (15-DaCompany Report #JRFUSA2000004632
 Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Duragesic	PS	Alza Corp	
TRANSDERMAL	11 PATCH, 1	Drug Toxicity	Health				
IN 1 TIME		Medication Error	Professional				
(S), TRANSD				Paroxetine (Paroxetine)	SS		
TRANSDERMAL	11 PATCH, 1						
IN 1 TIME				Carisoprodol (Carisoprodol)	SS		
(S), TRANSD				Methadone (Methadone)	SS		
				Alprazolam (Alprazolam)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Propoxyphene
 (Dextropropoxyphene) SS
 Gabapentin
 (Gabapentin) SS
 Meperidine
 Hydrochloride
 (Pethidine
 Hydrochloride) SS

Date:10/25/00ISR Number: 3608771-9Report Type:Periodic Company Report #A030491
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetes Mellitus	Consumer	Viagra	PS	Pfizer Agricultural Div	
	300.00 MG			Gabapentin	SS		ORAL
TOTAL:ORAL							

Atenolol C
 Diazepam C
 Trandolapril C
 Diltiazem C
 Indapamide C
 Gamolenic Acid C
 Potassium Bicarbonate C
 Amiloride C
 Allopurinol C
 Frusemide C
 Hydroxocobalamin C
 Naprosyn C
 Metformin C
 Gliclazide C

Date:10/26/00ISR Number: 3602007-0Report Type:Direct Company Report #
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatomegaly		Neurontin 200 Mg Tid	PS		ORAL
200 MG TID							

Lung Disorder

ORAL

Nortriptyline

C

Date:10/27/00ISR Number: 3603072-7Report Type:Expedited (15-DaCompany Report #001-0945-M0001107

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Health	Neurontin	PS	Parke Davis	
		Drug Dependence	Professional			Pharmaceuticals Ltd	
600 MG (300							
MG, BID)							

Date:10/27/00ISR Number: 3603073-9Report Type:Expedited (15-DaCompany Report #001-0945-M0001057

Age: Gender:Male I/FU:I

Outcome	PT
Other	Accidental Overdose
	Blood Creatinine
	Increased
	Creatinine Renal

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Freedom Of Information (FOI) Report

Clearance Decreased
Renal Impairment

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2700 MG	(DAILY)	Other	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:10/27/00ISR Number: 3603074-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000639
Age:46 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG,TID), PER ORAL	Acute Respiratory Distress Syndrome Dermatitis Eosinophilia Infection Pneumonia Pyrexia Respiratory Failure	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
			Dulcolax (Bisacodyl)	C		
			Colace (Docusate Sodium)	C		
			Senna(Senna)	C		
			Zantac (Ranitidine Hydrochloride)	C		
			Reglan (Metoclopramide)	C		
			Cacl (Heparin)	C		
			Atrovent (Ipratropium Bromide)	C		
			Albuterol (Salbutamol)	C		
			(Diazepam)	C		
			(Fentanyl)	C		
			Klonopin (Clonazepam)	C		
			(Methadone)	C		
			Decadron (Dexamethasone)	C		

Date:10/31/00ISR Number: 3604437-XReport Type:Direct
Age:66 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia		Gabapentin	PS		ORAL
300MG TID							
ORAL		Disturbance In Attention					
		Myalgia		Etanercept	C		
				Hydrochlorothiazide	C		
				Estrogens Conjugated	C		
				Ferrous Sulfate	C		
				Salsalate	C		
				Sulfasalazine	C		
				Hydroxychloroquine Sulfate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/01/00ISR Number: 3605449-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000887

Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Clonic Convulsion	Health	Neurontin	PS	Parke Davis	
SEE IMAGE		Epilepsy	Professional			Pharmaceuticals Ltd	ORAL
		Insomnia		Paxil (Paroxetine Hydrochloride)	C		
				Ultram (Tramadol Hydrochloride)	C		
				Aspirin (Acetylsalicylic Acid)	C		
				Baclofen (Baclofen)	C		
				(Potassium)	C		
				Lasix (Furosemide)	C		
				Prilosec (Omeprazole)	C		
				Midrin (Paracetamol, Dichloralphenazone, Isometheptene)	C		
				Lotanax (Terfenadine)	C		

Date:11/01/00ISR Number: 3605452-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000878

Age:64 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged	1800 MG (600	Depression	Health	Neurontin	PS	Parke Davis	
MG, TID)		Disturbance In Attention	Professional			Pharmaceuticals Ltd	
		Facial Nerve Disorder					
		Facial Pain		Oxycontin (Oxycodone Hydrochloride)	C		
		Fatigue		Clonopin (Clonazepam)	C		
		Neuralgia		Celebrex (Celecoxib)	C		
		Oedema Peripheral		Lorcet (Paracetamol, Hydrocodone Bitartrate)	C		
		Trigeminal Neuralgia					

Outcome	PT
Death	Arthropathy
Hospitalization -	Atrial Fibrillation
Initial or Prolonged	Brain Hypoxia
	Cerebellar Ataxia
	Choreoathetosis
	Coma
	Cyanosis
	Dementia
	Electroencephalogram
	Abnormal
	Epilepsy
	Joint Dislocation
	Loss Of Consciousness
	Metabolic Acidosis
	Pupil Fixed
	Respiratory Arrest

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Skin Disorder

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100 MG (ONE DOSE), PER ORAL		Foreign Health Professional	Neurontin (Phenytoin) (Clobazam) (Valproic Acid) (Insulin Porcine)	PS C C C C	Parke Davis Pharmaceuticals Ltd	ORAL

Date:11/01/00ISR Number: 3605778-2Report Type:Expedited (15-DaCompany Report #033-0945-M0000089
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL Required Intervention to Prevent Permanent PER ORAL Impairment/Damage PER ORAL PER ORAL		Anaemia Megaloblastic	Foreign Health Professional	Neurontin Asasantin (Acetylsalicylic Acid, Dipyridamole) Ventoline (Salbutamol) Mopral (Omeprazole)	PS SS SS SS	Parke Davis Pharmaceuticals Ltd	ORAL ORAL ORAL ORAL

Date:11/01/00ISR Number: 3605833-7Report Type:Expedited (15-DaCompany Report #001-0945-M0001134
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Disorientation Drug Interaction Feeling Abnormal Mania	Consumer	Neurontin Biaxin (Clarithromycin)	PS SS	Parke Davis Pharmaceuticals Ltd	

Pharyngitis Streptococcal
Stress
Vision Blurred

Unspecified
Antibiotic

SS

Date:11/02/00ISR Number: 3606194-XReport Type:Expedited (15-DaCompany Report #044-0945-M0000078

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG, (DAILY), PER ORAL	Dehydration Dizziness Hypotension Nephrotic Syndrome Oedema Peripheral Polyuria Proteinuria	Foreign Health Professional	Neurontin (Amitriptyline) Tegretol Retard (Carbamazepine) Frusemide (Furosemide) (Simvastatin)	PS C C C C	Parke Davis Pharmaceuticals Ltd	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/00ISR Number: 3606706-6Report Type:Direct
Age:71 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Constipation		Gabapentin	PS		

Date:11/03/00ISR Number: 3607276-9Report Type:Expedited (15-DaCompany Report #20001100032
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 10 MG QD PO		Anaemia Megaloblastic	Foreign	Prilosec	PS	Astrazeneca Lp	ORAL
Initial or Prolonged 2 DOSE QD PO		Blood Folate Decreased	Health	Asasantin	SS		ORAL
400 MG TID PO		Crepitations	Professional	Neurontin	SS		ORAL
RESPIRATORY (INHALATION)		Dyspnoea		Ventoline Inhalator	SS		
		Haemoglobin Decreased					
		Mean Cell Volume Increased		Bronchokod	C		
		Pallor		Vitamin B 1 And B6	C		
		Red Blood Cell Count Decreased		Doliprane	C		
		Tachycardia		Lasilix	C		
				Sodium Chloride	C		

Date:11/07/00ISR Number: 3608452-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000257
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 900 MG (300 MG, TID), PER		Back Pain	Health	Neurontin	PS	Parke Davis	
		Blood Pressure Fluctuation	Professional			Pharmaceuticals Ltd	ORAL
			Company				

ORAL

Bone Cyst

Representative

Condition Aggravated
Drug Withdrawal Syndrome
Tachycardia

Lisinopril
(Lisinopril) SS
Percocet
(Paracetamol,
Oxycodone
Hydrochloride,
Oxycodone C
Unspecified Anti
Depressant C

Date:11/07/00ISR Number: 3608453-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000966
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blindness Colour Blindness Optic Disc Haemorrhage Optic Neuritis Photopsia Retinal Haemorrhage Vision Blurred Visual Field Defect	Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/08/00ISR Number: 3609052-XReport Type:Expedited (15-DaCompany Report #2000-FF-S0618

Age:65 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 200 MG/200 MG X2/DAY/PO	Anaemia Anaemia Megaloblastic Blood Folate Decreased	Foreign Other	Aggrenox	PS	Boehringer Ingelheim Pharmaceuticals Inc	ORAL
BU 1.2 G/PO 10 MG/PO	Dyspnoea Pallor		Ventoline Neurontin Mopral Bronchokod Doliprane Lasilix Chlorure De Sodium Vitamine B1- B6	SS SS C C C C C		ORAL ORAL

Date:11/08/00ISR Number: 3609094-4Report Type:Expedited (15-DaCompany Report #033-0945-M0000081

Age:76 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Other 300 MG (DAILY)	Brain Oedema Sepsis Thrombocytopenia	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
			Unspecified Corticoids Medrol Mopral Cordarone	SS C C C		

Date:11/09/00ISR Number: 3610385-1Report Type:Expedited (15-DaCompany Report #JRFUSA2000006335

Age:48 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain

Initial or Prolonged
Disability
Other
Required
Intervention to
Prevent Permanent
Impairment/Damage

Abnormal Behaviour
Abnormal Dreams
Angina Pectoris
Anhedonia
Anxiety
Arachnoiditis
Asthenia
Back Pain
Bladder Disorder
Bronchitis
Bronchitis Acute
Cardiac Disorder
Cardiomegaly
Cervical Spinal Stenosis
Chest Pain
Colonic Polyp
Condition Aggravated
Constipation
Coronary Artery Disease
Cough
Decreased Appetite
Depression
Diarrhoea
Dizziness
Dry Mouth
Dyspnoea

Freedom Of Information (FOI) Report

Echocardiogram Abnormal
Ejection Fraction
Abnormal
Electrocardiogram
Abnormal
Electrocardiogram Qt
Prolonged
Emotional Disorder
Fall
Fibrosis
Gait Disturbance
Gastric Ulcer
Gastrooesophageal Reflux
Disease
Haematochezia
Hallucination
Hyperhidrosis
Hypoaesthesia
Hypotension
Influenza Like Illness
Injury
Insomnia
Joint Swelling
Malaise
Mental Disorder
Migraine
Mitral Valve Incompetence
Mononeuropathy
Muscle Spasms
Myalgia
Myocardial Infarction
Nasal Congestion
Nausea
Neck Pain
Nervous System Disorder
Nervousness
Nuclear Magnetic
Resonance Imaging
Abnormal
Pain
Pain In Extremity
Palpitations
Pancreatitis
Pharyngitis
Pharyngolaryngeal Pain
Polydipsia
Polyuria

Pruritus
Pyrexia
Rash Erythematous
Rectal Haemorrhage
Renal Colic
Rhinitis
Road Traffic Accident
Sinus Tachycardia
Spinal Osteoarthritis
Suicidal Ideation
Swelling
Syncope
Tricuspid Valve
Incompetence

Vicodin (Vicodin)	C
Procardia	
(Nifedipine)	C
Valium (Diazepam)	C
Prevacid	
(Lansoprazole)	C
Reglan	
(Metoclopramide)	C
Baclofen (Baclofen)	C
Vistaril	
(Hydroxyzine	
Embonate)	C
Vitamins (Vitamins)	C
Darvocet (Darvocet)	C
Parafon (Parafon	
Forte)	C
Flexeril	
(Cyclobenzaprine	
Hydrochloride)	C
Isocet (Axotal (Old	
Form))	C
Trazodone	
(Trazodone)	C
Amoxil (Amoxicillin)	C

Freedom Of Information (FOI) Report

Metoprolol	
(Metoprolol)	C
Carbamazepine	
(Carbamazepine)	C
Talacen (Fortagesic)	C
Nifrex-150 Forte	
(Unspecified)	C
Celexa (Citalopram	
Hydrobromide)	C
Lorazepam	
(Lorazepam)	C
Medrol Dose Pack	
(Methylprednisolone)	C
Lactulose	
(Lactulose)	C
Atarax (Hydroxyzine	
Hydrochloride)	C
Limbitrol	
(Limbitrol)	C
Baclofen (Baclofen)	C
Remeron	
(Mirtazapine)	C
Ms Contin (Morphine	
Sulfate)	C
Oxycontin (Oxycodone	
Hydrochloride)	C
Hemorrhoidal	
Suppository	
(Unspecified)	C
Midrin (Midrid)	C
Zithromax	
(Azithromycin)	C
Viokase	
(Pancrelipase)	C

Date:11/13/00ISR Number: 3610818-0Report Type:Expedited (15-DaCompany Report #001-0945-M0001170
 Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Asthma	Consumer	Neurontin	PS	Parke Davis	
Initial or Prolonged	Blindness Transient				Pharmaceuticals Ltd	
Other	Cerebrovascular Accident		Diamox			
	Convulsion		(Acetazolamide)	SS		
	Loss Of Consciousness		Depakote (Valproate			

Tremor
Visual Disturbance

Semisodium) SS
(Loratadine) C
(Fluticasone
Propionate) C
(Montelukast) C

Date:11/13/00ISR Number: 3610819-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000977
Age: Gender:Male I/FU:I

Outcome PT
Hospitalization - Amnesia
Initial or Prolonged Balance Disorder
Disturbance In Attention
Dizziness
Fall

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nervousness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1) 4800 MG (1200 MG, QID), PER ORAL; 2) UNK, UNKNOWN		Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
			(Prednisone)	C		
			(Calcium)	C		

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 Confusional State Hypotension	Consumer	Tenormin	PS	Astrazeneca Pharmaceuticals Ip	ORAL
150 MG BID PO		Pneumonia Sedation Speech Disorder Toxicologic Test Abnormal		Zyban	SS		
				Nifedipine	SS		
				Gabapentin	SS		

Date:11/13/00ISR Number: 3610971-9Report Type:Expedited (15-DaCompany Report #002-0945-M0000035
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
1200 MG (400 MG, TID)				Dilantin (Phenytoin)			

Sodium) SS ORAL
 100 MG EVERY
 SECOND DAY
 PER ORAL

Date:11/13/00ISR Number: 3610972-0Report Type:Expedited (15-DaCompany Report #033-0945-M0000088
 Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Bundle Branch Block Drug Interaction	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
PER ORAL		Dyspnoea Fatigue	Professional	Flecaine (Flecainide Acetate)	SS		ORAL
		Sinus Bradycardia		Depakine (Valproate Sodium)	C		

Date:11/13/00ISR Number: 3610973-2Report Type:Expedited (15-DaCompany Report #002-0945-M0000035
 Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
1200 MG (400 MG, TID)			Professional				
				Dilantin (Phenytoin Sodium)	SS		ORAL

100 MG EVERY
 SECOND DAY,
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

PER ORAL

Date:11/13/00ISR Number: 3611141-0Report Type:Expedited (15-DaCompany Report #001-0945-M0001165
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (300 MG, QID)		Cardiac Arrest Dysentery Pneumonia	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Klonopin (Clonazepam)	C		
				Desyrel Trazodone Hydrochloride)	C		
				Lithium	C		
				Prochlorperazine	C		
				Folic Acid	C		
				Inderal (Propranolol Hydrochloride)	C		
				Methadone	C		
				Duragesic Patch (Fentanyl)	C		

Date:11/13/00ISR Number: 3611143-4Report Type:Expedited (15-DaCompany Report #001-0945-M0001145
 Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 12000 MG (3000 MG, QID), PER ORAL		Status Epilepticus	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other (DAILY), PER	Cerebrovascular Accident Coma Convulsion Difficulty In Walking	Consumer	Neurontin (Gabapentin)	PS	Parke Davis Pharmaceuticals Ltd	ORAL
ORAL	Dizziness Fall Loss Of Consciousness Paralysis Speech Disorder		Cognex (Tacrine Hydrochloride0 Detrol (Tolterodine Tartrate) Selenium Vitamin E	C C C C C C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged DAILY PO DAILY PO	Intentional Misuse	Health Professional	Luvox Neurontin Zyprexa Tylenol	PS SS C C	Solvay Pharmaceuticals	ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/14/00ISR Number: 3611688-7Report Type:Expedited (15-DaCompany Report #033-0945-M000088
Age:76 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNK, PER ORAL		Bundle Branch Block Left Drug Interaction	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
UNK, PER ORAL		Dyspnoea Fatigue	Professional	Flecaine (Flecainide Acetate)	SS		ORAL
		Sinus Bradycardia		Depakine (Valproate Sodium)	C		

Date:11/14/00ISR Number: 3611783-2Report Type:Expedited (15-DaCompany Report #044-0945-M0000136
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200 MG (DAILY), PER ORAL (SEE IMAGE)		Aphasia Hypertonia	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
		Masked Facies	Professional				
		Paralysis					
		Parkinson'S Disease					
		Tremor		Thyroxine	C		

Date:11/14/00ISR Number: 3612066-7Report Type:Expedited (15-DaCompany Report #A036504
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Required 300.00 MG Intervention to TOTAL TID		Confusional State Dissociative Disorder	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
		Dysarthria		Gabapentin	SS		ORAL
		Headache					

Prevent Permanent
ORAL
Impairment/Damage

Insomnia
Memory Impairment
Mental Disorder
Paranoia
Speech Disorder

Lithium C
Hydrocodone +
Acetaminophen C
Estrogen C
Ergotamine +
Caffeine C

Date:11/14/00ISR Number: 3612196-XReport Type:Expedited (15-DaCompany Report #001-0945-M0001210
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Blood Electrolytes Abnormal	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:11/15/00ISR Number: 3612105-3Report Type:Expedited (15-DaCompany Report #001-0902-M0000076
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 90-120 MG (DAILY), PER ORAL	Blood Pressure Decreased Hiv Infection Cdc Group I Insomnia Nephrolithiasis Sexual Dysfunction Visual Disturbance	Consumer	Nardil Neurontin (Gabapentin) Viracept (Nelfinavir Mesilate) Epivir (Lamivudine)	PS SS C C	Parke Davis Div Warner Lambert Co	ORAL

Freedom Of Information (FOI) Report

Zerit (Stavudine) C
 Lipitor
 (Atrovastatin) C

Date:11/15/00ISR Number: 3612446-XReport Type:Expedited (15-DaCompany Report #044-0945-M0000078
 Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG		Blood Albumin Increased Dehydration	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
(DAILY), PER		Dizziness	Professional				
ORAL		Nephrotic Syndrome					
		Oedema Peripheral Orthostatic Hypotension Proteinuria		(Amitriptyline) Tegretol Retard (Carbamazepine) Frusemide (Furosemide) (Simvastatin)	C C C C		

Date:11/15/00ISR Number: 3612748-7Report Type:Expedited (15-DaCompany Report #001-0945-M0001107
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (300		Drug Dependence Feeling Abnormal	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

MG, TID), PER

ORAL

Date:11/15/00ISR Number: 3612749-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000966
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged 800 MG (DAILY); 1600 MG (DAILY); 3600 MG (DAILY); 4800	Blindness Colour Blindness Optic Disc Haemorrhage Optic Neuritis Photopsia Vision Blurred Visual Field Defect	Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd
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Date:11/15/00ISR Number: 3612842-0Report Type:Expedited (15-DaCompany Report #001-0945-M0001191
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 4800 MG (DAILY)		Drug Abuser Euphoric Mood Intentional Misuse	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:11/16/00ISR Number: 3611875-8Report Type:Expedited (15-DaCompany Report #A0122366A
Age:70 YR Gender:Female I/FU:F

Outcome Hospitalization - Initial or Prolonged	PT Abdominal Pain Abdominal Pain Upper Concussion Erythema
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fall					
		Feeling Abnormal					
		Gastric Mucosal Lesion					
		Hand Fracture		Dilantin	PS		
		Head Injury		Neurontin	SS		
		Overdose		Depakote	SS		
200MG SEE		Spinal Fracture		Lamictal	SS	Glaxo Wellcome	ORAL
TEXT							
				Lamictal	SS	Glaxo Wellcome	ORAL
100MG SEE							
TEXT							

Date:11/16/00ISR Number: 3612811-0Report Type:Expedited (15-DaCompany Report #001-0981-992632
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Amnesia	Consumer	Lipitor	PS	Pfizer Ireland	
Initial or Prolonged		Brain Damage	Health			Pharmaceuticals,	
Other		Cerebrovascular Accident	Professional			Tablet Plant	ORAL
10 MG							
(DAILY), PER		Condition Aggravated					
ORAL		Diabetic Neuropathy					
		Hypertension		Neurontin			
		Pain In Extremity		(Gabapentin)	SS		
		Peripheral Vascular		Glucophage	C		
		Disorder		Rezulin	C		
				Insulin	C		

Date:11/16/00ISR Number: 3613034-1Report Type:Expedited (15-DaCompany Report #049-0945-M0000019
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatitis	Foreign	Neurontin	PS	Parke Davis	
			Health			Pharmaceuticals Ltd	
			Professional				

Date:11/17/00ISR Number: 3613404-1Report Type:Expedited (15-DaCompany Report #00P-163-0099888-00 (0)
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 250 MG, 3 IN Initial or Prolonged 1 D, PER ORAL		Asthma	Consumer	Depakote	PS	Abbott Laboratories	ORAL
		Tremor	Other				
		Visual Disturbance		Gabapentin (Gabapentin)	SS		ORAL
300 MG, 1 IN 1 D, PER ORAL				Oxycodone	C		
				Sonata	C		
				Tizanidine	C		
				Tylox	C		
				Singulair	C		
				Nometasone Furoate	C		
				Fluticasone Propionate	C		
				Salbutamol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/00ISR Number: 3615207-0Report Type:Expedited (15-DaCompany Report #001-0945-M0001218
 Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG, DAILY), PER ORAL		Chills Dizziness Fall Feeling Abnormal	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
7.5 MG/750 MG (UNK), PER ORAL		Foot Fracture Influenza Like Illness Lower Limb Fracture		(Hydrocodone)	SS		ORAL
		Nausea Vomiting		Fosamax Norvasc Calcium	C C C		

Date:11/20/00ISR Number: 3615275-6Report Type:Expedited (15-DaCompany Report #A0122366A
 Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG/SEE Initial or Prolonged TEXT/ORAL		Abdominal Pain Upper Back Injury Convulsion	Consumer	Lamictal	PS	Glaxo Wellcome Inc	ORAL
100 MG 100 MG		Endoscopy		Phenytoin	SS		
200 MG/SEE TEXT/ORAL		Endoscopy Abnormal Fall Feeling Abnormal		Semisodium Valproate Lamictal	SS SS		ORAL
		Hand Fracture Nervous System Disorder					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion	Foreign Health Professional	Neurontin Meronem (Meropenem)	PS SS	Parke Davis Pharmaceuticals Ltd	
INTRAVENOUS	1500 MG						

(DAILY)

INTRAVENOUS

- Levolac (Lactulose) C
- Zantac (Ranitidine Hydrochloride) C
- Pulmicort (Budenonide) C
- Deprakine (Valproate Sodium) C
- Sabrillex (Vigabatrin) C
- Propulsin (Cisapride Monohydrate) C
- Orfiril (Valproate Sodium) C
- Salbuvent (Salbutamol Sulfate) C
- Oradexon (Dexamethasone) C
- Rifampicin C
- Topimax (Topiramate) C
- Hydantin (Phenytoin) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/00ISR Number: 3631160-8Report Type:Periodic
Age:73 YR Gender:Male I/FU:I

Company Report #2000026966US

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Tremor	Health Professional	Zyvox	PS	Pharmacia And Upjohn Co	
				Neurontin	SS		

Date:11/21/00ISR Number: 3615603-1Report Type:Expedited (15-DaCompany Report #001-0945-M0001057
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accidental Overdose	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Other		Creatinine Renal					
2700 MG		Clearance Decreased	Other				
(DAILY)		Renal Impairment					

Date:11/21/00ISR Number: 3615604-3Report Type:Expedited (15-DaCompany Report #001-0945-M0001206
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Upper Blood Electrolytes	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1800 MG (600		Decreased					
MG TID) PER		Disorientation					
ORAL		Drug Interaction		(Methadone)	SS		ORAL
PER ORAL		Drug Level Above		Morphine	SS		ORAL
PER ORAL		Therapeutic		Baclofen	SS		ORAL
PER ORAL		Hypertension		Zoloft (Sertraline Hydrochloride)	C		
		Memory Impairment		Clonidine	C		
		Migraine		Cardizem (Diltiazem			
		Muscle Contractions					

Involuntary
Tachycardia
Tremor

Hydrochloride) C
(Potassium) C
Theophylline C
Hydrochlorothiazide C
Flovent (Fluticasone
Propionate) C
(Fluocinonide) C
Combivent
(Ipratropium
Bromide, Salbutamol
Sulfate) C
Teagen
(Benzalkonium
Chloride) C
Vitamin B12
(Cyanocobalamin) C
Vasocon
(Phenylmercuric
Acetate, Sodium
Carbonate Anhydrous,
Naphazoline C
Docusate C
Compazine
(Prochlorperazine
Edisylate) C
(Capsaicin) C
Herbal Natural
Estrogen C

Freedom Of Information (FOI) Report

Vitamin E
 (Tocopherol) C
 Coenzyme Q10
 (Ubidecarenone) C
 Stresstab(Vitamins
 Nos) C
 (Garlic) C
 Very Green
 Supplement C
 Echinacea Extract C
 Vicodin
 (Paracetamol,
 Hydrocodone
 Bitartrate) C
 Bacitracin
 /Polymixin Ointment) C
 Benadryl
 (Diphenhydramine
 Hydrochloride) C
 Ibuprofen C
 Nystatin Cream C

Date:11/21/00ISR Number: 3615857-1Report Type:Expedited (15-DaCompany Report #00P-056-0099941-00 (0)
 Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	500 MG, 1 IN	Bradycardia Bundle Branch Block Left Drug Interaction	Foreign Health Professional	Depakene	PS	Abbott Laboratories Pharmaceutical Products Div	ORAL
	1 D, PER ORAL	Dyspnoea					
	PER ORAL	Fatigue		Flecainide Acetate (Flecainide Acetate)	SS		ORAL
	PER ORAL			Gabapentin (Gabapentin)	SS		ORAL

Date:11/21/00ISR Number: 3615920-5Report Type:Expedited (15-DaCompany Report #033-0945-M0000081
 Age:76 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Brain Oedema	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Other		Sepsis					
300 MG		Thrombocytopenia	Professional				
(DAILY),				Unspecified Corticoids	SS		
				Medrol	C		
				Mopral	C		
				Cordarone	C		

Date:11/21/00ISR Number: 3615921-7Report Type:Expedited (15-DaCompany Report #044-0945-M0000168
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Raynaud'S Phenomenon	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Other							
900 MG (300			Professional				
MG, TID), PER							
ORAL				Warfarin	C		
				Digoxin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Paracetamol C

Date:11/21/00ISR Number: 3616938-9Report Type:Expedited (15-DaCompany Report #001-0945-M001206
 Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG (600 MG, TID) PER ORAL	Abdominal Pain Upper Blood Electrolytes Decreased Disorientation	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
PER ORAL	Drug Level Above		Methadone	SS		ORAL
PER ORAL	Therapeutic		Morphine	SS		ORAL
PER ORAL	Dysgraphia		Baclofen	SS		ORAL
	Hypertension		Zoloft	C		
	Migraine		Clonidine	C		
	Movement Disorder		Cardizem	C		
	Pain		Potassium	C		
	Tachycardia		Theophylline	C		
	Tremor		Hydrochlorothiazide	C		
			Flovent	C		
			Fluocinonide	C		
			Combivent	C		
			Teargen	C		
			Vitamin B12	C		
			Vasocon	C		
			Docusate	C		
			Compazine	C		
			Capsaicin	C		
			Herbal Natural			
			Estrogen	C		
			Vitamin E	C		
			Coenzyme Q10	C		
			Stresstab	C		
			Garlic	C		
			Very Green			
			Supplement	C		
			Echinacea Extract	C		
			Vicodin	C		

Bacitracin/Polymixin C
 Ointment C
 Benadryl C
 Ibuprofen C
 Nystatin Cream C

Date:11/22/00ISR Number: 3616186-2Report Type:Periodic Company Report #98811.01
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Ineffective Drug Interaction	Consumer	Lorazepam	PS	Mylan Pharmaceuticals Inc	ORAL
1 MG, BID, ORAL				Neurontin Capsules 300 Mg Parke-Davis	SS	Parke-Davis	ORAL
600 MG QAM & QHS, 400 MG TID, ORAL				Rochephin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/22/00ISR Number: 3616394-0Report Type:Expedited (15-DaCompany Report #044-0945-M0000168

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	900 MG (300 MG, TID), PER ORAL	Raynaud'S Phenomenon	Foreign Health Professional Other	Neurontin (Warfarin) (Digoxin) (Paracetamol)	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:11/22/00ISR Number: 3616395-2Report Type:Expedited (15-DaCompany Report #033-0945-M0000081

Age:76 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300 MG (DAILY) CEREBRAL EDEMA	Brain Oedema Sepsis Thrombocytopenia	Foreign Health Professional Other	Neurontin Corticoids Medrol (Methylprednisolone) Mopral (Omeprazole) Cordarone (Amiodarone Hydrochloride)	PS	Parke Davis Pharmaceuticals Ltd	

Date:11/22/00ISR Number: 3617771-4Report Type:Expedited (15-DaCompany Report #A036897

Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Amnesia	Health	Procardia Xl	PS	Pfizer Laboratories	

Initial or Prolonged	Confusional State	Professional		Div Pfizer Inc	ORAL
ORAL					
	Hypotension		Bupropion	SS	ORAL
300.00 MG					
	Pneumonia				
TOTAL: BID: ORA					
L					
	Sedation				
	Speech Disorder		Gabapentin	SS	
	Toxicologic Test Abnormal		Atenolol	SS	

Date: 11/24/00 ISR Number: 3616346-0 Report Type: Expedited (15-DaCompany Report #249263
 Age: 55 YR Gender: Female I/FU: I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Breast Cancer Female	Health	Accutane	PS	Hlr Technology	ORAL
20 MG DAILY			Professional				
ORAL				Neurontin (Gabapentin)	SS		ORAL
100 MG DAILY							
ORAL							

Date: 11/24/00 ISR Number: 3617228-0 Report Type: Expedited (15-DaCompany Report #033-0945-M0000093
 Age: Gender: Male I/FU: I

Outcome	PT
Hospitalization -	Anaemia
Initial or Prolonged	Leukopenia
	Pancytopenia
	Pulmonary Oedema

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Thrombocytopenia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
400 MG (200 MG, BID),		Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
			Triatec (Ramipril)	C		
			Kardegic (Acetylsalicylate Lysine)	C		
			Trivastal (Piribedil)	C		
			Sermion (Nicergoline)	C		

Date:11/24/00ISR Number: 3617229-2Report Type:Expedited (15-DaCompany Report #358-0945-M0000006

Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	600 MG (DAILY), PER ORAL	Blood Disorder Clonic Convulsion Condition Aggravated Muscle Twitching	Foreign Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Marevan (Warfarin Sodium)	C		
				Tramal Retard (Tramadol Hydrochloride)	C		
				Panodil (Paracetamol)	C		
				Lanzo (Lansoprazole)	C		
				Spesicor (Metoprolol Tartrate)	C		
				Cozaar (Losartan Potassium)	C		
				Hydrex Semi (Hydrochlorothizide)	C		

Furesis Special (Furosemide)	C
Laxoberon (Sodium Picosulfate)	C
Oxepam (Oxazepam)	C
Insulin Mixtard (Insulin Injection, Isophane)	C
Zomax (Zomepirac Sodium)	C

Date:11/24/00ISR Number: 3617402-3Report Type:Expedited (15-DaCompany Report #047-0945-M0000004
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2400 MG	Blindness Unilateral Optic Neuritis	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Other (DAILY), PER			Professional				
ORAL				Rivotril (Clonazepam)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Several Unspecified Medications C

Date:11/27/00ISR Number: 3616029-7Report Type:Expedited (15-DaCompany Report #249263
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Breast Cancer Female		Neurontin	PS		
				Accutane Capsules	SS	Roche	

Date:11/29/00ISR Number: 3619050-8Report Type:Expedited (15-DaCompany Report #JRFUSA2000009351
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Aspiration	Health	Duragesic	PS	Alza Corp	
TRANSDERMAL	75 MCG/H, 1	Medication Error	Professional				
IN 1 TIME(S),							
TRANSD							
ORAL				Percocet (Oxycodone)	SS		ORAL
ORAL				Seroquel (Seroquel)	SS		ORAL
ORAL				Revia (Naltrexone)	SS		ORAL
ORAL				Neurontin (Gabapentin)	SS		ORAL
ORAL				Trazodone (Trazodone)	SS		ORAL
ORAL				Depakote (Valproate Semisodium)	SS		ORAL
ORAL				Amitriptyline (Amitriptyline0	SS		ORAL
ORAL				Celexa (Citalopram Hydrobromie)	SS		ORAL

Anti-Depressants
(Nos)
Antidepressants) SS
Fentanyl (0,05 Mg/Ml
Injection)
(Fentanyl) SS

INTRAVENOUS 1 IN 1

TIME(S), IV

Date:11/29/00ISR Number: 3619079-XReport Type:Expedited (15-DaCompany Report #001-0945-M0001245
Age:48 YR Gender:Male I/FU:I

Outcome	PT
Life-Threatening	Abdominal Pain
Hospitalization -	Abnormal Dreams
Initial or Prolonged	Aggression
Disability	Angina Pectoris
Other	Anxiety
	Arachnoiditis
	Arrhythmia
	Asthenia
	Back Pain
	Bronchitis
	Cardiac Disorder
	Cardiomegaly
	Cervical Spinal Stenosis
	Chest Pain
	Condition Aggravated
	Constipation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Decreased Appetite Depression Diarrhoea				
1200 MG		Dizziness Drug Abuser	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
(DAILY),		Dry Mouth				
UNKNOWN		Dyspnoea				
30 MG		Electrocardiogram Qt Prolonged	Propulsid (Cisapride)	SS		ORAL
(DAILY), PER		Gastric Ulcer				
ORAL, 40 MG		Hallucination				
(DAILY), PER		Hyperhidrosis				
ORAL		Hypotension				
75 MG		Influenza Like Illness Insomnia Mental Disorder	Pamelor (Nortriptyline Hydrochloride)	SS		ORAL
(DAILY), PER		Mitral Valve Incompetence				
ORAL		Muscle Spasms				
200 MG		Myalgia	(Promethazine)	SS		ORAL
(DAILY), PER		Myocardial Ischaemia				
ORAL		Neck Pain				
1000 MG		Neuropathy Peripheral Pancreatic Carcinoma Pancreatitis Pharyngitis Polydipsia Polyp Polyuria Pruritus	Mellaril (Thioridazine Hydrochloride) Compazine (Prochlorperazine Edisylate) Biaxin (Clarithromycin)	SS SS SS		
(DAILY)		Rash Papular				

Rectal Haemorrhage	Elavil	
Renal Colic	(Amitriptyline	
Rhinitis	Hydrochloride)	SS
Road Traffic Accident	Prilosec	
Suicidal Ideation	(Omeprazole)	C
Suicide Attempt	Paxil (Paroxetine	
Syncope	Hydrochloride)	C
Tachycardia	Prozac (Fluoxetine	
Tricuspid Valve	Hydrochloride	C
Incompetence	Vicodin	
Ultrasound Scan Abnormal	(Paracetamol,	
Urinary Retention	Hydrocodone	
Urinary Tract Infection	Bitartrate)	C
Ventricular Extrasystoles	Procardia	
Ventricular Fibrillation	(Nifedipine)	C
Vision Blurred	Valium (Diazepam)	C
Vomiting	Prevacid	
Weight Decreased	(Lansoprazole)	C
	Reglan	
	(Metoclopramide)	C
	(Baclofen)	C
	Vistaril	
	(Hydroxyzine	
	Embonate)	C
	Vitamins (Vitamins	
	Nos)	C
	Nitroglycerin	
	(Phenobarbital,	
	Atropine	
	Methonitrate,	
	Glyceyl Trinitrate,	C

Freedom Of Information (FOI) Report

Darvocet (Paracetamol, Dextropropoxyphene)	C
Parafon Forte (Chlorzoxazone, Paracetamol)	C
Flexeril (Cyclobenzaprine Hydrochloride)	C
Isocet (Caffeine, Butalbital, Paracetamol)	C
(Trazodone)	C
Amoxil (Amoxicillin)	C
(Metoprolol)	C
(Carbamazepine)	C
Talacen (Paracetamol, Pentazocine Hydrochloride)	C
Niferex-150 Forte (Vitamins Nos, Minerals Nos)	C
Celexa (Citalopram Hydrobromide)	C
(Lorazepam)	C
Medrol Dose Pak (Methylprednisolone)	C
(Lactulose)	C
Atarax (Hydroxyzine Hydrochloride)	C
Limbitrol (Chlordiazepoxide, Amitriptyline Hydrochloride)	C
Remeron (Mirtazapine)	C
Ms Contin (Morphine Sulfate)	C
Oxycontin (Oxycodone Hydrochloride)	C
Hemorrhoidal Suppository	
Unspecified	C
Midrin (Paracetamol, Dichoralphenazone, Isometheptene)	C

Zithromax
(Azithromycin) C
Viokase
(Pancrelipase) C

Date:11/30/00ISR Number: 3618967-8Report Type:Expedited (15-DaCompany Report #B0092629A

Age:65 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - RESPIRATORY Initial or Prolonged (INHALATION)	Anaemia Dyspnoea Exertional Pallor		Ventoline Asasantine Mopral	PS SS SS	Glaxo Wellcome	ORAL ORAL

2UNIT per day

10MG Per day

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

1200MG per day	Neurontin	SS	ORAL
	Bronchokod	C	
	Doliprane	C	
	Lasilix	C	
	Sodium Chlorure	C	
	B-Vitamins	C	

Date:11/30/00ISR Number: 3619921-2Report Type:Expedited (15-DaCompany Report #358-0945-M000005
Age:2 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Convulsion	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
INTRAVENOUS	1500 MG DAILY		MeroneM (Meropenem)	SS		
INTRAVENOUS			Levolac (Lactulose)	C		
			Zantac (Ranitidine Hydrochloride)	C		
			Pulmicort (Budesonide)	C		
			Deprakine (Valproate Sodium)	C		
			Sabrillex (Vigabatrin)	C		
			Propulsin (Cisapride Monohydrate)	C		
			Orfiril (Valproate Sodium)	C		
			Salbuvent (Salbutamol Sulfate)	C		
			Oradexon (Dexamethasone)	C		
			(Rifampicin)	C		
			Topimax (Topiramate)	C		
			Hydantin (Phenytoin)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Fall Muscle Twitching	Health Professional	Neurontin (Oxcarbazepine) Zoloft (Sertraline Hydrochloride) Ultram (Tramadol Hydrochloride) Elavil (Amitriptyline Hydrochloride) Clonidine (Clonidine) Lotrim (Clotrimazole) Vitamins (Vitamins Nos)	PS SS C C C C C C	Parke Davis Pharmaceuticals Ltd	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/00ISR Number: 3620128-3Report Type:Expedited (15-DaCompany Report #001-0945-M0001254

Age:30 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1) 200 MG (100 MG, BID); 2) 1500 MG (500MG, TID)	Back Injury Condition Aggravated Coordination Abnormal Dizziness Dysarthria Fall Syncope	Consumer	Neurontin Prozac (Fluoxetine Hydrochloride) Vioxx (Rofecoxib) (Diazepam) (Hydrochlorothiazide , Triamterene) Demerol (Pethidine Hydrochloride)	PS C C C C C C	Parke Davis Pharmaceuticals Ltd	

Date:11/30/00ISR Number: 3620130-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000462

Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER ORAL	Blood Glucose Increased Cellulitis Disturbance In Attention Dizziness Headache Nausea Speech Disorder Tremor Vision Blurred	Consumer	Neurontin (Insulin) Aleve (Naproxen Sodium)	PS C C	Parke Davis Pharmaceuticals Ltd	ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - RESPIRATORY	Anaemia	Foreign	Ventolin	PS	Glaxo Wellcome Inc	
Initial or Prolonged (INHALATION) 100 MCG	Dyspnoea Exertional					
INHALED	Pallor					
ORAL			Asasantin Capsule (Asasantin)	SS		ORAL
10 MG PER DAY			Omeprazole Capsule (Omeprazole)	SS		ORAL
ORAL						
400 MG ORAL			Gabapentin Capsule (Gabapentin)	SS		ORAL
			Carbocisteine	C		
			Paracetamol	C		
			Frusemide	C		
			Sodium Chloride	C		
			Compound Vitamin B	C		

Outcome
Life-Threatening
Hospitalization -

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Freedom Of Information (FOI) Report

Initial or Prolonged
Disability
Other

PT
Anhedonia
Anxiety
Arrhythmia
Asthenia
Back Pain
Bronchitis
Cardiac Disorder
Cardiomegaly
Chest Pain
Condition Aggravated
Cough
Dizziness
Drug Abuser
Dyspnoea
Electrocardiogram Qt
Corrected Interval
Prolonged
Electrocardiogram Qt
Prolonged
Emotional Disorder
Emotional Distress
Fall
Fear Of Disease
Fibrosis
Headache
Heart Rate Irregular
Hyperhidrosis
Hypoaesthesia
Injury
Joint Swelling
Left Ventricular Failure
Malaise
Mental Disorder
Mitral Valve Incompetence
Myalgia
Nasal Congestion
Nausea
Neck Pain
Nuclear Magnetic
Resonance Imaging
Abnormal
Pain
Pancreatic Carcinoma
Pancreatitis

Pharyngitis
Pharyngolaryngeal Pain
Polydipsia
Polyuria
Pyrexia
Rash Erythematous
Renal Colic
Rhinitis
Road Traffic Accident
Sinus Tachycardia
Spinal Osteoarthritis
Suicide Attempt
Syncope
Tricuspid Valve

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Incompetence Urinary Retention Urinary Tract Infection Ventricular Fibrillation Vomiting	Report Source	Product	Role	Manufacturer	Route
1200 MG				Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
(DAILY)							
30 MG				Propulsid	SS		ORAL
(DAILY), PER							
ORAL (SEE							
IMAGE)							
200 MG				Promethazine	SS		ORAL
(DAILY), PER							
ORAL							
75 MG				Pamelor	SS		ORAL
(DAILY), PER							
ORAL							
1000 MG				Mellaril	SS		
(DAILY)				Biaxin	SS		
				Elavil	SS		
				Compazine	SS		
				Prilosec	C		
				Trazodone	C		
				Isocet	C		
				Flexeril	C		
				Parafon Forte	C		
				Darvocet	C		
				Nitroglycerin	C		
				Vitamins	C		
				Vistaril	C		
				Baclofen	C		
				Zithromax	C		

Midrin	C
Hemorrhoidal	
Suppository	
Unspecified	C
Oxycontin	C
Ms Contin	C
Remeron	C
Limbitrol	C
Atarax	C
Lactulose	C
Viokase	C
Medrol Dose Pak	C
Lorazepam	C
Celexa	C
Niferex-150 Forte	C
Talacen	C
Carbamazepine	C
Metoprolol	C
Amoxil	C
Vicodin	C
Procardia	C
Valium	C
Reglan	C
Prevacid	C
Paxil	C
Prozac	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/05/00ISR Number: 3622411-4Report Type:Direct
Age:83 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Coma		Gabapentin	PS		ORAL
Other		Dialysis					
200MG PO		Mental Impairment					
AFTER							
DIALYSIS							
				Amiodarone	C		
				Calcium Carbonate	C		
				Docusate	C		
				Erythropoietin	C		
				Folic Acid	C		
				Metoprolol	C		
				Tamsulosin	C		
				Lactulose	C		
				Promethazine	C		
				Carbamazepine	C		

Date:12/05/00ISR Number: 3622509-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000301
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Neurontin	PS	Parke Davis	
Hospitalization -		Amblyopia				Pharmaceutical	
Initial or Prolonged		Amnesia				Research Div Warner	
		Back Pain				Lambert Co	ORAL
1800 MG (600		Confusional State					
MG,TID), PER		Coordination Abnormal					
ORAL / 2400		Depression					
MG (600 MG		Dizziness					
BID, 1200 MG		Drug Withdrawal Syndrome		Pain Medication			
		Dry Mouth		Unspecified	C		
		Dry Throat		Celexa(Citalopram			
		Dysarthria		Hydrobromide)	C		
		Euphoric Mood					

Fatigue
 Headache
 Hostility
 Muscle Twitching
 Oedema
 Thinking Abnormal
 Tremor
 Vomiting
 Weight Decreased

Date:12/05/00ISR Number: 3622510-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000403
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Pulmonary Fibrosis Respiratory Distress	Health Professional	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
4000 MG (800 MG TID, 1600 MG QHS), PER ORAL				Zoloft (Sertraline Hydrochloride) Depakote (Valproate Semisodium)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/05/00ISR Number: 3622666-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000264
 Age:71 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Confusional State Disorientation Renal Cell Carcinoma Stage Unspecified	Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
1800 MG (600 MG, TID), PER ORAL						

Zestril (Lisinopril)	C
Levoxyl (Levothyroxine Sodium)	C
Celebrex (Celecoxib)	C
Baby Aspirin (Acetylsalicylic Acid)	C

Date:12/06/00ISR Number: 3623444-4Report Type:Expedited (15-DaCompany Report #042-0945-M0000001
 Age:31 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Coordination Abnormal Headache Nystagmus Vertigo	Foreign Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
900 MG (DAILY), PER ORAL						

Sanepil (Phenobarbital, Phenytoin)	SS
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Date:12/06/00ISR Number: 3623667-4Report Type:Expedited (15-DaCompany Report #055-0945-M0000020
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Visual Acuity Reduced	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1200 MG (600 MG, BID), PER ORAL							

Date:12/07/00ISR Number: 3623654-6Report Type:Direct Company Report #
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Coordination Abnormal Disorientation		Neurontin 300mg Parke-Davis	PS	Parke-Davis	ORAL
NEURONTIN TID ORAL Dizziness							
		Lethargy Medication Error		Fioricet Generic Geneva Labs	SS	Geneva Labs	ORAL
FIORICET Q 4-6 HRS ORAL Rizotriptan C							

Date:12/07/00ISR Number: 3638462-XReport Type:Periodic Company Report #2000-09-0259
 Age:53 YR Gender:Male I/FU:I

Outcome	PT
	Alanine Aminotransferase Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
570 MG QDX5D, ORAL		Blood Lactate Dehydrogenase Increased Haptoglobin Decreased	Health	Temodar	PS	Schering Corp	ORAL
900 MG ORAL		Hyperbilirubinaemia Reticulocytosis	Professional	Neurontin Tablets	SS		ORAL

Date:12/11/00 ISR Number: 3626585-0 Report Type:Expedited (15-DaCompany Report #001-0945-M0001297
Age:52 YR Gender:Male I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID), PER ORAL; 600MG(200MG,T ID), PER		Abdominal Pain Cholelithiasis Loss Of Consciousness Malaise Nausea	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:12/11/00 ISR Number: 3626586-2 Report Type:Expedited (15-DaCompany Report #001-0945-M0000937
Age: Gender:Female I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		Impaired Gastric Emptying	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
240 MG (80				Oxycontin (Oxycodone Hydrochloride)	SS		

MG, TID)

YR

(Amitriptyline)	C
Reglan	
(Metoclopramide)	C
Klonopin	
(Clonazepam)	C
Prilosec	
(Omeprazole)	C
(Feros Sulfate)	C
Multivitamins	
(Ergocalciferol,	
Ascorbic Acid, Folic	
Acid, Thiamine	
Hydrochliride,	C

Date:12/11/00ISR Number: 3626929-XReport Type:Expedited (15-DaCompany Report #044-0945-M0000180

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vasculitis	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
100 MG			Professional				
(DAILY), PER							
ORAL							

(Amlodipine)	C
(Zomitriptan)	C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/11/00ISR Number: 3626930-6Report Type:Expedited (15-DaCompany Report #033-0945-M0000101
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1600 MG (DAILY), PER ORAL		Dyskinesia	Foreign Health Professional Other	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:12/11/00ISR Number: 3626931-8Report Type:Expedited (15-DaCompany Report #032-0945-M0000027
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL Other		Agranulocytosis	Foreign Health Professional Other	Neurontin (Valproate Sodium)	PS SS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:12/14/00ISR Number: 3630113-3Report Type:Expedited (15-DaCompany Report #001-0945-M0001324
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Bradycardia Hypothermia Pneumonia	Health Professional	Neurontin Olanzapine Quazepam	PS C C	Parke Davis Pharmaceuticals Ltd	

Date:12/14/00ISR Number: 3630345-4Report Type:Expedited (15-DaCompany Report #002-0945-M0000043
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Tunnel Vision	Foreign	Neurontin	PS	Parke Davis	

1200 MG
 (DAILY), PER
 ORAL

Visual Field Defect
 Health
 Professional
 Pharmaceuticals Ltd ORAL

Date:12/18/00ISR Number: 3632818-7Report Type:Expedited (15-DaCompany Report #001-0945-M0001218
 Age:68 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Chills Dizziness	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
900 MG (300 MG, DAILY), PER ORAL		Fall Feeling Abnormal					
7.5 MG/750 MG, PER ORAL		Influenza Like Illness Lower Limb Fracture		Hydrocodone	SS		ORAL
		Nausea Vomiting		Fosamax (Alendronate Sodium) Norvasc (Amlodipine Besilate) Calcium	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/18/00ISR Number: 3633147-8Report Type:Expedited (15-DaCompany Report #000928-SK359

Age: Gender:Not SpecifiI/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - PO	Drug Interaction	Health	Celebrex	PS	Gd Searle And Co	ORAL
Initial or Prolonged 300.00 MG BID	Mental Impairment	Professional	Neurontin	SS		ORAL

PO

Date:12/18/00ISR Number: 3633149-1Report Type:Expedited (15-DaCompany Report #000928-SK360

Age: Gender:Not SpecifiI/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - PO	Drug Interaction	Health	Celebrex	PS	Gd Searle And Co	ORAL
Initial or Prolonged 300.000 MG	Mental Impairment	Professional	Neurontin	SS		ORAL

BID PO

Date:12/18/00ISR Number: 3633153-3Report Type:Expedited (15-DaCompany Report #000928-SK361

Age: Gender:Not SpecifiI/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - PO	Drug Interaction	Health	Celebrex	PS	Gd Searle And Co	ORAL
Initial or Prolonged 300.00 MG BID	Mental Impairment	Professional	Neurontin	SS		ORAL

Renal Impairment

PO

Date:12/18/00ISR Number: 3633172-7Report Type:Expedited (15-DaCompany Report #000928-SK358

Age: Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Hospitalization - Drug Interaction Health Celebrex PS Gd Searle And Co ORAL
UNKNOWN PO
Initial or Prolonged Mental Impairment Professional Neurontin SS ORAL
300.000 MG
Renal Impairment
BID PO

Date:12/18/00ISR Number: 3633301-5Report Type:Expedited (15-DaCompany Report #002-0945-M0000044
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
PER ORAL			Professional				

Date:12/18/00ISR Number: 3633302-7Report Type:Expedited (15-DaCompany Report #002-0945-M0000031
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diplopia Myopia	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1600 MG (400 MG, QID), PER		Presbyopia	Professional				

ORAL

Date:12/18/00ISR Number: 3633381-7Report Type:Expedited (15-DaCompany Report #001-0945-M0001322
Age: Gender: I/FU:I

Outcome	PT
Congenital Anomaly	Complications Of Maternal Exposure To Therapeutic Drugs

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Cyst Foetal Disorder	Report Source	Product	Role	Manufacturer	Route
SEE TEXT, PLACENTAL			Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	OTHER
SEE TEXT, PLACENTAL				Tegretol (Carbamazepine)	SS		OTHER
				Folic Acid	C		

Date:12/18/00ISR Number: 3633384-2Report Type:Expedited (15-DaCompany Report #001-0945-M0001064
Age:48 YR Gender:Female I/FU:F

Outcome Dose Other	Duration	PT Colour Blindness Macular Oedema	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL			Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Demadex	C		
				Zoloft	C		
				Remeron	C		
				Premarin	C		
				Prevacid	C		
				Imitrex	C		

Date:12/19/00ISR Number: 3632885-0Report Type:Expedited (15-DaCompany Report #A0133661A
Age: Gender:Male I/FU:I

Outcome Dose Hospitalization - 50MG Per day	Duration	PT Convulsion	Report Source	Product	Role	Manufacturer	Route
				Lamictal	PS	Glaxo Wellcome	ORAL

Initial or Prolonged Coordination Abnormal Neurontin SS ORAL
YR
Disability Drug Interaction

Date:12/19/00 ISR Number: 3633682-2 Report Type:Direct Company Report #
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 400 MG PO	Coordination Abnormal	Health	Neurontin	PS		ORAL
Initial or Prolonged 1 MG PO	Tremor	Professional	Lamotrigine	SS		ORAL
			Folate	C		

Date:12/19/00 ISR Number: 3633778-5 Report Type:Expedited (15-DaCompany Report #001-0945-M0001323
Age:31 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required Intervention to 300 MG Prevent Permanent (DAILY) PER Impairment/Damage ORAL	Chest Pain	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/20/00ISR Number: 3633823-7Report Type:Expedited (15-DaCompany Report #A0122366A

Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100MG See Initial or Prolonged text		Abdominal Pain Upper		Lamictal	PS	Glaxo Wellcome	ORAL
		Concussion					
		Constipation		Dilantin	SS		
		Decreased Appetite		Neurontin	SS		
		Drug Ineffective		Depakote	SS		
200MG See text		Facial Bones Fracture		Lamictal	SS	Glaxo Wellcome	ORAL
		Fall					
		Hand Fracture					
		Insomnia					
		Nausea					
		Overdose					
		Spinal Fracture					

Date:12/20/00ISR Number: 3635081-6Report Type:Expedited (15-DaCompany Report #2000TRE0088

Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aspiration	Health Professional	Revia	PS	Dupont Merck Pharmaceutical Co	ORAL
PO				Percocet (Oxycodone /Acetaminophen)	SS		ORAL
PO				Duragesic(Fentanyl)	SS		
TRANSDERMAL	75 MCG/H ONCE						
TD				(Seroquel)	SS		ORAL
PO				Neurontin (Gabapentin)	SS		ORAL
PO				(Trazodone)	SS		ORAL
PO				(Antidepressant-Unsp			

			ecified)	SS	
			Depakote (Valproate Semisodium)	SS	ORAL
PO					
			Amitriptyline (Amitriptyline)	SS	ORAL
PO					
			Celexa (Citalopram Hydrobromide)	SS	ORAL
PO					
			(Fentanyl)	SS	
INTRAVENOUS	0.05 MG/ML				
INJECTION					
ONCE IV					

Date:12/20/00ISR Number: 3635094-4Report Type:Expedited (15-DaCompany Report #A0133661A
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 50 MG PER DAY		Coordination Abnormal	Health	Lamictal	PS	Glaxo Wellcome Inc	ORAL
Initial or Prolonged ORAL Disability		Drug Interaction	Professional				
		Epilepsy		Gabapentin (Formulation Unknown) (Gabapentin)	SS		ORAL
ORAL	YR						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/20/00ISR Number: 3635529-7Report Type:Expedited (15-DaCompany Report #2000031428US
Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Alopecia Deafness	Consumer	Depo-Medrol	PS	Pharmacia And Upjohn Co	
EPIDURAL	EPIDURAL	Madarosis Visual Acuity Reduced		Neurontin (Gabapentin) Cardura (Doxazosin Mesilate)	SS C		

Date:12/20/00ISR Number: 3635742-9Report Type:Expedited (15-DaCompany Report #001-0945-M0001228
Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Diarrhoea Lymphoma Oesophageal Pain Osteoporosis	Consumer	Neurontin Fosamax Unspecified Vitamins Lipitor	PS C C C	Parke Davis Pharmaceuticals Ltd	

Date:12/21/00ISR Number: 3635867-8Report Type:Expedited (15-DaCompany Report #A039777
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage		Difficulty In Walking Fall Pain	Consumer	Norvasc Neurontin Lipitor Coumadin Pain Medication	PS SS SS SS C	Pfizer Agricultural Div	

Date:12/22/00ISR Number: 3635929-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000616
Age:4 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Autism	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
300 MG (100 MG, TID), PER ORAL		Developmental Delay					
		Encephalopathy					
		Psychomotor Hyperactivity					
		Speech Disorder		Depakote (Valproate Semisodium)	SS		
		Vomiting		(Hydroxyzine Hydrochloride)	C		
				(Phenobarbital)	C		
				(Lactulose)	C		
				Senokot (Senna Fruit)	C		
				Ryna-12s (Chlorpheniramine, Pseudoephedrine Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/22/00ISR Number: 3637300-9Report Type:Expedited (15-DaCompany Report #033-0945-M0000104
Age:71 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG DAILY	Delirium Mania	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
PER ORAL			Laroxyl (Amitriptyline Hydrochloride)	C		
			Eucalcic (Calcium Carbonate)	C		
			Plavix (Clopidogrel)	C		
			Lasilix (Furosemide)	C		
			Tahor (Atorvastatin)	C		
			Tadenan (Pygeum Africanum)	C		
			Speciafoldine (Folic Acid)	C		
			Josir (Tamsulosin Hydrochloride)	C		

Date:12/26/00ISR Number: 3638427-8Report Type:Expedited (15-DaCompany Report #001-0945-M0001357
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Epistaxis Pulmonary Haemorrhage Respiratory Tract Haemorrhage	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:12/26/00ISR Number: 3638430-8Report Type:Expedited (15-DaCompany Report #001-0073-M0000496
Age:1 DY Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Complications Of Maternal Exposure To Therapeutic	Health Professional	Dilantin	PS	Parke Davis Div Warner Lambert Co	

SEE TEXT,

PLACENTAL
Drugs
Small For Dates Baby
Neurontin
(Gabapentin) SS
SEE TEXT,
PLACENTAL
Unspecified Nasal
Spray C
Unspecified Inhaler C
Folic Acid C
Ampicillin C
Terbutaline C

Date:12/26/00ISR Number: 3638773-8Report Type:Expedited (15-DaCompany Report #A0122366A
Age:70 YR Gender:Female I/FU:F

Outcome PT
Hospitalization - Concussion
Initial or Prolonged Drug Effect Decreased
Eating Disorder
Fall
Feeling Abnormal
Gastric Mucosal Lesion
Hand Fracture

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
100 MG SEE		Overdose Sleep Disorder Spinal Fracture	Consumer	Lamictal	PS	Glaxo Wellcome Inc	ORAL
TEXT ORAL				Phenytoin (Formulation Unknown)	SS		
100 MG				Gabapentin (Gabapentin_)	SS		
100 MG				Semisodium Valproate (Divalproex Sodium)	SS		
200 MG SEE				Lamictal Tablet (Lamotrigine)	SS		ORAL
TEXT ORAL							

Date:12/26/00ISR Number: 3640195-0Report Type:Expedited (15-DaCompany Report #358-0945-M0000007
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG		Amnesia	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
(DAILY), PER			Professional				
ORAL							

Date:12/26/00ISR Number: 3640196-2Report Type:Expedited (15-DaCompany Report #044-0945-M0000195
Age:1 DY Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Complications Of Maternal	Foreign	Neurontin	PS	Parke Davis	

PLACENTAL	Exposure To Therapeutic	Health		Pharmaceuticals Ltd
SEE TEXT,	Drugs	Professional	Phenytoin	
	Developmental Delay		(Phenytoin)	SS
	Visual Disturbance			
PLACENTAL			(Carbamazepine)	SS
SEE TEXT,				
PLACENTAL				

Date:12/26/00ISR Number: 3640197-4Report Type:Expedited (15-DaCompany Report #044-0945-M0000193
Age:90 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Dizziness	Foreign	Neurontin	PS	Parke Davis	
Initial or Prolonged	Fatigue	Health			Pharmaceuticals Ltd	ORAL
600 MG						
(DAILY), PER	Sleep Attacks	Professional				

ORAL

(Fluoxetine)	C
Aspirin	
(Acetylsalicylic	
Acid)	C

Date:12/26/00ISR Number: 3640361-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000595
Age: Gender:Male I/FU:I

Outcome
Hospitalization -
Initial or Prolonged
Other

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
1600 MG (800 MG, BID), PER ORAL	Angina Pectoris Chest Pain Coordination Abnormal Coronary Artery Embolism Coronary Artery Occlusion Deep Vein Thrombosis	Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
750 MG (250 MG, TID), PER ORAL	Difficulty In Walking Drug Level Above Therapeutic Memory Impairment		Depakote (Valproate Semisodium)	SS		ORAL
YEARS 30 MG (DAILY),	Myocardial Infarction Pneumonia		Coumadin (Warfarin Sodium) Remeron (Mirtazapine)	SS SS		
			Lipitor (Atorvastatin) Lasix (Furosemide) K-Dur (Potassium Chloride) Synthroid (Levothyroxine Sodium) Ambien (Zolpidem Tartrate) Plavix (Clopidogrel)	C C C C C C		

Date:01/02/01
 ISR Number: 3640315-8
 Report Type:Expedited (15-DaCompany Report #049-0945-M0000031
 Age:65 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Life-Threatening Other 1600 MG Required (DAILY) PER Intervention to ORAL Prevent Permanent Impairment/Damage	Arterial Disorder Hepatic Failure Hepatocellular Damage Pancreatitis Necrotising Pulmonary Oedema	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
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Date:01/02/01ISR Number: 3640317-1Report Type:Expedited (15-DaCompany Report #046-0945-M0000059
Age:24 YR Gender:Male I/FU:I

Outcome Dose Hospitalization - Initial or Prolonged 2400 MG DAILY	Duration	PT	Report Source	Product	Role	Manufacturer	Route
PER ORAL;3200 MG (DAILY), PER ORAL		Abdominal Abscess Hypersensitivity	Foreign Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Trileptal (Oxcarbazepine)	SS		
				Topimax (Topiramate)	SS		
				Ergenyl (Valproate Sodium)	SS		
				Pentasa (Mesasalazine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/02/01ISR Number: 3640912-XReport Type:Expedited (15-DaCompany Report #PHFR2000GB02178
 Age:2 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic	Foreign Health	Tegretol	PS	Novartis Pharmaceuticals Corp	
TRANSPLACENTAL	4 TABS	DAILY, Drugs	Professional				
TRANSPLACENTA		Developmental Delay	Other				
L		Vision Abnormal Neonatal		Phenytoin	SS		
TRANSPLACENTAL	2 TABS	DAILY,					
TRANSPLACENTA							
L				Gabapentin	SS		
TRANSPLACENTAL		TRANSPLACENTA					
L							

Date:01/02/01ISR Number: 3640986-6Report Type:Expedited (15-DaCompany Report #001-0945-M0001170
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cerebrovascular Accident Convulsion	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Other		Loss Of Consciousness Tremor Visual Disturbance		Diamox	SS		
				Depakote	SS		
				Loratadine	C		
				Fluticasone			
				Propionate	C		
				Montelukast	C		

Date:01/03/01ISR Number: 3641648-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000196
 Age:40 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Arthralgia Asthenia

Other

Blood Potassium Decreased
Carpal Tunnel Syndrome
Cholelithiasis
Cholestasis
Decreased Activity
Dental Caries
Depression
Difficulty In Walking
Dry Mouth
Fall
Fatigue
Gallbladder Disorder
Gallbladder Pain
Headache
Hypoaesthesia
Hypothyroidism
Joint Dislocation
Lethargy
Liver Function Test
Abnormal
Lymphadenopathy
Malnutrition
Movement Disorder
Nervous System Disorder
Oedema Peripheral
Osteoporosis
Ovarian Cyst

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pain In Extremity Pruritus Skin Discolouration	Report Source	Product	Role	Manufacturer	Route
900 MG(300 MG TID):2400MG(8 00MG TID):2100MG(7 00MG 80 MG PER ORAL 1600 MG 2500 MG 1 OR 2 (Q 4 H PRN) 80 MG 8 MG PER ORAL 200 MG PER ORAL 100 MG PER ORAL 2000 MG PER ORAL	1 WK	Skin Ulcer Tendon Disorder Vomiting Weight Decreased Weight Increased	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Baclofen	SS		ORAL
				Ms Contin	SS		
				Propulsid	SS		
				Methadone	SS		
				Percocet	SS		
				Valium	SS		
				Zanaflex	SS		ORAL
				Zoloft	SS		ORAL
				Hydrochlorothiazide	SS		ORAL
				Veetids	SS		ORAL
				Synthroid	SS		
				Oxy Ir (Oxycodone			

8-10 DAILY	Hydrochloride)	SS
80 MG	Lasix	SS
80 MG	Ritalin	SS
20 MCG	K-Dur	SS
	Seroquel	SS
	Ketamine	SS
	Klonopin	SS
	Corgard	SS
	Relafen	SS
	Celebrex	SS
800 MG		
4 MG	Carafate	SS
200 MG	Dextromethorphan	SS
	Nadolol	SS
	Tegaderm	SS

Date:01/04/01ISR Number: 3641829-7Report Type:Expedited (15-DaCompany Report #001-0945-M0001204
Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Chloasma	Consumer	Neurontin	PS	Parke Davis	
Initial or Prolonged	Ecchymosis				Pharmaceutical	
	Neuroma				Research Div Warner	
	Skin Discolouration				Lambert Co	ORAL

400 MG (100
MG, QID) PER

ORAL

Celebrex (Celecoxib)	C
Tetracycline	C
Fastin (Phentermine Hydrochloride)	C
Limbitrol (Chlordiazepoxide, Amitriptyline Hydrochloride)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/04/01ISR Number: 3642095-9Report Type:Periodic
Age:67 YR Gender:Female I/FU:I

Company Report #001-0945-M0000363

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Headache Increased Appetite	Consumer	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
1200 MG (600 MG, BID) PER ORAL				Premarin (Estrogens Conjugated) Depakote (Valproate Semisodium) Prinivil (Lisinopril) Centrum Silver (Ascorbic Acid)Tocoperyl Acetate, Retinol, Zinc, Calcium, Calcium	C C C C C		

Date:01/04/01ISR Number: 3649424-0Report Type:Expedited (15-DaCompany Report #001-0945-M0001074
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Angioneurotic Oedema Drug Interaction	Company Representative Other	Neurontin Celebrex (Celecoxib)	PS SS	Parke Davis Pharmaceuticals Ltd	ORAL
UNK, PER ORAL							

Date:01/05/01ISR Number: 3642314-9Report Type:Expedited (15-DaCompany Report #A0135985A
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other		Abdominal Pain		Lotronex	PS	Glaxo Wellcome	ORAL
1MG Twice per							
day	10	MON	Condition Aggravated				
			Diarrhoea	Neurontin	SS		
			Epilepsy	Trazodone	C		
			Migraine	Phenergan	C		
			Nightmare	Celebrex	C		
			Weight Decreased	Vicodin	C		
				Soma	C		
				Epidural	C		

Date:01/05/01ISR Number: 3642723-8Report Type:Expedited (15-DaCompany Report #001-0945-M0001381
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Postmenopausal	Health	Neurontin	PS	Parke Davis	
		Haemorrhage	Professional			Pharmaceuticals Ltd	
SEE IMAGE				Vioxx	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/05/01ISR Number: 3642728-7Report Type:Expedited (15-DaCompany Report #358-0945-M0000007

Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (DAILY), PER ORAL		Visual Pathway Disorder	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Seroxat (Paroxetine Hydrochloride) Triptyl (Amitripty-Line Hydrochloride)	C C		

Date:01/08/01ISR Number: 3643521-1Report Type:Expedited (15-DaCompany Report #A0135985A

Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1 MG/TWICE PER DAY, ORAL		Abdominal Distension Abdominal Pain	Consumer	Lotronex	PS	Glaxo Wellcome Inc	ORAL
		Condition Aggravated Constipation Diarrhoea Fear Grand Mal Convulsion Irritable Bowel Syndrome Migraine Nightmare Weight Decreased		Gabapentin Trazodone Promethazine Hcl Celecoxib Vicodin Carisoprolol Local Anesthetic	SS C C C C C C		

Date:01/09/01ISR Number: 3643569-7Report Type:Direct

Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Depressed Mood		Gabapentin (400 Mg)			

Initial or Prolonged Suicidal Ideation Caps, Parke-Davis) PS Parke-Davis ORAL
400 MG PO BID

Date:01/09/01ISR Number: 3643578-8Report Type:Direct Company Report #
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin	PS		
Other		Oedema Peripheral		Zyprexa	SS		

Date:01/10/01ISR Number: 3650510-XReport Type:Periodic Company Report #001102-SK642
Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Consumer	Celebrex	PS	Gd Searle And Co	ORAL
UNKNOWN PO		Back Pain		Neurontin	SS		ORAL
200.000 MG		Dysgeusia					
QHS PO							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/10/01ISR Number: 3652472-8Report Type:Periodic
Age:72 YR Gender:Female I/FU:I

Company Report #001026-SK879

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG QD PO		Constipation	Consumer	Celebrex	PS	Gd Searle And Co	ORAL
1800 MG PO		Dry Skin		Neurontin	SS		ORAL
TOPICAL	TOP	Dyspepsia		Cosmetics	SS		
MORB/MORT NEC		Paraesthesia		Cardiac Therapy	SS		
				Vitamins	C		
				Omeprazole	C		

Date:01/10/01ISR Number: 3652476-5Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001026-SK896

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG BID PO		Dyspepsia	Health	Celebrex	PS	Gd Searle And Co	ORAL
100 MG PO		Hypertension	Professional	Neurontin	SS		ORAL
				Tramadol			
				Hydrochloride	C		
				Hydrocodone			
				Bitartrate	C		
				Acetaminophen	C		
				Quinapril			
				Hydrochloride	C		
				Hydrochlorothiazide	C		
				Alendronate	C		

Date:01/11/01ISR Number: 3646470-8Report Type:Expedited (15-DaCompany Report #001-0945-M0001385
Age:50 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300		Balance Disorder Disturbance In Attention	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

MG TID) PER
 ORAL; 1200
 (400 TID) PER
 ORAL
 Dizziness
 Grand Mal Convulsion
 Nervousness
 Speech Disorder
 Tremor
 Premarin (Estrogens
 Conjugated) C
 Provera
 (Medroxyprogesterone
 Acetate) C

Date:01/11/01ISR Number: 3646508-8Report Type:Expedited (15-DaCompany Report #033-0945-M0100001
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (100		Condition Aggravated Hepatic Enzyme Increased	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
MG, QID), PER		Hepatitis B	Professional				
ORAL		Hepatitis C					
PER ORAL		Hiv Infection		Norvir (Ritonavir)	SS		ORAL
PER ORAL		Prothrombin Time		Zerit (Stavudine)	SS		ORAL
PER ORAL		Prolonged		Depamide (Valpromide)	SS		ORAL
0.75 MG				Hivid (Zalcitabine)	SS		ORAL
(DAILY), PER							
ORAL				Zoloft (Sertraline)			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG				Hydrochloride)	SS		ORAL
(DAILY), PER							
ORAL							
Date:01/11/01ISR Number: 3646509-XReport Type:Expedited (15-DaCompany Report #033-0945-M0100002							
Age:57 YR Gender:Male I/FU:I							
Hospitalization -	Initial or Prolonged	Agranulocytosis	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1500 MG		Multiple Myeloma					
Other		Pyrexia	Professional				
(DAILY), PER							
ORAL							
INTRA	VENOUS			Axepim (Cefepime Hydrochloride)	SS		
400 MG				Oflocet (Ofloxacin)	SS		ORAL
(DAILY), PER							
ORAL							
20 MG				Moprol (Omeprazole)	SS		ORAL
(DAILY), PER							
ORAL							
50 MG				Laroxyl (Amitriptyline Hydrochloride)	SS		ORAL
(DAILY), PER							
ORAL							
100 MG				Aldactone (Spironolactone)	SS		ORAL

(DAILY), PER

ORAL

Date:01/12/01ISR Number: 3647192-XReport Type:Expedited (15-DaCompany Report #2001COU0016
Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Drug Effect Decreased	Consumer	Coumadin	PS	Dupont Merck Pharmaceutical Co	ORAL
2.5-5 MG DLY							
PO				Neurontin (Gabapentin)	SS		ORAL
PO							

Date:01/12/01ISR Number: 3647234-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100033
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Sickle Cell Anaemia With Crisis	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
4200 MG (1400 MG, TID)				Augmentin (Clavulanate Potassium, Amoxicillin Trihydrate)	C		
				Flovent (Fluticasone Propionate)	C		
				Proventil (Salbutamol Sulfate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/16/01ISR Number: 3648660-7Report Type:Expedited (15-DaCompany Report #EMADSS2001000052
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Face Oedema	Foreign	Duragesic	PS	Alza Corp	
Other		Urticaria Papular	Health				
TRANSDERMAL	50 MCG/H	1	Professional				
IN 72 HOUR(S)							
TRANSD				Gabapentin (Gabapentin)	SS		ORAL
300 MG 3 IN 1							
DAY(S) ORAL							
				Carbamazepine (Carbamazepine)	SS		ORAL
200 MG 3 IN 1							
DAY(S) ORAL							

Date:01/16/01ISR Number: 3649660-3Report Type:Periodic Company Report #2000UW01295
Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Zomig	PS	Astrazeneca	
Other		Drug Interaction		Sertraline	SS	Pharmaceuticals Lp	
				Gabapentin	SS		
				Cyproheptadine	C		

Date:01/17/01ISR Number: 3648107-0Report Type:Direct Company Report #
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Gabapentin	PS		ORAL
1200 MG TID		Insomnia					
ORAL		Sense Of Oppression					

Date:01/17/01ISR Number: 3649282-4Report Type:Expedited (15-DaCompany Report #001-0073-M0100016

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Visual Acuity Reduced	Consumer	Dilantin	PS	Parke Davis Div Warner Lambert Co	
				Neurontin (Gabapentin)	SS		

Date:01/17/01ISR Number: 3649325-8Report Type:Expedited (15-DaCompany Report #001-0945-M0100045

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Monoparesis	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

3200 MG

(DAILY) 2000

MG (DAILY)

Date:01/17/01ISR Number: 3649697-4Report Type:Expedited (15-DaCompany Report #031-0945-M0100003

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Post-Traumatic Stress Disorder	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

900 MG

(DAILY), PER

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Freedom Of Information (FOI) Report

ORAL

Amitriptyline C

Date:01/17/01ISR Number: 3649895-XReport Type:Expedited (15-DaCompany Report #A100542

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Convulsion	Health Professional	Zoloft	PS	Pfizer	
Intervention to Prevent Permanent Impairment/Damage				Neurontin	SS	Pharmaceuticals Inc	ORAL
TOTAL: BID:							

ORAL

Tylenol #3 C
Xanax C
Dilantin C

Date:01/17/01ISR Number: 3649942-5Report Type:Expedited (15-DaCompany Report #A0135986A

Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain	Consumer	Lotronex	PS	Glaxo Wellcome Inc	ORAL
1 MG / TWICE		Colitis					
PER DAY /		Colonic Polyp					

ORAL

Constipation	Gabapentin	SS
Dysgeusia	Paroxetine	
Dyspepsia	Hydrochloride	C
Nausea	Aspirin+Butalbital+C	
Oral Candidiasis	affn.	C
Rectal Haemorrhage	Sumatriptan	
	Succinate	C

Date:01/18/01ISR Number: 3651104-2Report Type:Expedited (15-DaCompany Report #055-0945-M0000020

Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Visual Acuity Reduced	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1200 MG (600 MG, BID), PER			Professional				
ORAL							

Date:01/19/01ISR Number: 3652097-4Report Type:Expedited (15-DaCompany Report #033-0945-M0100009
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Complications Of Maternal Exposure To Therapeutic	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
PLACENTAL		Drugs Foetal Growth Retardation	Professional				

Date:01/22/01ISR Number: 3652942-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100059
Age:93 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Confusional State
Other	Dyspnoea
	Gastrointestinal Disorder
	Mental Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Myocardial Infarction Pain	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL			Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Fosamax (Alendronate Sodium)	SS		
				Demerol (Pethidine Hydrochloride)	SS		
				Lotrel (Amlodipine Besylate, Benazepril Hydrochloride)	C		
				Norvasc (Amlodipine Besilate)	C		
				(Levothyroxine)	C		
				Beta Blocker Eye Drop	C		
				Unspecified Eye Drop	C		

Date:01/23/01ISR Number: 3654063-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100050
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL;		Fibromyalgia Muscular Weakness Musculoskeletal Disorder	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
2400 MG (DAILY), PER ORAL				Norvasc(Amlodipine Besilate)	C		
				Synthroid(Levothyrox ine Sodium)	C		
				Trazodone	C		
				Trilafon(Perphenazin			

e) C
 Depakote(Valproate Semisodium) C
 Klonopin(Clonazepam) C
 Efflexor(Venlafaxine Hydrochloride) C

Date:01/24/01ISR Number: 3654153-3Report Type:Expedited (15-DaCompany Report #001-0945-M0001380
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (DAILY) PER ORAL		Chest Pain Circadian Rhythm Sleep Disorder Dizziness	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
600 MG DAILY PER ORAL		Feeling Drunk Halo Vision Oxygen Saturation		Neurontin Capsules 300 Mg (Gabapentin)	SS		ORAL
300 MG (DAILY) PER ORAL		Decreased Sedation Tinnitus		Neurontin Capsules, 300 Mg (Gabapentin)	SS		ORAL
				Ziac			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Hydrochlorothiazide
 , Bisoprolol
 Fumarate) C
 Xanax (Alprazolam) C

Date:01/25/01ISR Number: 3655720-3Report Type:Expedited (15-DaCompany Report #A036897
 Age:58 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Amnesia Blood Pressure Decreased	Consumer Health	Procardia Xl	PS	Pfizer Laboratories Div Pfizer Inc	ORAL
300.00 MG	Pneumonia	Professional	Bupropion	SS		ORAL
TOTAL: BID: ORA	Sedation					
L	Speech Disorder					
	Toxicologic Test Abnormal		Gabapentin Atenolol	SS SS		

Date:01/25/01ISR Number: 3655725-2Report Type:Direct Company Report #
 Age:70 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1 CAPSULE 1 PER DAY ORAL	Abdominal Pain Upper Dyspepsia Nausea Vomiting		Neurontin 300 Mg Parke-Davis	PS	Parke-Davis	ORAL

Date:01/25/01ISR Number: 3656817-4Report Type:Expedited (15-DaCompany Report #044-0073-M0000028
 Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged SEE IMAGE	Urticaria	Foreign Health	Dilantin	PS	Parke Davis Div Warner Lambert Co	ORAL

900 MG Professional Gabapentin SS ORAL
 (DAILY), PER
 ORAL Carbamazepine C

Date:01/26/01ISR Number: 3655965-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100066
 Age:83 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Confusional State	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1200 MG (400 MG, TID), PER ORAL		Dizziness Grand Mal Convulsion Hallucination Oedema					

Date:01/26/01ISR Number: 3655975-5Report Type:Expedited (15-DaCompany Report #001-0981-M0100322
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coeliac Disease	Health Professional	Lipitor	PS	Pfizer Ireland Pharmaceuticals, Tablet Plant	
20 MG (DAILY) 900 MG (TID)				Neurontin (Gabapentin)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Zyrtec (Cetirizine Hydrochloride) SS
 Oxycodone Hydrochloride C
 Tramadol Hydrochloride C
 Metoclopramide C
 Pantoprazole C
 Carisoprodol C
 Venlafaxine Hydrochloride C
 Sucralfate C
 Metoprolol Succinate C
 Carbamazepine C

Date:01/26/01ISR Number: 3656004-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000616
 Age:4 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Autism	Health	Neurontin	PS	Parke Davis	
Other		Dementia	Professional			Pharmaceuticals Ltd	ORAL
SEE IMAGE		Psychomotor Hyperactivity		Depakote (Valproate Semisodium)	SS		
		Speech Disorder		Hydroxyzine Hydrochloride	C		
		Vomiting		Phenobarbital	C		
				Lactulose	C		
				Senokot	C		
				Ryna-12s	C		

Date:01/29/01ISR Number: 3656958-1Report Type:Periodic Company Report #A037518
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Neuralgia	Consumer	Tikosyn	PS	Pfizer	
Other			Health Professional			Pharmaceuticals Production Corp Ltd	ORAL

1000.00 MCG

TOTAL: BID: ORA

L

2400.00 MG

TOTAL:TID

Neurontin

SS

Lanoxin

C

Aspirin

C

Diltiazem

C

Naproxen

C

Lidopatch

C

Ultram

C

Restoril

C

Paxil

C

Date:01/30/01ISR Number: 3656653-9Report Type:Direct
Age:52 YR Gender:Male I/FU:I

Company Report #

Outcome
Life-Threatening
Hospitalization -
Initial or Prolonged
Disability

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Hospitalization - Initial or Prolonged "SEE IMAGE	Depression Disturbance In Attention	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd
	Fatigue Insomnia Oedema Peripheral Trigeminal Neuralgia		Celebrex (Celecoxib) Oxycontin (Oxycodone Hydrochloride) Clonopin (Clonazepam) Lorcet (Paracetamol, Hydrocodone Bitartrate)	SS C C C	

Date:01/30/01ISR Number: 3657930-8Report Type:Expedited (15-DaCompany Report #033-0945-M0100011
Age:48 YR Gender:Male I/FU:I

Outcome Dose Hospitalization - Initial or Prolonged 1600 MG Other (DAILY), PER ORAL	PT Blood Creatine Phosphokinase Increased Diabetic Ketoacidosis Hyperglycaemia Myocardial Infarction Pancreatic Pseudocyst Pancreatitis Necrotising	Report Source Foreign Health Professional	Product Neurontin Depakine Chrono (Valproic Acid, Valproate Sodium)	Role PS SS	Manufacturer Parke Davis Pharmaceuticals Ltd	Route ORAL ORAL
2000 MG (DAILY), PER ORAL			Gardenal			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Phenobarbital) SS ORAL

150 MG
(DAILY), PER

ORAL

Glucophage
(Metformin
Hydrochloride) C
Humulin Nph (Insulin
Human Injection,
Isophane) C
Mopral (Omeprazole) C
Fungizone
(Amphotericin B) C

Date:01/30/01ISR Number: 3658310-1Report Type:Expedited (15-DaCompany Report #032-0945-M000026
Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE TEXT, PER		Condition Aggravated Drug Ineffective	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
ORAL		Drug Interaction	Professional				
		Fall Neuropathy Peripheral Pain Sedation Vertigo		Rifampicin Isoniazid Ethambutol Pyrazinamide Morphine Sulfate Fentanyl	SS SS SS SS SS C		

Date:01/30/01ISR Number: 3659137-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000950
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (100 MG X3 QPM)		Condition Aggravated Depression	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
		Disorientation					

PER ORAL	Dizziness					
	Fall		Serzone (Nefazodone			
400 MG (200	Hypotension		Hydrochloride)	SS		ORAL
MG BID) PER	Post-Traumatic Stress					
ORAL	Disorder					
	Syncope		Neurontin			
400 MG (100			(Gabapentin)	SS		ORAL
MG X1 QAM &						
X3 QPM) PER						
ORAL						
			Serzone (Nefazodone			
375 MG			Hydrochloride)	SS		ORAL
(DAILY) PER						
ORAL						
			Klonopin			
			(Clonazepam)	C		

Date:01/30/01ISR Number: 3659920-8Report Type:Expedited (15-DaCompany Report #001-0945-M0100130
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion	Health	Neurontin	PS	Parke Davis	
Initial or Prolonged		Coordination Abnormal	Professional			Pharmaceuticals Ltd	
UNKNOWN							
Disability		Drug Interaction		Lamictal			
50 MG DAILY				(Lamotrigine)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

PER ORAL

200 MG (400

MG, BID) PER

ORAL

Lamictal
(Lamotrigine)

SS

ORAL

Date:01/31/01ISR Number: 3657867-4Report Type:Expedited (15-DaCompany Report #001-0073-M0100040

Age:48 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged PER ORAL, 400	Arthritis Convulsion	Consumer	Dilantin	PS	Parke Davis Div Warner Lambert Co	ORAL
Other MG (DAILY), PER ORAL	Dysgraphia Grand Mal Convulsion Reading Disorder		Neurontin (Gabapentin)	SS		ORAL
SEE IMAGE			Unspecified Arthritis Medication	C		

Date:02/01/01ISR Number: 3659970-1Report Type:Expedited (15-DaCompany Report #A101069

Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150.00 MG Required TOTAL:DAILY	Blood Pressure Decreased Disorientation	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	
Intervention to 400.00 MG Prevent Permanent TOTAL:PID	Dizziness Fall Insomnia		Neurontin	SS		
Impairment/Damage 375.00 MG	Oral Intake Reduced		Serzone	SS		

Syncope

TOTAL: BID

Proamatine SS
 Ambien C
 Unspecified Blood Pressure Medication C
 Klonopin C

Date: 02/01/01 ISR Number: 3660088-2 Report Type: Expedited (15-DaCompany Report #001-0945-M0001381
 Age: 57 YR Gender: Female I/FU: F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG, TID), PER ORAL ; 1800 MG (600 MG, TID)		Postmenopausal Haemorrhage	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Vioxx (Rofecoxib)	C		

Date: 02/01/01 ISR Number: 3660544-7 Report Type: Expedited (15-DaCompany Report #044-0945-M0100024
 Age: Gender: Female I/FU: I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Tenderness Tenderness	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/01ISR Number: 3660545-9Report Type:Expedited (15-DaCompany Report #055-0945-M0100001

Age:71 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (400 MG, TID), PER ORAL	Arrhythmia	Foreign Consumer	Neurontin (Diltiazem Hydrochloride)	PS C	Parke Davis Pharmaceuticals Ltd	ORAL

Date:02/02/01ISR Number: 3660343-6Report Type:Direct

Company Report #

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Hypotonia Loss Of Consciousness Sedation	Health Professional	Clozaril Depakote Neurontin Ativan	PS SS SS SS		

Date:02/05/01ISR Number: 3660381-3Report Type:Periodic

Company Report #A038643

Age:52 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 600.00 MG	Penis Disorder	Consumer	Zyrtec Neurontin	PS SS	Pfizer Inc	ORAL

TOTAL: BID:ORA

L

Synthorid C

Date:02/05/01ISR Number: 3660479-XReport Type:Direct
Age: Gender: I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 400MG PO @HS Initial or Prolonged	Dizziness Dysarthria Gait Disturbance Medication Error Vision Blurred		Neurontin	PS		ORAL

Date:02/05/01ISR Number: 3661087-7Report Type:Expedited (15-DaCompany Report #A101296

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Depression Post-Traumatic Stress Disorder	Health Professional	Zoloft Neurontin	PS SS	Pfizer Pharmaceuticals Inc	ORAL

1600.00 MG

TOTAL: QID:

ORAL

Vicodin	C
Colace	C
Seroquel	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/05/01ISR Number: 3661327-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100100

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Unspecified Medications	SS		

Date:02/05/01ISR Number: 3661537-6Report Type:Expedited (15-DaCompany Report #001-0991-M0001451

Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Abdominal Pain	Consumer	Rezulin	PS	Parke Davis Pharmaceuticals Ltd	
Other		Ammonia Increased					
		Anhedonia		Vasotec (Enalapril Maleate)	SS		
		Anxiety					
		Blood Urea Increased		Hyoscyamine (Hyoscyamine)	SS		
		Cholelithiasis					
		Cough		Amitriptyline (Amitriptyline)	SS		
		Depression					
		Dermatitis		Neurontin (Gabapentin)	SS		
		Emotional Distress					
300 MG							
(DAILY),		Fatigue					
UNKNOWN		Fungal Rash					
		Gallbladder Disorder		Diabeta (Glibenclamide)	C		
		Hepatic Cirrhosis					
		Hepatic Steatosis		Glucophage (Metformin Hydrochloride)	C		
		Hepatocellular Damage					
		Hypergammaglobulinaemia					
		Hypertension		Spiroinolactone (Spiroinolactone)	C		
		Ill-Defined Disorder					
		Irritable Bowel Syndrome		Zyrtec (Cetirizine Hydrochloride)	C		
		Metrorrhagia					
		Nausea		Demulen (Mestranol, Etyndiol Diacetate)	C		
		Neuralgia					
		Oedema Peripheral		Trazodone (Trazodone)	C		
		Oligomenorrhoea					
		Osteoarthritis		Vancenase (Beclometasone			
		Pain					

Platelet Count Decreased
Pollakiuria
Polyp
Purpura
Red Blood Cell Count
Decreased
Scar
Sedation
Serum Ferritin Increased
Skin Ulcer
Thermal Burn

Dipropionate)

C

Date:02/06/01ISR Number: 3660931-7Report Type:Direct

Company Report #

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Glossodynia		Neurontin	PS		ORAL
300MG PO TID		Vulvovaginal Discomfort		Premarin	C		
				Lysine	C		
				Dhea	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Calcium	C
Thyrolar	C
Levoxyl	C
Triazolam	C
Temazepam	C
Estrogen/Testosteron	
e/Progesterone	C
Vit E	C
Docusate	C

Date:02/06/01ISR Number: 3660950-0Report Type:Direct
 Age:49 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 100MG TID PO			Complex Regional Pain Syndrome Condition Aggravated	Neurontin 100mg	PS		ORAL

Date:02/06/01ISR Number: 3661566-2Report Type:Expedited (15-DaCompany Report #002-0945-M0100008
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - PER ORAL Initial or Prolonged			Myocardial Infarction Foreign Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:02/08/01ISR Number: 3662303-8Report Type:Direct
 Age:42 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 400 MG QID ORAL			Abnormal Behaviour Aggression	Neurontin	PS		ORAL

Anxiety	Fentanyl Patch	C
Aspiration	Prozac	C
Brain Damage	Inderal	C
Delusional Disorder,	Zyprexa	C
Persecutory Type	Klonopin	C
Hostility	Phenergan	C
Thinking Abnormal		
Vomiting		

Date:02/09/01ISR Number: 3663421-0Report Type:Expedited (15-DaCompany Report #001-0945-M0001385
Age:50 YR Gender:Female I/FU:F

Outcome	PT
Other	Balance Disorder
	Convulsion
	Decreased Appetite
	Disturbance In Attention
	Dizziness
	Dysarthria
	Epileptic Aura
	Head Injury
	Memory Impairment
	Nausea

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Adverse Event	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), PER ORAL		Nervousness Speech Disorder Tremor Weight Decreased	Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1200 MG (400 MG, TID), PER ORAL				Neurontin (Gabapentin)	SS		ORAL
3600 MG (900 MG, QID)				Neurontin(Gabapentin)	SS		
				Premarin (Estrogens Conjugated) Provera (Medroxyprogesterone Acetate) Zyprexa (Olanzapine) Depakote (Valproate Semisodium)	C C C C		

Date:02/09/01
 ISR Number: 3663763-9
 Report Type:Expedited (15-DaCompany Report #001-0945-M0100133
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
1800 MG (600 MG, TID)				Cyclobenzaprine Vicodin (Paracetamol,	SS		

Hydrocodone
 Bitartrate) C
 Ultram (Tramadol
 Hydrochloride C
 Paracetamol,
 Dextropropoxyphene C
 Naprosyn (Naproxen) C

Date:02/09/01ISR Number: 3663766-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100134
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nephritis Interstitial	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Unspecified Antibiotic	C		

Date:02/09/01ISR Number: 3663787-1Report Type:Expedited (15-DaCompany Report #047-0945-M0100001
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Folate Deficiency Polyneuropathy	Foreign Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
		Transient Ischaemic Attack		Analgesics	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/09/01ISR Number: 3663987-0Report Type:Expedited (15-DaCompany Report #A102512

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Coeliac Disease	Health Professional	Zyrtec	PS	Pfizer Inc	
Intervention to				Lipitor	SS		
20.00 MG							
Prevent Permanent							
TOTAL:DAILY				Neurontin	SS		
Impairment/Damage							
900.00 MG							
TOTAL:TID							
				Oxycontin	C		
				Ultram	C		
				Reglan	C		
				Protonix	C		
				Soma	C		
				Effexor Xr	C		
				Carafate	C		
				Toprol Xl	C		
				Tegretol	C		

Date:02/12/01ISR Number: 3664294-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100153

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Clonic Convulsion	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/12/01ISR Number: 3664402-3Report Type:Expedited (15-DaCompany Report #2001SE01042

Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Cardiac Failure	Foreign Health Professional	Pulmicort	PS	Astrazeneca Pharmaceuticals Lp	
Initial or Prolonged		Coma					
		Overdose		Allopurinol Choay	SS		

2.5 MG BID	Respiratory Disorder	Other	Hemi-Daonil	SS
300 MG DAILY			Fonzyllane	SS
100 MG TID			Lasilix	SS
			Neurontin	SS
			Acuitel	C
			Burinex	C
			Monicor	C
			Aspirine	C

Date:02/13/01ISR Number: 3665130-0Report Type:Expedited (15-DaCompany Report #PHNU2001DE00577
Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia Convulsion	Foreign Health	Trileptal	PS	Novartis Pharmaceuticals Corp	ORAL
600 MG, BID, ORAL		Headache	Professional				
400 MG, QID		Overdose Vertigo	Company Representative	Neurontin (Gabapentin) Regimen	SS		
		Vomiting	Other	Orfiril "Desitin"	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/13/01ISR Number: 3665203-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000488

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Pain	Health	Neurontin	PS	Parke Davis	ORAL
4800 MG (1200		Pulmonary Hypertension	Professional			Pharmaceuticals Ltd	
MG, TID), PER		Pulmonary Oedema					
ORAL		Sleep Apnoea Syndrome					
				Oxycontin (Oxycodone Hydrochloride)	SS		
				Unspecified Medication	C		

Date:02/13/01ISR Number: 3665430-4Report Type:Direct

Company Report #

Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Diplopia		Gabapentin	PS		
		Eye Pain		Olanzapine	SS		

Date:02/13/01ISR Number: 3666436-1Report Type:Expedited (15-DaCompany Report #358-0945-M0100001

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1200 MG		Confusional State	Foreign	Neurontin	PS		ORAL
Initial or Prolonged (DAILY), PER		Dizziness	Health				
ORAL		Drug Interaction	Professional				
		Dysstasia		Panacod (Codeine Phosphate, Paracetamol)	SS		
		Movement Disorder		Diltiazem Hydrochloride	C		
		Pneumonia		Nizax (Nizatidine)	C		
				Amitriptyline			

Hydrochloride	C
Levofloxacin	C
Tazocin (Tazobactam Sodium)	C
Klexane (Heparin-Fraction, Sodium Salt)	C
Levoac	C

Date:02/14/01ISR Number: 3666754-7Report Type:Expedited (15-DaCompany Report #HQ7038409FEB2001
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Consumer	Lorazepam	PS		ORAL
0.5 MG AS							
NEEDED, ORAL				Dilantin (Phenytoin Sodium)	SS		
				Neurontin (Gabapentin)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/01ISR Number: 3666772-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000998

Age:72 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (300 Other MG BID), PER ORAL	Cerebrovascular Accident Coma Complex Partial Seizures Convulsion Dizziness Postural	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Cognex (Tacrine Hydrochloride)	C		
			Detrol (Tolterodine Tartrate)	C		
			(Selenium)	C		
			Vitamin E (Tocopherol)	C		
			Estrogen	C		
			Lorazepam	C		
			Zantac (Ranitidine Hydrochloride)	C		

Date:02/15/01ISR Number: 3665971-XReport Type:Direct

Age:76 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Necrosis Ischaemic Pulmonary Oedema		Gabapentin	PS		

Date:02/15/01ISR Number: 3665997-6Report Type:Direct

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

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent Impairment/Damage	Fatigue Liver Function Test Abnormal		Pioglitazone	PS		
			Furosemide 40 Mg Tab	SS		
			Glyburide 5mg Tab	SS		
			Spironolactone 25mg Tab	SS		
			Gabapentin 600mg Tab	SS		

Fluoxetine Hcl 20mg
Cap SS
Lisinopril 40mg Tab SS

Date:02/15/01ISR Number: 3666590-1Report Type:Expedited (15-DaCompany Report #033-0945-M0100017
Age:69 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG (100 MG TID) PER ORAL	Accidental Overdose Coma	Foreign Health Professional	Neurotin (Gabapentin)	PS		ORAL
ORAL			Lasilix (Furosemide)	SS		ORAL
ORAL			Allopurinol	SS		ORAL
ORAL			Pulmicort (Budesonide)	SS		
2.5 MG DAILY PER ORAL			Hemi-Daonil (Glibenclamide)	SS		ORAL
600 MG DAILY			Fonzyllane (Buflomedil Hydrochloride)	SS		ORAL

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Freedom Of Information (FOI) Report

PER ORAL

Date:02/15/01ISR Number: 3666595-0Report Type:Expedited (15-DaCompany Report #2001SE01042
 Age:69 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiac Failure Coma Overdose	Foreign Health Professional	Pulmicort Allopurinol Choay Hemi-Daonil	PS SS SS		
2.5 MG BID							
300 MG DIALY		Respiratory Disorder	Other	Fonzylane	SS		
				Lasilix	SS		
100 MG TID				Neurontin	SS		
				Acuitel	C		
				Burinex	C		
				Monicor	C		
				Aspirine	C		

Date:02/15/01ISR Number: 3666979-0Report Type:Expedited (15-DaCompany Report #001-0981-M0100322
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Coeliac Disease	Health Professional	Atorvastatin (Atorvastatin)	PS		
20 MG (DAILY)				Neurontin (Gabapentin)	SS		
900 MG (TID)				Zyrtec (Cetirizine Hydrochloride)	SS		
				(Oxycodone Hydrochloride)	C		
				(Tramadol Hydrochloride)	C		
				(Metoclopramide)	C		
				(Pantoprazole)	C		
				(Carisoprodol)	C		
				(Venlafaxine Hydrochloride)	C		

(Sucralfate) C
(Metoprolol Succinate) C
(Carbamazepine) C

Date:02/15/01ISR Number: 3667107-8Report Type:Expedited (15-DaCompany Report #001-0945-M0100148
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Burning Sensation Complex Regional Pain Syndrome	Consumer Health Professional	Neutrotin (Gabapentin)	PS		ORAL
100 MG (DAILY), PER ORAL		Condition Aggravated		Neurontin (Gabapentin)	SS		ORAL
300 MG QD		Electromyogram Abnormal Hypoaesthesia Pain Paraesthesia Thermogram Abnormal		(Oxycodone Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/16/01ISR Number: 3667759-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000781

Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Health Professional	Neurontin (Gabapentin)	PS	Parke Davis Pharmaceuticals Ltd	ORAL

SEE IMAGE

Propoxyphene (Dextropropoxyphene)	SS
Nortriptyline(Nortriptyline)	SS
Methadone (Methadone)	C
Topamax (Topiramate)	C
Xanax (Alprazolam)	C
Effexor (Venlafaxine Hydrochloride)	C
(Diphenhydramine)	C
Norpropoxyphene	C

Date:02/16/01ISR Number: 3667760-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000782

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Toxicity Overdose Sudden Death	Health Professional	Neurontin (Gabapentin)	PS	Parke Davis Pharmaceuticals Ltd	ORAL

7200 MG (2400

MG, TID), PER

ORAL

150 MCG (Q 3

DAYS)

Duragesic (Fentanyl)	SS
Nortriptyline	C
Oxycontin (Oxycodone Hydrochloride)	C
Paxil (Paroxetine Hydrochloride)	C
Allopurinol	

(Allopurinol) C
Sudafed
(Pseudoephedrine
Hydrochloride) C
Clonapin
(Clonazepam) C
Benadryl
(Diphenhydramine
Hydrochloride) C
Lomotil (Atropine
Sulfate,
Diphenoxylate
Hydrochloride) C
Vioxx (Rofecoxib) C

Date:02/20/01ISR Number: 3667183-2Report Type:Direct
Age:51 YR Gender:Female I/FU:I

Company Report #

Outcome PT
Hospitalization - Atrial Fibrillation
Initial or Prolonged Cardiac Failure
Disability Oedema
Pneumonia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Renal Failure Respiratory Failure Sinusitis	Report Source	Product	Role	Manufacturer	Route
300 MGM TID				Neurontin	PS		ORAL
				Tilgade	C		
				Accolate	C		
				Serevent	C		
				Allergy Shots	C		

Date:02/20/01ISR Number: 3668598-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100158
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Anuria	Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
"SEE IMAGE"				Prevacid (Lansoprazole)	C		
				Muscle Relaxant	C		

Date:02/21/01ISR Number: 3668636-3Report Type:Expedited (15-DaCompany Report #055-0945-M0100002
Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (400 MG, TID), PER ORAL		Convulsion	Foreign Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Valproic Acid	C		
				Valproate Sodium	C		

Date:02/21/01ISR Number: 3669398-6Report Type:Expedited (15-DaCompany Report #NSADSS2001002487
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Level Above Therapeutic	Health Professional	Topamax (Unspecified) (Topiramate)	PS		
				Alprazolam (Alprazolam)	SS		
				Gabapentin (Gabapentin)	SS		
				Carbamazepine (Carbamazepine)	SS		

Date:02/22/01ISR Number: 3669600-0Report Type:Expedited (15-DaCompany Report #001-0073-M0100040
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	PER ORAL, 40	Arthritis Convulsion	Consumer	Dilantin	PS	Parke Davis Div Warner Lambert Co	ORAL
Other	MG (DAILY),	Dysgraphia					
PER ORAL		Grand Mal Convulsion					
		Reading Disorder		Neurontin (Gabapentin)	SS		ORAL
				Unspecified			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Arthritis Medication C

Date:02/22/01ISR Number: 3669601-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100197
 Age:43 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Dependence Medication Error	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
900 MG (300 MG, TID), PER ORAL							

Date:02/22/01ISR Number: 3669602-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100169
 Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged HIGH DOSE		Drug Ineffective Drug Withdrawal Syndrome	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Malaise Tremor							

Date:02/22/01ISR Number: 3669972-7Report Type:Expedited (15-DaCompany Report #002-0945-M0100015
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Clumsiness Confusional State	Foreign Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
300 MG DAILY PER ORAL; 1500 MG DAILY PER ORAL							

Iron	C
Clonidine	C

Date:02/23/01ISR Number: 3670105-1Report Type:Expedited (15-DaCompany Report #001-0991-M0001451
Age:44 YR Gender:Female I/FU:F

Outcome	PT
Disability	Abdominal Mass
Other	Anhedonia
	Anxiety
	Back Pain
	Benign Laryngeal Neoplasm
	Cholelithiasis
	Cough
	Depression
	Emotional Disorder
	Fungal Rash
	Hepatic Cirrhosis
	Hepatic Steatosis
	Hepatocellular Damage
	Hepatomegaly
	Hypertension
	Ill-Defined Disorder
	Injury
	Irritable Bowel Syndrome
	Liver Function Test
	Abnormal

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nausea Oedema Peripheral Oligomenorrhoea	Consumer	Rezulin	PS	Parke Davis Pharmaceuticals Ltd	
SEE IMAGE		Pain Pharyngolaryngeal Pain					
		Pollakiuria Psoriasis Purpura Thermal Burn		Vasotec (Enalapril Maleate) Hyoscyamine (Hyoscyamine) Amitriptyline (Amitriltyline) Neurontin (Gabapentin)	SS SS SS SS		
SEE IMAGE				Diabeta (Glibenclamide) Glucophage (Metformin Hydrochloride) Spironolactone (Spironolactone) Zyrtec (Cetirizine Hydrochloride) Demulen (Mestranol, Etyndiol Diacetate) Trazodone (Trazodone) Vancenase (Beclometasone Dipropionate) Axid (Nizatidine) Glyburide (Glibenclamide)	C C C C C C		

Date:02/26/01ISR Number: 3671118-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100190
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (100		Delusion Dyspnoea	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

MG, BID), PER	Euphoric Mood			
ORAL	Feeling Abnormal			
400 MG (200	Hallucination	Neurontin		
MG, BID), PER	Heart Rate Increased	(Gabapentum)	SS	ORAL
ORAL	Hyperreflexia			
800 MG (400	Insomnia			
MG, BID), PER	Oral Intake Reduced	Neurontin		
ORAL	Paranoia	(Gabapentin)	SS	ORAL
1800 MG (400				
MG, BID), PER		Neurontin		
ORAL		(Gabapentin)	SS	ORAL
900 MG				
(DAILY), PER		Neurontin		
ORAL		(Gabapentin)	SS	ORAL
500 MG, PER				
ORAL		Neurontin		
		(Gabapentin)	SS	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

400 MG, PER	Neurontin (Gabapentin)	SS	ORAL
ORAL			
200 MG, PER	Neurontin (Gabapentin)	SS	ORAL
ORAL			

Date:02/26/01ISR Number: 3671137-XReport Type:Expedited (15-DaCompany Report #044-0945-M0100024
Age:49 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 6.4 GM (DAILY), PER	Abdominal Tenderness Accidental Overdose	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
ORAL						
2.4 GM (DAILY), PER			Neurontin (Gabapentin)	SS		ORAL
ORAL						
			(Amitriptyline)	C		

Date:02/27/01ISR Number: 3671673-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100214
Age:31 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 900 MG (300 MG TID) PER	Diabetes Mellitus	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

ORAL

Prozac (Fluoxetine Hydrochloride)	C
Diovan (Valsartan)	C
Imitrex (Sumatriptan Succinate)	C
Unspecified Birth Control Pills	C

Date:02/27/01ISR Number: 3671675-XReport Type:Expedited (15-DaCompany Report #001-0945-M0001385
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Balance Disorder Decreased Appetite	Consumer Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
900 MG 300 MG		Disturbance In Attention	Professional				
TID PER ORAL;		Dysarthria					
2400 MG DAILY		Epileptic Aura					
ORAL; 3600 MG		Grand Mal Convulsion					
(900 MG QID)		Head Injury		Premarin (Estrogens Conjugated)	C		
		Memory Impairment		Provera			
		Nausea		(Medroxyprogesterone Acetate)	C		
		Nervousness		Zyprexa (Olanzapine)	C		
		Photopsia		Depakote (Valproate Semisodium)	C		
		Thinking Abnormal		Heroin (Diamorphine)	C		
		Tremor		Methadone			
		Weight Decreased		(Methadone)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Librium
(Chlordiazepoxide
Hydrochloride) C

Date:02/28/01ISR Number: 3671474-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000462
Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER ORAL		Aphonia Blood Glucose Increased Cellulitis Disturbance In Attention	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
75 MG (25 MG, TID), PER ORAL		Dizziness Headache Nausea Tremor Vision Blurred		Baclofen (Baclofen) (Insulin) Aleve (Naproxen Sodium)	SS C C		ORAL

Date:02/28/01ISR Number: 3671767-5Report Type:Expedited (15-DaCompany Report #044-0945-M0100043
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchopneumonia Metastases To Lung Respiratory Arrest	Foreign Health Professional Company Representative	Neurontin Analgesia Chemotherapy	PS C C	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671832-2Report Type:Periodic Company Report #001-0945-990640
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER		Pancreatitis	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
ORAL							

Date:02/28/01ISR Number: 3671833-4Report Type:Periodic Company Report #001-0945-M0001337
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG, UNKNOWN		Hypoventilation Respiratory Failure	Literature Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Albuterol (Salbutamol)	C		
				Ipratropium (Ipratropium)	C		
				Clonazepam (Clonazepam)	C		
				Zolpidem (Zolpidem)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/01ISR Number: 3671834-6Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-M0001344

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671835-8Report Type:Periodic
Age:77 YR Gender:Female I/FU:I

Company Report #001-0945-M0100029

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Alopecia	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Other		Deafness Visual Acuity Reduced		Depo-Medrol (Methylprednisolone Acetate) Cardura (Doxazosin Mesilate)	SS C		

Date:02/28/01ISR Number: 3671836-XReport Type:Periodic
Age:87 YR Gender:Male I/FU:F

Company Report #001-0945-980074

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Dizziness	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

900 MG (300
MG, TID), PER

ORAL

Synthroid (Levothyroxine Sodium)	C
Mutivitamins (Ergocalciferol, Ascorbic Acid, Thiamine Hydrochloride, Furosemide)	C C

Date:02/28/01ISR Number: 3671837-1Report Type:Periodic
Age:55 YR Gender:Female I/FU:F

Company Report #001-0945-991016

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 - 1200 MG (DAILY), PER ORAL	Pyrexia	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Amitriptyline C

Date:02/28/01ISR Number: 3671838-3Report Type:Periodic
Age: Gender:Male I/FU:F

Company Report #001-0945-991017

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Stevens-Johnson Syndrome	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/01ISR Number: 3671839-5Report Type:Periodic
Age:35 YR Gender:Male I/FU:F

Company Report #001-0945-991099

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blindness	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
1200 MG (300 MG, QID), UNKNOWN							

				Flexeril (Cyclobenzaprine Hydrochloride)	C		
--	--	--	--	--	---	--	--

Date:02/28/01ISR Number: 3671840-1Report Type:Periodic
Age:48 YR Gender:Female I/FU:F

Company Report #001-0945-991197

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1600 MG (400 MG, QID), PER ORAL		Dental Caries Drug Interaction Fatigue Hypotension Oral Intake Reduced Sedation Stupor Thinking Abnormal	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Tylenol Pm (Diphenhydramine, Paracetamol)	SS		
				Propulsid (Cisapride)	C		
				Prevacid (Lansoprazole)	C		
				Calcium	C		
				Estratest (Methyltestosterone, Estrogens Esterified)	C		
				Baclofen	C		
				Celebrex (Clecocixib)	C		
				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic			

Acid, Thiamine
 Hydrochloride, C
 Celexa (Citalopram
 Hydrobromide) C
 Xanax (Alprazolam) C
 Vistaril
 (Hydroxyzine
 Embonate) C

Date:02/28/01ISR Number: 3671841-3Report Type:Periodic
 Age:20 YR Gender:Female I/FU:F

Company Report #001-0945-991268

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
2700 MG (900 MG, TID), PER ORAL				Depakote (Valproate Semisodium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/01ISR Number: 3671842-5Report Type:Periodic
 Age:26 YR Gender:Male I/FU:I

Company Report #001-0945-990606

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other			Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1800 MG (300 MG, 2TID), PER ORAL		Blindness Transient Convulsion					
600 MG, (100 MG, 2TID), PER ORAL				Neurontin Capsules 100 Mg (Gabapentin)	SS		ORAL
				Lamictal (Lamotrigine)	C		
				Magnesium (Magnesium)	C		
				Vitamin C And 3 (Ascorbic Acid, Tocopherol)	C		
				Vitamin B6 (Pyridoxine Hydrochloride)	C		

Date:02/28/01ISR Number: 3671843-7Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-M0001186

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671844-9Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-M0001187

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Death		Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Date:02/28/01ISR Number: 3671845-0Report Type:Periodic Company Report #001-0945-M0001188							
Age: Gender:Unknown I/FU:I							
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671846-2Report Type:Periodic Company Report #001-0945-M0001189							
Age: Gender:Unknown I/FU:I							
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671847-4Report Type:Periodic Company Report #001-0945-M0001190							
Age: Gender:Unknown I/FU:I							
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671850-4Report Type:Periodic Company Report #001-0945-M0001246
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671851-6Report Type:Periodic Company Report #001-0945-M0001266
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blindness	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

1) 900 MG

(300 MG, TID,

PER ORAL; 2)

600 MG (300

MG, BID), PER

Novolin 70/30
(Insulin Human,
Insulin Human
Injection, Isophane) C
Lipitor

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Atorvastatin) C
 Monopril (Fosinopril Sodium) C
 (Furosemide) C
 Zantac (Ranitidine Hydrochloride) C

Date:02/28/01ISR Number: 3671852-8Report Type:Periodic Company Report #001-0945-M0001286
 Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG TO 2400 MG	Amnesia Disturbance In Attention	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
(PRN), PER		Hypersensitivity					
ORAL		Mood Altered					
		Myeloid Metaplasia					
		Pneumonia		Unspecified Antibiotic	SS		
INTRAVENOUS	INTRAVENOUS			Altace (Ramipril)	C		
				Revacid (Lansoprazole)	C		
				(Triamterene)	C		
				(Theophylline)	C		

Date:02/28/01ISR Number: 3671853-XReport Type:Periodic Company Report #001-0945-M0000994
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Hypertension	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Ativan (Lorazepam)	C		
				Unspecified Medications	C		

Date:02/28/01ISR Number: 3671854-1Report Type:Periodic Company Report #001-0945-M0001132
Age:30 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Pyrexia	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671855-3Report Type:Periodic Company Report #001-0945-M0001133
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Renal Failure	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671856-5Report Type:Periodic Company Report #001-0945-M0001141
Age:56 YR Gender:Male I/FU:I

Outcome	PT
Other	Alopecia Arthralgia Hair Colour Changes

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hyperhidrosis
Rash Pruritic

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
UNK PER ORAL		Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
			Unspecified Opiates	SS		

Date:02/28/01ISR Number: 3671857-7Report Type:Periodic Company Report #001-0945-M0001176
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 MG BID) PER ORAL		Deep Vein Thrombosis Uterine Haemorrhage	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Depo-Provera (Medroxyprogesterone Acetate)	SS		
				Risperdal (Risperidone)	SS		ORAL
				Serzone (Nefazodone Hydrochloride) (Trazodone)	C C		

Date:02/28/01ISR Number: 3671858-9Report Type:Periodic Company Report #001-0945-M0001179
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671859-0Report Type:Periodic Company Report #001-0945-M0001182
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671860-7Report Type:Periodic Company Report #001-0945-M0001183
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671861-9Report Type:Periodic Company Report #001-0945-M0001184
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/01ISR Number: 3671862-0Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-M0001185

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671863-2Report Type:Periodic
Age:36 YR Gender:Female I/FU:I

Company Report #001-0945-M0000685

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2700 MG (900	Intentional Misuse Loss Of Consciousness	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

MG, TID), PER

ORAL

Ag-1549	C
Combivir	C
Viracept	C

Date:02/28/01ISR Number: 3671864-4Report Type:Periodic
Age:52 YR Gender:Male I/FU:I

Company Report #001-0945-M0000697

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Brain Neoplasm Dizziness	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

SEE TEXT, PER

ORAL

Dysphemia	Company
Intentional Misuse Petit Mal Epilepsy Tremor	Representative

Date:02/28/01ISR Number: 3671865-6Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-M0000792

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorgasmia	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
200 MG							
(DAILY), PER							
ORAL							

Date:02/28/01ISR Number: 3671866-8Report Type:Periodic Company Report #001-0945-M0000817
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Drug Level Above Therapeutic Sudden Death	Health Professional	Neurontin Hydrocodone	PS C	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671867-XReport Type:Periodic Company Report #001-0945-M0000821
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Coordination Abnormal	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
PRN;1500 MG							

(DAILY), PER

ORAL

Percocet
(Paracetamol,

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

2 TABLET(S)	Oxycodone Hydrochloride	SS	ORAL
(DAILY), PER			
ORAL			
2 TABLET(S)	Tylenol (Paracetamol)	SS	ORAL
(DAILY), PER			
ORAL			
	Coumadin	C	

Date:02/28/01ISR Number: 3671869-3Report Type:Periodic Company Report #001-0945-M0000877
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Neurontin	PS	Parke Davis	
Other		Brain Neoplasm	Professional			Pharmaceuticals Ltd	
5200 MG		Intentional Misuse					
(DAILY)							

Date:02/28/01ISR Number: 3671870-XReport Type:Periodic Company Report #001-0945-M0000884
 Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Asthenia	Consumer	Neurontin	PS	Parke Davis	
Initial or Prolonged		Blood Pressure Decreased				Pharmaceuticals Ltd	ORAL
900 MG (300		Circulatory Collapse					
MG TID), PER		Hallucination					
ORAL		Muscle Twitching		Cefalexin	C		
		Speech Disorder		Percocet	C		
		Syncope		Sr Morphine	C		
		Tremor		(Morphine)	C		
				Ativan (Lorazepam)	C		

Zoloft (Sertraline
Hydrochloride) C
Nitrostat (Glyceryl
Trinitrate) C

Date:02/28/01ISR Number: 3671874-7Report Type:Periodic Company Report #001-0945-M0000897
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER ORAL		Suicide Attempt	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Sinemet	C		
				Digoxin	C		

Date:02/28/01ISR Number: 3671875-9Report Type:Periodic Company Report #001-0945-M0000938
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (300 MG QAM 600 MG QPM), PER ORAL		Breast Cancer Female	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Wellbutrin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/01ISR Number: 3671876-0Report Type:Periodic
Age:31 YR Gender:Female I/FU:I

Company Report #001-0945-M0000967

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
600 MG			Company Representative				

Date:02/28/01ISR Number: 3671883-8Report Type:Periodic
Age:33 YR Gender:Female I/FU:I

Company Report #001-0945-M0000460

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hallucination	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
300 MG (100 MG, TID), PER							
ORAL							

Buspar C

Date:02/28/01ISR Number: 3671884-XReport Type:Periodic
Age:21 YR Gender:Female I/FU:I

Company Report #001-0945-M0000497

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
PER ORAL		Impulse-Control Disorder					
		Irritability		Prolixin	C		
		Mood Altered		Cogentin	C		
		Pressure Of Speech		Zyprexa	C		
				Vitamin E	C		
				Tegretol	C		
				Buspar	C		
				Carbatrol	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Encephalopathy	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG (100 MG, TID), PER ORAL		Ageusia Blindness Cataract Condition Aggravated Visual Disturbance	Consumer	Neurontin Glucophage Prilosec Niaspan Vasotec Lipitor Elavil	PS C C C C C C	Parke Davis Pharmaceuticals Ltd	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/01ISR Number: 3671890-5Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-M0000573

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Encephalopathy	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671895-4Report Type:Periodic
 Age:37 YR Gender:Male I/FU:I

Company Report #001-0945-M0000575

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 1200 MG (300 MG BID, 600 MG MG QHS)		Weight Increased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Serzone	C
Serevent	C
Atrovent	C
Symmetrel	C
Risperdal	C
Zyprexa	C
Klonopin	C
Effexor Xr	C
Dalmane	C

Date:02/28/01ISR Number: 3671898-XReport Type:Periodic
 Age:36 YR Gender:Female I/FU:I

Company Report #001-0945-M0000577

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other SEE IMAGE		Ammonia Increased Weight Increased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Depakote	C
Risperdal	C
Ditropan	C

Pepcid C
 Albuterol Inhaler C
 Cipro Eye Drops C
 Haldol Decanoate C
 Haldol C

Date:02/28/01ISR Number: 3671901-7Report Type:Periodic Company Report #001-0945-M0000578
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Glucose Increased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Other		Mania					
1500 MG (600							
MG AM AND 900							
MG QHS)							

Depakote C
 Cogentin C
 Valium C
 Serentil C
 Albuterol C
 Atrovent C
 Depo Provera C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/01ISR Number: 3671904-2Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-M0000642

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Leukocytosis	Health Professional Company Representative	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671905-4Report Type:Periodic
Age:76 YR Gender:Male I/FU:I

Company Report #001-0945-M0000681

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Alanine Aminotransferase Increased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
900 MG (300 MG, TID), PER ORAL		Blood Alkaline Phosphatase Increased Coordination Abnormal Dizziness		Glyburide Glucophage Procrit Vasotec Prilosec Vitamin E Multivitamin Calcium Many Unspecified Medications	C C C C C C C C		

Date:02/28/01ISR Number: 3671910-8Report Type:Periodic
Age:80 YR Gender:Female I/FU:I

Company Report #001-0945-M0000199

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE TEXT PER ORAL, 800 MG		Exostosis Gingival Bleeding Haematocrit Decreased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

(DAILY) PER

Haemoglobin Decreased

ORAL, SEE

Rectal Haemorrhage

TEXT PER ORAL

Lanoxin (Digoxin)	C
Premarin (Estrogens Conjugated)	C
Oxygen Via Nasal Cannula (Oxygen)	C
Darvocet (Paracetamol)	C
Darvocet (Paracetamol, Dextropropoxyphene)	C
Requip (Digoxin)	C
Prilosec (Omeprazole)	C
(Meprobamate)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/01ISR Number: 3671912-1Report Type:Periodic
Age:25 YR Gender:Female I/FU:I

Company Report #001-0945-M0000262

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 MG TID) PER ORAL	Abdominal Pain Deafness Vomiting	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
			Depakote (Valproate Semisodium) Dhea (Prasterone) Phenergan (Promethazine Hydrochloride) Zithromax (Azithromycin) Narcotics	C C C C C		

Date:02/28/01ISR Number: 3671914-5Report Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #001-0945-M0000282

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1500 MG	Crying Suicide Attempt	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
			Elavil (Amitriptyline Hydrochloride)	C		

Date:02/28/01ISR Number: 3671915-7Report Type:Periodic
Age:57 YR Gender:Female I/FU:I

Company Report #001-0945-M0000295

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG(300	Asthenia Dizziness	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

MG BID) PER
 ORAL
 400 MG
 (DAILY) PER
 ORAL

Dysarthria
 Hypertension
 Vomiting

Neurontin Capsules
 400 Mg (Gabapentin) SS ORAL

ORAL

Oxycontin (Oxycodone Hydrochloride)
 (Atenolol) C
 C

Date:02/28/01ISR Number: 3671917-0Report Type:Periodic Company Report #001-0945-M0000323
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	600 MG (300)	Pancreatitis	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

MG BID) PER
 ORAL

Lopressor
 (Metoprolol Tartrate) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/01ISR Number: 3671919-4Report Type:Periodic
Age:10 YR Gender:Male I/FU:I

Company Report #001-0945-M0000330

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG DAILY PER ORAL		Asthenia Back Pain Dysarthria Facial Palsy Hiccups Hypotonia Micturition Urgency Muscle Rigidity Muscle Twitching	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:02/28/01ISR Number: 3671921-2Report Type:Periodic
Age:29 YR Gender:Female I/FU:I

Company Report #001-0945-M0000332

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 700 MG Other (DAILY) PER ORAL		Alanine Aminotransferase Increased Aspartate Aminotransferase Increased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Ambien (Zolpidem Tartrate) Benadryl (Diphenhydramine Hydrochloride) Multivitamins(Vitami ns Nos, Minerals Nos) (Docusate) Nicotine Patch(Nicotine) Nicorette Gum (Nicotine Resin)	C C C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Hypoaesthesia Tunnel Vision	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
300 MG							

(DAILY) PER

ORAL

Glucophage (Metformin Hydrochloride)	C
Diabeta (Glibenclamide)	C

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Oedema	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
2400 MG			Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/01ISR Number: 3671924-8Report Type:Periodic
Age:30 YR Gender:Female I/FU:I

Company Report #001-0945-M0000386

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diplopia Disturbance In Attention Intentional Misuse Vision Blurred	Health Professional	Neurontin Trileptal (Oxcarbazepine)	PS SS	Parke Davis Pharmaceuticals Ltd	
2700 MG							
(DAILY),							
UNKNOWN; 5400							
MG (DAILY)							
UNKNOWN; 3600							

Kepra (Lefitracetam) C

Date:02/28/01ISR Number: 3671925-XReport Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-M0000025

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pancreatitis	Other	Neurontin	PS	Parke-Davis Pharm. Research	

Date:02/28/01ISR Number: 3671926-1Report Type:Periodic
Age:32 YR Gender:Female I/FU:I

Company Report #001-0945-M0000028

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 300 MG		Pancreatitis	Health Professional	Neurontin	PS	Parke-Davis Pharm. Research	

(DAILY)

Zoloft (Sertraline
Hydrochloride) C
Vicodin
(Paracetamol,
Hydrocodone)

Bitartrate)

C

Date:02/28/01ISR Number: 3671927-3Report Type:Periodic
Age:58 YR Gender:Female I/FU:I

Company Report #001-0945-M0000085

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE IMAGE		Convulsion Pain	Consumer	Neurontin	PS	Parke-Davis Pharm. Research	ORAL
		Sedation		(Lithium) Synthroid (Levothyroxine Sodium)	C		
				Serax (Oxazepam)	C		
				Premarin (Estrogens Conjugated)	C		
				Dilantin (Phenytoin Sodium)	C		

Date:02/28/01ISR Number: 3671928-5Report Type:Periodic
Age:38 YR Gender:Female I/FU:I

Company Report #001-0945-M0000127

Outcome	PT	Report Source
Other	Deafness	Health Professional Company

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
200 MG (100 MG, BID), UNK		Neurontin 100 Mg (Gabapenn)	PS	Parke-Davis Pharm. Research	
		Procardia (Nifedipine)	C		
		Monopril (Fosinopril Sodium)	C		
		Lipitor (Atorvastin)	C		

Date:02/28/01ISR Number: 3671938-8Report Type:Periodic Company Report #001-0945-M0000132
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Visual Disturbance	Health Professional	Neurontin	PS	Parke-Davis Pharm. Research	ORAL
400 MG (QHS), PER ORAL				Ativan(Lorazepam)	C		

Date:02/28/01ISR Number: 3671941-8Report Type:Periodic Company Report #001-0945-M0000149
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hepatomegaly	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:02/28/01ISR Number: 3671943-1Report Type:Periodic Company Report #001-0945-M0000158
 Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Eye Haemorrhage Vitreous Disorder	Health Professional	Neurontin	PS	Parke-Davis Pharm. Research	
300 MG TID OR							
QID							
				Rezulin(Troglitazone)	C		
				Accupril (Quinapril Hydrochloride)	C		

Date:02/28/01ISR Number: 3671945-5Report Type:Periodic Company Report #001-0945-M0000164
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cerebrovascular Accident	Health Professional Company Representative	Neurontin	PS	Parke-Davis Pharm. Research	

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 100.00 Required TOTAL:DAILY:0 Intervention to RAL		Depression Suicide Attempt	Other	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
Prevent Permanent 900.00 MG Impairment/Damage TOTAL:DAILY:0 RAL				Gabapentin	SS		ORAL
				Naproxen Chlorzoxazone	C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Liver Function Test Abnormal	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
				Neurontin	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Wellbutrin SS

Date:02/28/01ISR Number: 3677190-1Report Type:Periodic Company Report #A007753
 Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea	Consumer	Zoloft	PS	Pfizer	ORAL
100.00 MG		Drug Ineffective				Pharmaceuticals Inc	
TOTAL;BID;		Insomnia					
ORAL				Neurontin	SS		ORAL
600.00 MG							
TOTAL;DAILY;O							
RAL				Trazodone	C		

Date:02/28/01ISR Number: 3677336-5Report Type:Periodic Company Report #A035591
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Myalgia	Health Professional	Zoloft	PS	Pfizer	ORAL
75.00 MG						Pharmaceuticals Inc	
TOTAL;DAILY				Neurontin	SS		
				Insulin	C		
				Ambien	C		
				Estradiol	C		
				Synthroid	C		

Date:02/28/01ISR Number: 3677404-8Report Type:Periodic Company Report #A004325
 Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tachycardia	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
25.00 MG							
TOTAL:DAILY:O							
RAL				Adderall	SS		ORAL
ORAL				Neurontin	SS		ORAL
ORAL							

Date:02/28/01ISR Number: 3677522-4Report Type:Periodic Company Report #A001056
Age:13 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain Agitation	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
25.00 MG							
TOTAL: DAILY		Drug Ineffective					
ORAL		Dysgeusia					
300.00 MG		Hypertonia		Lithium	SS		ORAL
TOTAL:DAILY:O		Muscle Twitching					
RAL		Myalgia					
900.00 MG		Osteoarthritis		Neurontin	SS		ORAL
TOTAL:TID:ORA		Pain					
L		Thinking Abnormal					
1.00 MG		Vasodilatation		Haldol	SS		ORAL
TOTAL:BID:ORA							
L				Adderall	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tenex C
 Claritin C

Date:02/28/01ISR Number: 3677651-5Report Type:Periodic Company Report #A032315
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged Required	Agitation Apathy Drug Ineffective	Consumer	Zoloft Neurontin	PS SS	Pfizer Pharmaceuticals Inc	
Intervention to Prevent Permanent Impairment/Damage						

Date:02/28/01ISR Number: 3677680-1Report Type:Periodic Company Report #A037371
 Age:81 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Disability 25.00 MG TOTAL:DAILY:0 RAL	Hypoaesthesia Myasthenic Syndrome	Consumer	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
1800.00 MG TOTAL:DAILY:0 RAL			Neurontin	SS		ORAL

Date:02/28/01ISR Number: 3679439-8Report Type:Periodic Company Report #A031899
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other						
	Depersonalisation Speech Disorder	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	

Gabapentin SS
Medication
Unspecified C

Date:02/28/01ISR Number: 3679763-9Report Type:Periodic Company Report #A037782
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	
80.00 MG				Gabapentin	SS		
TOTAL:DAILY				Clonazepam	C		

Date:03/02/01ISR Number: 3672875-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100193
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence Feeling Abnormal	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
400 MG (100 MG, QID) SEE IMAGE				Unspecified Allergy/Sinus	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Celebrex (Celecoxib) C

Date:03/02/01ISR Number: 3673093-7Report Type:Expedited (15-DaCompany Report #049-0945-M0100016

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Extrapyramidal Disorder	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:03/05/01ISR Number: 3672613-6Report Type:Expedited (15-DaCompany Report #WAES 01027276

Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 7 DAY		Blood Alkaline		Vasotec	PS	Merck & Co., Inc	ORAL
Initial or Prolonged		Phosphatase Increased		Vioxx	SS		ORAL
		Blood Bilirubin Increased		Gabapentin	SS		ORAL
		Blood Lactate		Floxacillin Sodium	SS		ORAL
9 DAY		Dehydrogenase Increased		Leflunomide	SS		ORAL
		Blood Potassium Increased		Furosemide	SS		ORAL
7 DAY		Cardiomegaly		Metronidazole	SS		ORAL
9 DAY		Culture Urine Positive		Morphine	C		
		Depressed Level Of Consciousness		Vitamin E [Therapy	C		
		Electroencephalogram Abnormal		Unspecified]	C		
		Fatigue		Albuterol Sulfate And Ipratropium			
		Headache		Bromide	C		
		Infection		Calcium Carbonate			
		Liver Disorder		And Cholecalciferol	C		
		Pain		Loperamide	C		
		Pharyngolaryngeal Pain		Prednisolone	C		
		Proteus Infection		Acetaminophen	C		
		Renal Failure		Cisapride	C		
		Skin Ulcer		Cetirizine			
		Varicose Ulceration		Hydrochloride	C		
				Morphine	C		

Folic Acid C
 Lansoprazole C
 Zolpidem Tartrate C

Date:03/05/01ISR Number: 3673975-6Report Type:Expedited (15-DaCompany Report #358-0945-M0100001
 Age:75 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (400 MG, TID), PER ORAL	Asthenia Confusional State Difficulty In Walking Dizziness Drug Interaction Pneumonia	Foreign Health Professional	Neurontin Panacod (Codeine Phosphate Paracetamol) Diltiazem Hydrochloride Nizax (Nizatidine) Amitriptyline Hydrochloride Levofloxacin	PS C C C C C C	Parke Davis Pharmaceuticals Ltd	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tazoabactim Sodium C
 Klexane
 (Heparin-Fraction,
 Sodium Salt) C
 Lactulose C

Date:03/05/01ISR Number: 3673978-1Report Type:Expedited (15-DaCompany Report #061-0945-M0100004
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Circulatory Collapse Loss Of Consciousness Pulse Absent	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
1200 MG (600 MG, BID) , PER ORAL							

Date:03/05/01ISR Number: 3674648-6Report Type:Expedited (15-DaCompany Report #2001SE01042
 Age:69 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiac Failure Coma Overdose Respiratory Disorder	Foreign Health Professional Other	Pulmicort Allopurinol Choay Hemi-Daonil	PS SS SS	Astrazeneca Pharmaceuticals Lp	
2.5 MG BID 300 MG DAILY 400 MG TID							
Fonzylane SS							
Lasilix SS							
Neurontin SS							
Acuitel C							
Burinex C							
Monicor C							
Aspirine C							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Alkaline Phosphatase Increased Cardiomegaly	Foreign Other	Vasotec	PS	Merck Research Laboratories Div Merck Co Inc	ORAL
20 MG, PO	7	DAY		Tab Vioxx (Rofecoxib)	SS		ORAL
25 MG, PO				Tab Metronidazole	SS		ORAL
800 MG, PO	9	DAY		Cap Gabapentin	SS		ORAL
2400 MEQ, PO				Tab Furosemide	SS		ORAL
40 MG, PO	7	DAY		Tab Leflunomide	SS		ORAL
20 MG, PO				Tab Floxacillin Sodium	SS		ORAL
2200 MG, PO	9	DAY		[Therapy Unspecified]	C		
		Metabolic Disorder		Acetaminophen	C		
		Pharyngolaryngeal Pain		Albuterol Sulfate			
		Proteus Infection		(+) Ipratropium Bromid	C		
		Renal Failure		Calcium Carbonate			
		Urinary Tract Infection		(+) Cholecalciferol	C		
				Cetirizine Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Folic Acid C
 Lansoprazole C
 Loperamide Morphine C
 Prednisolone C
 Vitamin E C
 Zolpidem Tartrate C

Date:03/06/01ISR Number: 3675378-7Report Type:Expedited (15-DaCompany Report #049-0945-M0100012
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Coma	Foreign	Neurontin	PS	Parke Davis	
Other		Hypoglycaemia	Consumer Health Professional			Pharmaceuticals Ltd	

Date:03/06/01ISR Number: 3675381-7Report Type:Expedited (15-DaCompany Report #002-0945-M0100017
 Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hyperacusis	Foreign	Neurontin	PS	Parke Davis	ORAL
600 MG (300		Movement Disorder	Health			Pharmaceuticals Ltd	
MG, BID), PER		Poverty Of Speech	Professional				
ORAL				(Amitriptyline)	C		

Date:03/07/01ISR Number: 3675532-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100219
 Age:85 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Disorientation	Health	Neurontin			
Initial or Prolonged		Mental Impairment	Professional	(Gabapentin)	PS	Parke Davis	
3200-3600 MG		Musculoskeletal Stiffness				Pharmaceuticals Ltd	ORAL

Myalgia

(DAILY), PER

ORAL

Vicodin
(Paracetamol,
Hydrocodone
Bitartrate)

C

Date:03/07/01ISR Number: 3675614-7Report Type:Expedited (15-DaCompany Report #2001COU0301

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged PO	Drug Interaction International Normalised Ratio Decreased International Normalised	Consumer	Coumadin	PS	Dupont Merck Pharmaceutical Co	ORAL
0-300 MG	Ratio Increased Prothrombin Time Prolonged		Neurontin (Gabapentin)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/08/01ISR Number: 3676588-5Report Type:Expedited (15-DaCompany Report #049-0945-M0100013

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 400 (DAILY), Required PER ORAL	Difficulty In Walking Malaise Oedema Peripheral	Foreign Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Intervention to Prevent Permanent Impairment/Damage	Oral Intake Reduced Pain In Extremity					

Date:03/08/01ISR Number: 3676590-3Report Type:Expedited (15-DaCompany Report #358-0945-M0100001

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (400 MG, TID), PER ORAL	Asthenia Confusional State Difficulty In Walking Dizziness	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
PER ORAL	Drug Interaction Pneumonia		Amitriptyline Hydrochloride	SS		ORAL
PER ORAL			Panacod (Codeine Phosphate, Paracetamol)	SS		ORAL
			Diltiazem Hydrochloride	C		
			Nizax (Nizatidine)	C		
			Levofloxacin	C		
			Tazobactam Sodium	C		
			Klexane (Heparin-Fraction, Sodium Salt)	C		
			Lactulose	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200.00 MG :		Depression Oedema Peripheral Post-Traumatic Stress Disorder	Health Professional	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
BID : ORAL 1600.00 M G TOTAL: QID : ORAL				Neurontin	SS		ORAL
				Vicodin Colace Seroquel	C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG (100 MG, TID); 600 MG (200 MG, TID)		Facial Palsy	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				... Tylenol W/Codeine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Codeine Phosphate,
Paracetamol) C
Flexaril
(Cyclobenzaprine
Hydrochloride) C
Paxil (Paroxetine
Hydrochloride) C
Trazodone C

Date:03/09/01ISR Number: 3676428-4Report Type:Direct
Age: Gender: I/FU:I

Company Report #USP 081393

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Neurontin	PS	Parke-Davis	
				Noroxin	SS	Merck & Co	

Date:03/09/01ISR Number: 3678203-3Report Type:Expedited (15-DaCompany Report #044-0945-M0100053
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diabetes Mellitus	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Other		Diabetes Mellitus					
900 MG		Inadequate Control	Professional				
(DAILY), PER			Company				
ORAL			Representative				

Date:03/09/01ISR Number: 3678205-7Report Type:Expedited (15-DaCompany Report #044-0945-M0100052
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sepsis	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Other		Urinary Tract Infection					
1800 MG			Professional				
(DAILY), PER							

ORAL

Other

Tylex (Phenylephrine
 Hydrochloride,
 Paracetamol,
 Carbinoxamine
 Maleate) C
 (Naproxen) C
 (Temazepam) C
 Oramorph (Morphine
 Sulfate) C

Date:03/09/01ISR Number: 3678208-2Report Type:Expedited (15-DaCompany Report #044-0945-M0100051
 Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Sepsis	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
		Urinary Tract Infection					
900 MG			Professional				
(DAILY), PER			Other				

ORAL

(Amitriptyline) C
 Co-Proxamol
 (Paracetamol,
 Dextropropoxyphene
 Hydrochloride) C
 Frumil (Furosemide,

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Amiloride
 Hydrochloride) C
 Asa (Acetylsalicylic
 Acid) C
 Imdur (Isorbide
 Mononitrate) C
 (Nicorandil) C
 (Atenolol) C

Date:03/09/01ISR Number: 3678210-0Report Type:Expedited (15-DaCompany Report #044-0945-M0100029
 Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 100 MG		Drug Interaction - Narcolepsy	Foreign Health Professional Other	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
(DAILY), PER				Sinemet (Levodopa, Carbidopa) (Paroxetine)	SS SS		
ORAL							

Date:03/12/01ISR Number: 3680258-7Report Type:Expedited (15-DaCompany Report #001-0945-M0001041
 Age:14 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2400 MG (800 MG, TID)		Amnesia Anxiety Condition Aggravated	Health Professional	Neurontin	PS	Pfizer Pharmaceuticals	
		Convulsion Dizziness Mental Impairment Psychomotor Retardation Sedation Tearfulness Visual Acuity Reduced		Lamictal (Lamotrigine)	C		

Date:03/12/01ISR Number: 3680317-9Report Type:Expedited (15-DaCompany Report #S01-USA-00393-01
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	40 MG QD PO	Anorectal Disorder Colitis	Health Professional	Celexa	PS	Forest Laboratories Inc	ORAL
		Colitis Ulcerative Diarrhoea Haemorrhagic		Celexa (Citalopram Hydrobromide)	SS		ORAL
	30 MG QD PO	Haematocrit Decreased Haemoglobin Decreased Rectal Haemorrhage Sinusitis		Neurontin (Gabapentin) Risperdal (Risperidone)	SS SS		

Date:03/12/01ISR Number: 3681439-9Report Type:Expedited (15-DaCompany Report #055-0945-M0100003
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	400 MG (DAILY), PER	Condition Aggravated	Foreign Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
	ORAL						

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ORAL

450 MG

Clozapine (Clozapine)	SS
Levothyroxine (Levothyroxine)	C
Lorazepam	C
Zolpidem	C
Celecoxib (Celecoxib)	C

Date:03/19/01ISR Number: 3684352-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100041
 Age:31 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged SEE IMAGE, PER ORAL	Abnormal Behaviour Asthenia Blood Pressure Increased Dizziness Fatigue Feeling Jittery Malaise Medication Error Pain Palpitations Sedation Tremor	Consumer	Neurontin Ultram (Tramadol Hydrochloride)	PS C	Parke Davis Pharmaceuticals Ltd	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/19/01ISR Number: 3684500-8Report Type:Expedited (15-DaCompany Report #001-0945-M0100302

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other		Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:03/19/01ISR Number: 3684502-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100293

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged (DAILY), PER Other ORAL	Arterial Disorder - Chest Discomfort Myocardial Infarction Pain In Extremity	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:03/19/01ISR Number: 3684633-6Report Type:Expedited (15-DaCompany Report #033-0945-M0100008

Age:54 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Scintillating Scotoma	Foreign Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

SEE IMAGE

Date:03/20/01ISR Number: 3685701-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100041

Age:31 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (600 MG, BID), PER	Asthenia Blood Pressure Increased Dizziness	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

ORAL	Drug Withdrawal Syndrome				
(300 MG,	Fatigue		Neurontin		
BID), PER	Malaise		(Gabapentin) 600 Mg	SS	ORAL
	Pain				
ORAL	Palpitations				
300 MG, BID,	Tremor		Neurontin		
PER ORAL	Visual Disturbance		(Gabapentin) 600 Mg	SS	ORAL
			Neurontin		
PER ORAL			(Gabapentin)	SS	ORAL
			Ultram (Tramadol		
			Hydrochloride)	C	

Date:03/20/01ISR Number: 3685748-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100261
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Alkaline	Health	Neurontin	PS	Parke Davis	
3200 MG		Phosphatase Increased	Professional			Pharmaceuticals Ltd	
(DAILY)		Liver Function Test					
		Abnormal		(Unspecified Pain			
				Pills)	C		
				(Fluorouracil)	C		

Date:03/20/01ISR Number: 3685820-3Report Type:Expedited (15-DaCompany Report #001-0945-M0001325
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Asthenia	Health	Neurontin	PS	Parke Davis	
		Hypoaesthesia	Professional			Pharmaceuticals Ltd	
1600 MG (400							
MG ,4 TIMES							
DAILY) PER							
				Narcotic (Narcotic)	C		

Date:03/20/01ISR Number: 3686136-1Report Type:Expedited (15-DaCompany Report #044-0945-M0100069
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Post Procedural	Foreign	Neurontin	PS	Parke Davis	
		Complication	Health			Pharmaceuticals Ltd	
UNKNOWN			Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/20/01ISR Number: 3686137-3Report Type:Expedited (15-DaCompany Report #033-0945-M0100030

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	600 MG	Atrioventricular Block Cardiac Arrest Malaise Ventricular Fibrillation	Foreign Health Professional	Neurontin Fosphenytoin Sodium	PS SS	Parke Davis Pharmaceuticals Ltd	
(SINGLE DOSE),							
INTRAVENOUS							

(Carbamazepine) C
 (Clonazepam) C
 (Alprazolam) C
 (Clobazam) C
 (Valproate Sodium) C
 (Lamotrigine) C
 (Domperidone) C
 (Topiramate) C

Date:03/20/01ISR Number: 3686318-9Report Type:Expedited (15-DaCompany Report #044-0945-M0100067

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG, PER	Grand Mal Convulsion Necrotising Fasciitis	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Other		Sepsis					
ORAL							

Amitriptyline C
 Fluoxetine C
 Fentanyl C
 Paracetamol C
 Oxygen, Nitrous Oxide C
 Diclofenac Sodium C
 Potassium Chloride C
 Potassium Chloride C
 Diazepam C

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening	Coma	Foreign	Neurontin	PS	Parke Davis	
Other	Hypoglycaemia	Health			Pharmaceuticals Ltd	
900 MG		Professional	Hynorex Retard	C		
			Norvasc 5	C		
			(Lorazepam)	C		
			(Belladonna			
			Alkaloids)	C		
			(Ass)	C		
			(Risperidone)	C		
			Clavulante			
			Potassium,			
			Amoxicillin			
			Trihydrate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/20/01ISR Number: 3686375-XReport Type:Expedited (15-DaCompany Report #061-0945-M0100004

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Pulse Absent Syncope	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
300 MG (BID), PER ORAL				Citalopram	C		

Date:03/21/01ISR Number: 3687004-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100237

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other DAILY, PER ORAL		Dizziness Gastrointestinal Disorder Nausea	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
		Nervousness Tremor		Unknown Heart Medication	C		

Date:03/21/01ISR Number: 3687108-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100300

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (QID);2400 MG(QID)		Blood Albumin Decreased Cardiac Failure Congestive Pco2 Increased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Zostrix (Capsaicin) Morphine (Morphine) Glucophage (Metformin)	C C		

Hydrochloride) C
 Glucotrol
 (Glipizide) C
 Potassium
 (Potassium) C
 (Furosemide) C
 Lomotil (Atropine
 Sulfate,
 Dihphenoxyate
 Hydrochloride) C
 Ambien (Zolpidem
 Tartrate) C

Date:03/21/01ISR Number: 3687339-2Report Type:Expedited (15-DaCompany Report #001-0073-M0100097
 Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Drug Ineffective	Health Professional	Dilantin	PS	Parke Davis Div Warner Lambert Co	ORAL
400 MG DAILY PER ORAL		Medication Error					
				Neurontin (Gabapentin)	SS		ORAL
1100 MG DAILY PER ORAL							
				Clozapine (Clozapine)	SS		
450 MG				Levothyroxine			

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchopneumonia	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Other		Metastases To Lung					
900 MG, PER		Pulmonary Function Test	Professional				
ORAL; 1800		Abnormal	Company				
MG, PER ORAL		Respiratory Arrest	Representative	Analgesia	C		
				Chemotherapy	C		
				Ramipril	C		
				Diamorphine	C		
				Temazepam	C		
				Midazolam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/26/01ISR Number: 3689728-9Report Type:Expedited (15-DaCompany Report #001-0945-M0001276

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	900 MG (TID), PR ORAL	Condition Aggravated Corneal Oedema Dizziness Eye Disorder Photophobia Sedation Vision Blurred	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
				Celebrex (Celecoxib)	C		
				Enbrel	C		
				Prilosec (Omeprazole)	C		
				Estratest (Methyltestosterone, Estrogens Esterfied)	C		
				Folic Acid	C		
				Methotrexate	C		
				Pindolol	C		

Date:03/26/01ISR Number: 3689865-9Report Type:Expedited (15-DaCompany Report #044-0945-M0100072

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged		Hallucination	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Amitriptyline (Amitriptyline)	C		
				Carbamazepine (Carbamazepine)	C		
				Opioids	C		

Date:03/26/01ISR Number: 3689925-2Report Type:Expedited (15-DaCompany Report #044-0945-M0100053

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG		Diabetes Mellitus Inadequate Control	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

(DAILY), PER

Professional

ORAL

Company

Representative

Insulin
Capsaicin
Solfadeine

C
C
C

Date:03/26/01ISR Number: 3690205-XReport Type:Expedited (15-DaCompany Report #PHRM2000FR01473

Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG/DAY,		Drug Interaction Grand Mal Convulsion	Foreign Health	Tegretol-Xr	PS	Novartis Pharmaceuticals Corp	ORAL
ORAL			Professional				
2400 MG/DAY,			Other	Neurontin (Gabapentin) Capsule	SS		ORAL
ORAL				Nutritional Supplements	SS		ORAL
ORAL				Urbanyl (Clobazam)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/27/01ISR Number: 3690667-8Report Type:Direct
Age:15 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged X 46 TABS Required ORAL	Convulsion Intentional Misuse Vomiting		Wellbutrin 150mg Glaxo-Wellcome	PS	Glaxo-Wellcome	ORAL
Intervention to Prevent Permanent X 50 TABS Impairment/Damage ORAL			Neurontin 300mg Parke-Davis	SS	Parke-Davis	ORAL
			Celexa	C		

Date:03/27/01ISR Number: 3690780-5Report Type:Expedited (15-DaCompany Report #049-0945-M0000031
Age:65 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 1600 MG Life-Threatening (DAILY), PER Hospitalization - ORAL Initial or Prolonged Other 10 MG, Required UNKNOWN	Abdominal Pain Arterial Disorder Bronchopneumonia Gangrene Hepatic Failure Liver Disorder	Foreign Health Professional Other	Neurontin	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage	Oral Intake Reduced Pancreatitis Necrotising Pulmonary Oedema Refusal Of Treatment By Patient Respiratory Failure Sepsis Thrombotic Microangiopathy Vomiting White Blood Cell Count Decreased		Beloc-Zok	C		

Date:03/28/01ISR Number: 3690399-6Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin 300mg	PS		
Other		Dermatitis		Depakote 500 Mg	SS		

Date:03/28/01ISR Number: 3691281-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000751
Age:70 YR Gender:Male I/FU:F

Outcome	PT
	Abdominal Pain
	Abdominal Tenderness
	Autoantibody Positive
	Benign Prostatic Hyperplasia
	Blister
	Ear Pain
	Electric Shock
	Eye Irritation
	Eye Pain
	Facial Pain
	Fatigue
	Feeling Abnormal

B12 (Cyanocobalamin)	C
Folate (Folate Sodium)	C
Zinc (Zinc)	C
Chromium (Chromium)	C
Saw Palmetto (Serenoa Repens)	C
Coenzyme (Q12)	C
Ginseng (Ginseng)	C
Ginko (Ginko Tree Leaves Extract)	C
Selenium (Selenium)	C
Glutathione (Glutathione)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/28/01ISR Number: 3691987-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100314

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Conversion Disorder Grand Mal Convulsion	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
1800 MG (600 MG, THREE TIMES DAILY) THREE TIMES DAILY (200 MG) 200 UNKNOWN (200 UNKNOWN, ONE EVERY MORNING)				Phenytoin	SS		
				Sertraline	SS		

Date:03/29/01ISR Number: 3692443-9Report Type:Expedited (15-DaCompany Report #A105990

Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 200.00 MG Prevent Permanent TOTAL:DAILY:O Impairment/Damage RAL		Hypertrophy Breast Weight Increased	Consumer	Zoloft	PS	Pfizer Pharmaceuticals Inc	ORAL
				Neurontin	SS		
				Lipitor	SS		
				Celebrex	SS		
				Clonazepam	C		
				Desipramine	C		
				Seroquel	C		

Date:03/29/01ISR Number: 3692446-4Report Type:Expedited (15-DaCompany Report #001-0945-M0001276
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Condition Aggravated Corneal Oedema	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
900 MG (TID), PER ORAL		Dizziness Eye Disorder Photosensitivity Reaction Sedation Vision Blurred		Celebrex (Celecoxib) Enbrel Prilosec (Omeprazole) Estratest (Methyltestosterone, Estrogens Esterified) Folic Acid Methotrexate Pindolol	C C C C C C C C		

Date:03/29/01ISR Number: 3692727-4Report Type:Expedited (15-DaCompany Report #044-0945-M0100072
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death Hospitalization - Initial or Prolonged		Hallucination	Foreign	Neurontin Amitriptyline Carbamazepine Opioids	PS C C C	Parke Davis Pharmaceuticals Ltd	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/01ISR Number: 3692768-7Report Type:Expedited (15-DaCompany Report #047-0945-M0100002

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Congenital Anomaly		Benign Congenital Hypotonia	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
2400 MG,							
PLACENTAL		Cleft Lip	Professional				
40 MG,		Cleft Palate	Other	Frisium (Clobazam)	SS		
PLACENTAL		Complications Of Maternal					
		Exposure To Therapeutic Drugs		Rivotril (Clonazepam)	C		

Date:03/29/01ISR Number: 3692769-9Report Type:Expedited (15-DaCompany Report #055-0945-M0100008

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Pneumonia	Foreign	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
Other		Throat Cancer	Consumer				
800 MG							
(DAILY)							

Date:04/02/01ISR Number: 3693514-3Report Type:Direct Company Report #USP 081400

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Neurontin	PS	Parke-Davis	
				Neurontin	SS	Parke-Davis	

Date:04/02/01ISR Number: 3698078-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100322

Age:5 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Aggression	Consumer	Neurontin	PS	Parke Davis	ORAL
600 MG	Agitation				Pharmaceuticals Ltd	
(DAILY), PER	Condition Aggravated					
ORAL	Disturbance In Social					
	Behaviour		Levothyroxine			
	Drug Effect Decreased		(Levothyroxine)	C		
	Emotional Disorder					

Date:04/03/01ISR Number: 3695126-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100336
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Confusional State	Consumer	Neurontin	PS	Parke Davis	ORAL
200 MG (100		Difficulty In Walking				Pharmaceuticals Ltd	
MG, PER ORAL		Speech Disorder					
				Duragesic (Fentanyl)	SS		
				Vicodin			
				(Paracetamol,			
				Hydrocodone			
				Bitartrate)	SS		
400MG (DAILY)				Neurontin	SS		
PO							
				Lipitor			
				(Atorvastatin)	C		
				Premarin (Estrogens			
				Conjugated)	C		
				Detrol (Tolterodine			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tartrate) C
 Vioxx (Rofecoxib) C
 Sinemet (Levodopa,
 Carbidopa) C

Date:04/03/01ISR Number: 3695127-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100365
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Upper Oedema Peripheral	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
300 MG (TID), PER ORAL		Pain In Extremity					

Levbid (Hyoscyamine
 Sulfate) C
 Coumadin (Warfarin
 Sodium) C
 Elavil
 (Amitriptyline
 Hydrochloride) C
 Allegra D
 (Pseudoephedrine,
 Fexofenadine
 Hydrochloride) C

Date:04/03/01ISR Number: 3695193-8Report Type:Expedited (15-DaCompany Report #001-0945-M0100363
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Breast Cancer Female Mania	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
1800 MG TID							

Diazepam C
 Nefazodone C
 Oxycodone
 Hydrochloride C
 Hydroxychloroquine
 Sulfate C
 Celecoxib C

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 150.00 MG Required TOTAL DAILY Intervention to 400.00 MG Prevent Permanent TOTAL BID Impairment/Damage 375.00 MG TOTAL BID	Blood Pressure Decreased Decreased Appetite Dehydration Depression Disorientation Dizziness Fall Insomnia Syncope	Health Professional	Zoloft Neurontin Serzone Proamatine Ambien Blood Pressure Medication Klonopin	PS SS SS SS C C C	Pfizer Pharmaceuticals Inc	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/03/01ISR Number: 3695691-7Report Type:Expedited (15-DaCompany Report #044-0945M0100051

Age:84 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG Other (DAILY), PER ORAL	Sepsis Urinary Tract Infection	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Amitriptyline	C
Co-Proxamol (Paracetamol, Dextropropoxyphene Hydrochloride)	C
Frumil (Furosemide)	C
Frumil (Furosemide, Amiloride Hydrochloride)	C
Asa (Acetylsalicylic Acid)	C
Imdur (Isosorbide Mononitrate)	C
Nicorandil	C
Atenolol	C

Date:04/03/01ISR Number: 3695711-XReport Type:Expedited (15-DaCompany Report #044-0945-M0100052

Age:72 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG Other (DAILY), PER ORAL	Sepsis Urinary Tract Infection	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Tylox (Phenylephrine Hydrochloride, Paracetamol, Carbinoxamine Maleate)	C
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Naproxen C
 Temazepam C
 Oramorph (Morphine Sulfate) C

Date:04/03/01ISR Number: 3696066-7Report Type:Expedited (15-DaCompany Report #2001SIN0055
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged "1000 MG" DLY		Drug Interaction Narcolepsy	Foreign Health Professional	Sinemet	PS	Dupont Pharmaceuticals Co	ORAL
PO 100 MG QHS PO			Other	Neurontin (Gabapentin)	SS		ORAL
20 MG DLY PO				Seroxat "Smith Kline Beecham" (Paroxetine Hydrochloride)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/03/01ISR Number: 3696166-1Report Type:Expedited (15-DaCompany Report #044-0945-M0100117

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Dermatitis Pruritus	Foreign Health Professional	Neurontin Oxycodone	PS C	Parke Davis Pharmaceuticals Ltd	ORAL

Date:04/03/01ISR Number: 3696878-XReport Type:Expedited (15-DaCompany Report #001-0945-M0100326

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (TID) , PER ORAL		Condition Aggravated Confusional State	Health Professional	Neurontin Unspecified Chemotherapy Agent (Oxycodone Hydrochloride)	PS C C	Parke Davis Pharmaceuticals Ltd	ORAL

Date:04/03/01ISR Number: 3696883-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100327

Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (BID), PER ORAL		Diabetes Mellitus	Consumer	Neurontin Tenormin (Atenolol) Accupril (Quinapril Hydrochloride) Megace (Megestrol Acetate) Darvocet (Paracetamol, Dextropropoxyphene)	PS C C C C	Parke Davis Pharmaceuticals Ltd	ORAL

Dalmane (Flurazepam
Hydrochloride) C

Date:04/04/01ISR Number: 3698070-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100345

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Chondropathy Hypertrophy Breast Knee Arthroplasty	Consumer	Neurontin (Sertraline)	PS SS	Parke Davis Pharmaceuticals Ltd	ORAL
200 MG (DAILY), PER ORAL		Weight Increased					

Lipitor
(Atorvastatin
Calcium) SS
Celecoxib
(Celecoxib) SS
Clonazepam
(Clonazepam) SS
Desipramine
(Desipramine) SS
Quetiapine
(Quetiapine) SS

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/04/01ISR Number: 3698083-XReport Type:Expedited (15-DaCompany Report #001-0945-M0100328

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Abdominal Pain Upper Dyspepsia	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1000 MG		Dyspnoea					
(DAILY), PER		Flatulence					
ORAL		Influenza		Trazodone			
		Muscle Spasms		(Trazodone)	C		
		Myalgia		Diphenhydramine			
		Pain		(Diphenhydramine)	C		
				Zolpidem (Zolpidem)	C		
				Buspirone			
				(Buspirone)	C		

Date:04/04/01ISR Number: 3698211-6Report Type:Expedited (15-DaCompany Report #001-0073-M0100016

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Condition Aggravated Visual Acuity Reduced	Consumer Health	Dilantin	PS	Parke Davis Div Warner Lambert Co	ORAL
500 MG, PER			Professional				
ORAL				Gabapentin			
				(Gabapentin)	SS		
900 MG (THREE							
TIMES A DAY)							

Date:04/06/01ISR Number: 3700189-3Report Type:Expedited (15-DaCompany Report #S01-USA-00393-01

Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abnormal Faeces	Health	Celexa	PS	Forest Laboratories	

Initial or Prolonged	Colitis Ulcerative	Professional		Inc	ORAL
40 MG QD PO					
	Dehydration		Celexa (Citalopram		
	Diarrhoea Haemorrhagic		Hydrobromide)	SS	ORAL
30 MG QD PO					
	Haematocrit Decreased		Neurontin		
	Haemoglobin Decreased		(Gabapentin)	SS	
	Haemorrhage		Risperdal		
	Pain		(Risperidone)	SS	
	Rectal Haemorrhage		Asacol (Mesalazine)	SS	
	Sinusitis		Prednisone	SS	
	Vomiting				

Date:04/06/01ISR Number: 3700210-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100354
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Parkinson'S Disease	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

300 MG
(DAILY) PER
ORAL

Date:04/06/01ISR Number: 3700215-1Report Type:Expedited (15-DaCompany Report #002-0945-M0100044
Age: Gender:Female I/FU:I

Outcome	PT	Report Source
Hospitalization -	Dizziness	Foreign
Initial or Prolonged	Mood Altered	Health

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
1100 MG (TWICE DAILY); PER ORAL		Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:04/06/01ISR Number: 3700447-2Report Type:Expedited (15-DaCompany Report #044-0945-M0100090
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN		Cardiac Failure Hypoglycaemia	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
		Lower Respiratory Tract Infection Transient Ischaemic Attack	Professional				

Date:04/06/01ISR Number: 3700448-4Report Type:Expedited (15-DaCompany Report #044-0945-M0100088
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN		Hypoglycaemia	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
			Professional				

Date:04/06/01ISR Number: 3718580-8Report Type:Periodic Company Report #2000036882US
Age:58 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4 MG, ORAL		Alopecia Drug Interaction	Consumer	Detrol Neurontin (Gabapentin)	PS SS	Pharmacia And Upjohn Co	ORAL

Date:04/09/01ISR Number: 3701075-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100366
Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (TID)		Dysphagia Vomiting	Health Professional	Neurontin Clonidine	PS C	Parke Davis Pharmaceuticals Ltd	

Date:04/09/01ISR Number: 3701084-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100367
Age: Gender:I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 7200 MG (TID)		Unevaluable Event	Health Professional	Neurontin Neurontin(Gabapentin)	PS SS	Parke Davis Pharmaceuticals Ltd	
9600 MG (QID)							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/09/01ISR Number: 3701116-5Report Type:Expedited (15-DaCompany Report #031-0945-M0100007

Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Condition Aggravated Dyspnoea Lung Disorder	Foreign Consumer Health Professional	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
900 MG (TID), PER ORAL			Other Pulmonal Medication	C		

Date:04/09/01ISR Number: 3701432-7Report Type:Expedited (15-DaCompany Report #046-0945-M0100016

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Coagulopathy Drug Interaction	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
900 MG (TID), Other PER ORAL	Pain In Extremity		Warfarin (Warfarin Sodium)	SS		

Date:04/10/01ISR Number: 3702446-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100390

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Cardiac Failure Congestive	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
900 MG (TID) Other	Disorientation Fluid Retention Renal Failure		Dyazide (Hydrochloroathiazid e, Triamterene)	C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypertension Nephrotic Syndrome	Health Professional	Neurontin (Valproate Semisodium) (Clonidine)	PS C C	Parke Davis Pharmaceuticals Ltd	

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysarthria	Consumer	Oxycontin	PS	Purdue Pharma Lp	ORAL
PO		Encephalopathy	Health Professional Other	Neurontin (Gabapentin) Urecholine(Bethanech ol Chloride) Percocet Robaxin (Methocarbamol) Paxil (Paroxetine) Betaseron Baclofen (Lioresal)	SS SS C C C C C		

FDA - Adverse Event Reporting System (AERS)

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Neurontin	
(Gabapentin)	C
Celexa(Citalopram	
Hydrobromide)	C
Valium (Diazepam)	C
Urecholine	
(Bethanechol	
Chloride)	C
Verapamil Hcl	C
Trazodone Hcl	C
Imodium	C
Prilosec	
(Omeprazole)	C

Date:04/10/01ISR Number: 3702748-0Report Type:Expedited (15-DaCompany Report #049-0945-M0100013
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Difficulty In Walking	Foreign	Neurontin	PS	Parke Davis	
Initial or Prolonged	Malaise	Consumer			Pharmaceuticals Ltd	ORAL
400 MG						
Required	Oedema Peripheral					
(DAILY), PER						
Intervention to	Oral Intake Reduced					
ORAL						
Prevent Permanent	Pain In Extremity					
Impairment/Damage						

Date:04/10/01ISR Number: 3703611-1Report Type:Expedited (15-DaCompany Report #002-0945-M0100045
 Age:17 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death	Drug Interaction	Foreign	Neurontin	PS	Parke Davis	
	Drug Level Above	Health			Pharmaceuticals Ltd	ORAL
PER ORAL						
	Therapeutic	Professional	Zyprexia			
			(Olanzapine)	SS		
			Hydromorphone(Hydrom			
			orphone)	SS		

Date:04/10/01ISR Number: 3703612-3Report Type:Expedited (15-DaCompany Report #002-0945-M0100047
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma	Foreign	Neurontin	PS	Parke Davis	
		Drug Interaction	Health			Pharmaceuticals Ltd	
		Drug Level Above	Professional	Oxycodone (Oxycodone)	SS		
		Therapeutic		Clonazepam			
		Overdose		(Clonazepam)	SS		
		Respiratory Depression					

Date:04/11/01ISR Number: 3703250-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100398
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Blood Pressure Increased Emotional Disorder	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Other		Hypoaesthesia Palpitations		Fluphenazine (Fluphenazine) Alprazolam (Alprazolam)`	C C		

Freedom Of Information (FOI) Report

Levothyroxine
 (Levothyroxine) C
 Conjugated
 Estrogen(Medroxyprog
 esterone Acetate,
 Estrogens
 Conjugated) C
 Aspirin
 (Acetylsalicylic
 Acid) C
 Soy C
 Acetaminophen W/
 Codeine (Codeine,
 Paracetamol) C
 Rolaid
 (Dyhydroxyaluminum
 Sodium Carbonate) C
 Temazepam(Temazepam) C
 Albuterol
 (Salbutamol) C

Date:04/11/01ISR Number: 3703742-6Report Type:Expedited (15-DaCompany Report #200112306EU
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	800 MG/DAY PO 9 DAY	Blood Alkaline Phosphatase Increased	Foreign Other	Noritrate	PS	Dermik Laboratories Inc	ORAL
	20 MG /D AY	Cardiomegaly		Leflunomide (Arava)	SS		ORAL
	PO	Chest Pain					
	2.4 G /D AY	Depressed Level Of Consciousness		Gabapentin (Neurontin)	SS		
	40 MG/DAY PO	Ear Pain		Furosemide (Furix)	SS		ORAL
	20 MG/ DAY PO 1 WK	Electroencephalogram Abnormal		Enalapril Maleate (Renitec)	SS		ORAL
	2.2 G / DAY 9 DAY	Fatigue Gamma-Glutamyltransferase		Flucloxacillin Sodium (Heracillin)	SS		

Increased

Headache
 Hepatic Enzyme Increased
 Hypocalcaemia
 Infected Skin Ulcer
 Pharyngolaryngeal Pain
 Pleural Effusion
 Proteus Infection
 Renal Failure
 Weight Decreased

Rofecoxib (Vioxx) SS
 Folic Acid, Calcium Phosphate (Folacin) C
 Vitamin E C
 Paracetamol
 (Alvedon) C
 Prednisone C
 Morfin C
 Cisapride
 (Prepulsid) C
 Cetirizine (Zyrlex) C
 Lansoprazole (Lanzo) C
 Zolpidem Tartrate
 (Stilnoct) C
 Loperamide C
 Colecalciferol,
 Calcium Carbonate
 (Calcichew D3) C
 Ipratropium Bromide,
 Salbutamol Sulfate
 (Combivent) C
 Spironolactone C

ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/11/01ISR Number: 3714040-9Report Type:Periodic Company Report #2001-BP-00558
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness Fatigue	Consumer	Flomax	PS	Boehringer Ingelheim Pharmaceuticals Inc	ORAL
0.4 MG/1							
CAPSULES/QD/P							
O				Neurontin	SS		ORAL
600 MG/PO				Aspirin	C		

Date:04/12/01ISR Number: 3704357-6Report Type:Expedited (15-DaCompany Report #044-0945-M0100069
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Post Procedural Complication	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:04/13/01ISR Number: 3704672-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100410
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG		Depression Mood Swings	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
(DAILY)		Uterine Leiomyoma					
150 MG				Zoloft (Sertraline)	SS		
(DAILY); 200							
MG (DAILY)							

Date:04/13/01ISR Number: 3704705-7Report Type:Expedited (15-DaCompany Report #044-0945-M0100130
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Optic Neuritis	Foreign Other	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
PER ORAL				Dosulepin	C		
				Paracetamol	C		
				Diclofenac Sodium	C		

Date:04/13/01ISR Number: 3704707-0Report Type:Expedited (15-DaCompany Report #044-0945-M0100131
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Corneal Disorder Visual Field Defect	Foreign Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Epanuon (Phenytoin)	SS		
				Prednisolone (Prednisolone)	C		
				Cipramil (Citalopram Hydrobromide)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/13/01ISR Number: 3705117-2Report Type:Expedited (15-DaCompany Report #A107350
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 150.00 MG		Depression Uterine Leiomyoma	Consumer	Zoloft	PS	Pfizer Pharmaceuticals Inc	
TOTAL:DAILY				Gabapentin	SS		
600.00 MG							
TOTAL:DAILY							

Date:04/13/01ISR Number: 3705196-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100413
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG (TID), PER ORAL		Grand Mal Convulsion Rib Fracture	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:04/13/01ISR Number: 3705203-7Report Type:Expedited (15-DaCompany Report #055-0945-M0100008
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 800 MG Initial or Prolonged (DAILY) Other		Cerebrovascular Accident Pneumonia Throat Cancer	Foreign Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other	Convulsion Visual Disturbance	Health Professional	Diamox Depakote (Valproate Semisodium) Neurontin (Gabapentin) Imitrex (Sumatriptan Succinate) Loratadine (Loratadine) Fluticansoe Propionate (Fluticasone Propionate) Montelukast (Montelukast)	PS SS SS C C C C	Lederle Laboratories Div American Cyanamid Co	

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Suicide Attempt		Ativan Clonopine Prozac Trazodone	PS SS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Synthroid SS
 Gabapentin SS
 "Silver Bullet" Otc SS

Date:04/19/01ISR Number: 3708150-XReport Type:Expedited (15-DaCompany Report #001-0945-M0100442
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bleeding Time Prolonged Ecchymosis	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
SEE IMAGE		Epistaxis Platelet Adhesiveness Abnormal					

Date:04/19/01ISR Number: 3708319-4Report Type:Expedited (15-DaCompany Report #061-0945-M0100019
 Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - Initial or Prolonged INTRAVENOUS		Dementia Pneumonitis	Foreign Health Professional	Neurontin Voriconazole	PS SS	Parke Davis Pharmaceuticals Ltd	
				Ambisome (Amphotericine B, Liposome) Effexor (Fenlaxine Hydrochloride)	SS SS		

Date:04/19/01ISR Number: 3708320-0Report Type:Expedited (15-DaCompany Report #044-0945-M0100084
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression Depression	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
400 MG PER		Paranoia	Professional	Non-Opiate Pain Killers	C		

Date:04/20/01ISR Number: 3708654-XReport Type:Expedited (15-DaCompany Report #044-0945-M0100136
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypoglycaemia	Foreign Other	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
2.4 GM, PER							
ORAL				(Morphine Sulfate)	C		

Date:04/20/01ISR Number: 3708658-7Report Type:Expedited (15-DaCompany Report #033-0945-M0100052
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Bradycardia Haematemesis	Foreign Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
800 MG (BID),							
PER ORAL		Hypotension	Other				
		Melaena Ulcer		Feldene (Piroxicam) Aspegic (Acetylsalicylate Lysine)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lioresal (Baclofen) C
 Sotalol (Sotalol Hydrochloride) C
 Chronodalate (Nifedipine) C

Date:04/20/01ISR Number: 3708885-9Report Type:Expedited (15-DaCompany Report #044-0945-M0100070
 Age:89 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Neutropenia	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Other	600 MG, PER		Professional				
ORAL							

Frumil (Furosemide Amiloride) C
 Bumetanide (Bumetanide) C
 Lansoprazole (Lansoprazole) C
 Warfarin (Warfarin) C

Date:04/20/01ISR Number: 3708895-1Report Type:Expedited (15-DaCompany Report #200110296BFR
 Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG DAILY		Chorea	Foreign	Precose	PS	Bayer Corp	
Initial or Prolonged YR		Difficulty In Walking	Other	Ikorel	SS		
YR		Dysphagia		Sectral	SS		
YR				Furosemide	SS		
YR				Vastarel "Biopharma"	SS	Biopharma	
				Neurontin	SS		
				Amlor	C		
				Kardegic	C		
				Nitriderm Tts	C		

Date:04/24/01ISR Number: 3710542-XReport Type:Expedited (15-DaCompany Report #NSADSS2001011239
Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Colectomy Total	Health	Risperdal	PS	Janssen Research Fdn	
Initial or Prolonged	Colitis Ulcerative	Professional	Celexa (Citalopram			
Required	Dehydration		Hydrobromide)	SS		ORAL
SEE IMAGE						
Intervention to	Haematocrit Decreased		Neurontin			
Prevent Permanent	Haemoglobin Decreased		(Gabapentin)	SS		
Impairment/Damage	Laboratory Test Abnormal					
	Sinusitis					
	Vomiting					

Date:04/24/01ISR Number: 3710807-1Report Type:Expedited (15-DaCompany Report #PHBS2001NO03843
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Complications Of Maternal	Foreign	Tegretol	PS	Novartis	
Initial or Prolonged	Exposure To Therapeutic	Health			Pharmaceuticals Corp	
TRANSPLACENTAL	TRANSPLACENTA					
	Drugs	Professional				
L						
	Exomphalos	Other	Neurontin			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Gabapentin) SS

TRANSPLACENTAL TRANSPLACENTA

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Date:04/25/01ISR Number: 3710837-XReport Type:Direct
Age:70 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Sedation		Gabapentin	PS		

Date:04/26/01ISR Number: 3711898-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100479
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Suicidal Ideation	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

1200 MG DAILY
Other
PER ORAL;

2400 MG DAILY

PER ORAL

Ibuprofen
(Ibuprofen) C
Lorazepam
(Lorazepam) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C

Date:04/26/01ISR Number: 3711901-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100486
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Colitis Ulcerative	Health	Neurontin	PS	Parke Davis	

Initial or Prolonged UNKNOWN	Dehydration	Professional		Pharmaceuticals Ltd
Other Required DAILY PER	Diarrhoea Haemorrhagic Haematocrit Decreased		Celexa (Citalopram Hydrobromide)	SS ORAL
Intervention to ORAL	Haemoglobin Decreased			
Prevent Permanent Impairment/Damage UNKNOWN	Pain Rectal Haemorrhage		Risperdal (Risperidone)	SS
UNKNOWN	Vomiting		Asacol (Mesalazine)	SS
UNKNOWN			Prednisone (Prednisone)	SS
UNKNOWN			Solu-Medrol (Methylprednisolone Sodium Succinate)	SS
UNKNOWN			Cyclosporine	SS

Date:04/26/01ISR Number: 3712600-2Report Type:Periodic Company Report #A105423
 Age:56 YR Gender:Male I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Back Pain Drug Ineffective	Consumer	Viagra	PS	Pfizer Agricultural Div	ORAL
50 MG TOTAL PRN ORAL		Pain					
600.00 MG TOTAL, BID, ORAL		Sedation		Neurontin	SS		ORAL
				Baby Aspirin Vitamin E	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Folic Acid C
One A Day C

Date:04/27/01ISR Number: 3715572-XReport Type:Periodic Company Report #001-0945-M0000251
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dysgeusia Glossodynia	Consumer	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
1200 MG (BID), PER ORAL							

Toprol (Metoprolol
Succinate) C

Date:04/27/01ISR Number: 3715573-1Report Type:Periodic Company Report #001-0945-M0100403
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Hyperacusis	Consumer	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
600 MG (BID), PER ORAL							

Temazepam C

Date:04/27/01ISR Number: 3715574-3Report Type:Periodic Company Report #001-0945-M0100203
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Nausea	Consumer	Neurontin	PS	Parke Davis Pharmaceutical	

1200 MG (600

MG, BID), PER

ORAL

Date:04/27/01ISR Number: 3715576-7Report Type:Periodic Company Report #001-0945-M0100178
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Withdrawal Syndrome Intentional Misuse	Consumer	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL

7000 MG

(DAILY), PER

ORAL

Date:04/27/01ISR Number: 3715577-9Report Type:Periodic Company Report #001-0945-M0100113
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Herpes Zoster		Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lambert Co

Date:04/30/01ISR Number: 3714079-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100487
Age:34 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 9000 MG	Balance Disorder Blood Pressure Increased	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
	Depression Dysphagia Nausea		Valium (Diazepam) Extra Strength Tylenol Pm	SS SS		ORAL
25-30 (HS) , PER ORAL	Overdose Renal Failure Suicidal Ideation Tremor		Unspecified Antidepressants	C		

Date:04/30/01ISR Number: 3714180-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100502
Age:47 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG (300 MG,BID) PER ORAL	Asthenia Blood Potassium Decreased Blood Sodium Decreased Convulsion	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
600 MG (300 MG, BID) PER ORAL	Dizziness Muscle Spasms Weight Increased		Neurontin (Gabapentin)	SS		ORAL
			Depakote (Valproate Semisodium) Mysoline (Primidone) Klonopine(Klonazepam)	C C C		

Date:04/30/01ISR Number: 3714350-5Report Type:Expedited (15-DaCompany Report #039-0945-M0100003
Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - UNKNOWN, PER Initial or Prolonged ORAL		Bradycardia Cardiac Pacemaker Insertion	Foreign Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Date:04/30/01ISR Number: 3714402-XReport Type:Expedited (15-DaCompany Report #HQ0164626APR2001
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hypotension Hypothermia	Health Professional Other	Artane	PS	Lederle Laboratories Div American Cyanamid Co	ORAL
5 MG 1XPER 1 DAY ORAL				Depakine (Valproate Sodium)	SS		ORAL
1000 MG 1X PER 1 DAY ORAL				Haldol (Haloperidol)	SS		ORAL
1 MG 1XPER 1 DAY ORAL				Neurontin (Gabapantin)	SS		ORAL
1600 MG 1 X							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

PER 1 DAY

ORAL

Tranxene
(Clorazepate
Dipotassium)

SS

ORAL

ORAL

Date:05/01/01ISR Number: 3714483-3Report Type:Direct
Age:86 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Gabapentin	PS		

Date:05/01/01ISR Number: 3714688-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100134
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nephritis Interstitial	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Unspecified Antibiotic	C		

Date:05/01/01ISR Number: 3714695-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000724
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Tooth Discolouration	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
50 MG							
Other (DAILY),							

UNKNOWN

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG (300 MG, TID), PER ORAL	Drug Ineffective Dysphemia Hypoxia Mental Impairment	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Lasix (Furosemide)	C
Seroquel (Quetiapine)	C
Depakote (Valproate Semisodium)	C
Colace (Docusate Sodium)	C
Protonix (Pantoprazole)	C
K-Dur (Potassium Chloride)	C
Ferrous Sulfate (Ferrous Sulfate)	C
Thorazine (Chlorpromazine Hydrochloride)	C
Prozac (Fluoxetine Hydrochloride)	C
Cogentin (Benzatropine	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Mesilate0 C
 Oxygen (Oxygen) C

Date:05/01/01ISR Number: 3715162-9Report Type:Expedited (15-DaCompany Report #047-0945-M0100003
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
PLACENTAL		Drugs Umbilical Hernia	Professional Other	Tegretol (Carbamazepine)	SS		
PLACENTAL							

Date:05/01/01ISR Number: 3722785-XReport Type:Periodic Company Report #2001049457US
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Weight Increased	Consumer	Celebrex	PS	Gd Searle And Co	ORAL
ORAL				Clonazepam	SS		ORAL
ORAL				Desipramine	SS		ORAL
ORAL				Seroquel	SS		ORAL
ORAL				Zoloft	SS		ORAL
200MG. QD.							
ORAL				Neurontin	SS		ORAL
ORAL				Lipitor(Atorvastatin)	C		

Date:05/03/01ISR Number: 3716959-1Report Type:Expedited (15-DaCompany Report #055-0945-M0100012
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Cerebrovascular Accident Foreign Consumer Neurontin PS Parke Davis Pharmaceuticals Ltd ORAL
 400 MG
 (DAILY), PER
 ORAL

Date:05/03/01ISR Number: 3716974-8Report Type:Expedited (15-DaCompany Report #055-0945-M0100013
 Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 800 MG (DAILY), PER ORAL		Blood Pressure Increased Cough Dyspnoea	Foreign Consumer	Neurontin (Insulin)	PS C	Parke Davis Pharmaceuticals Ltd	ORAL

Date:05/03/01ISR Number: 3716980-3Report Type:Expedited (15-DaCompany Report #049-0945-M0100044
 Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 300 MG, PER ORAL		Angioneurotic Oedema Dyspnoea Renal Failure Stevens-Johnson Syndrome	Foreign Consumer Other	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/03/01ISR Number: 3717418-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100522

Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Priapism	Literature Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
900 MG (DAILY)			Professional				
6 MG (DAILY)				Risperidone (Risperidone)	SS		
7.5 MG (DAILY)				Olanzapine (Olanzapine)	SS		
300 MG (DAILY)				Fluvoxamine (Fluvoxamine)	SS		
25 MG (DAILY)				Oxazepam (Oxazepam)	SS		

Date:05/03/01ISR Number: 3717419-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100390

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 900 MG Initial or Prolonged (TID);300 MG (100 MG, TID)		Cardiac Failure Congestive Condition Aggravated Disorientation Fluid Retention Renal Failure	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
				Nitrostat (Glyceryl Trinitrate)	C		
				Lasix (Furosemide)	C		
				Toprol-Xl (Metoprolol Succinate)	C		
				K-Dur (Potassium			

Chloride) C
 Allopurinol C
 (Allopurinol)
 Bufferin
 (Acetylsalicylic
 Acid, Magnesium
 Carbonate, Aluminium
 Glycinate) C
 Paxil (Paroxetine
 Hydrochloride) C
 Zaroxolyn
 (Metolazone) C
 Digoxin (Digoxin) C

Date:05/04/01ISR Number: 3717938-0Report Type:Expedited (15-DaCompany Report #002-0945-M0100056
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Facial Palsy Headache	Foreign Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
PER ORAL							

Date:05/04/01ISR Number: 3718208-7Report Type:Expedited (15-DaCompany Report #044-0945-M0100051
 Age:84 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

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG		Sepsis Urinary Tract Infection	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
(DAILY), PER			Professional				
ORAL			Other				
				Amitriptyline	C		
				Co-Proxamol (Paracetamol, Dextropropoxyphene Hydrochloride)	C		
				Frumil (Furosemide, Amiloride Hydrochloride)	C		
				Asa (Acetylsalicylic Acid)	C		
				Imdur (Isosorbide Mononitrate)	C		
				Nicorandil	C		
				Atenolol	C		

Date:05/04/01ISR Number: 3718351-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100513
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Malaise	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	

Date:05/04/01ISR Number: 3718352-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100514
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (TID)		Cardiac Failure Congestive	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL

Other
PER ORAL

Dizziness

Dysuria
Fall
Head Banging
Malaise
Memory Impairment
Myocardial Infarction
Renal Failure
Syncope
Urinary Incontinence

Lasix (Furosemide) C
Thyroid (Thyroid) C
Premarin (Estrogens
Conjugated) C
Allopurinol
(Allopurinol) C
Flexeril
(Cyclobenzaprine
Hydrochloride) C
Prilosec
(Omeprazole) C
Potassium
(Potassium) C
Duragesic Patches C

Date:05/04/01ISR Number: 3718395-0Report Type:Expedited (15-DaCompany Report #001-0945-M0100197
Age:43 YR Gender:Male I/FU:F

Outcome PT
Other Anxiety
Condition Aggravated
Dizziness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Dependence Drug Withdrawal Syndrome Medication Error	Report Source	Product	Role	Manufacturer	Route
SEE IMAGE		Nausea Pain	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
		Pruritus		Prozac (Fluoxetine Hydrochloride)	C		

Date:05/04/01ISR Number: 3718677-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100228
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 900 MG, (300 Prevent Permanent MG, TID) Impairment/Damage		Coordination Abnormal Difficulty In Walking	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
		Muscular Weakness		Celebrex (Celecoxib) Vitamin C (Ascorbin Acid)	C C		

Date:05/07/01ISR Number: 3719036-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100525
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly 1600 MG, PLACENTAL		Cleft Palate Complications Of Maternal Exposure To Therapeutic Drugs	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	
1600 MG PLACENTAL; 1500 MG, PLACENTAL				Tegretol (Carbamazepine)	SS		

Synthroid
(Levothyroxine
Sodium) C

Date:05/08/01ISR Number: 3719663-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100059
Age:93 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 Other MG, TID) PER ORAL	Abnormal Behaviour Apnoea Chronic Obstructive Pulmonary Disease	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
	Confusional State Drug Effect Decreased Dyspnoea Feeling Hot And Cold Gastrointestinal Disorder Myocardial Infarction Refusal Of Treatment By Patient		Fosamax (Alendronate Sodium) Lotrel (Amlodipine Besylate, Benazepril Hydrochloride) Norvasc (Amlodipine Besilate) Syntrhoid (Levothyroxine Sodium) Oxygen (Oxygen) Beta Blocker Eye Drops Unspecified Eye Drops	SS C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/08/01ISR Number: 3719822-5Report Type:Expedited (15-DaCompany Report #046-0945-M0100018

Age:47 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Cardiomegaly	Foreign	Neurontin	PS	Parke Davis	
Initial or Prolonged	Chest Pain	Other			Pharmaceuticals Ltd	ORAL
2400 MG, PER	Ear Pain					
ORAL	Fatigue		Furix (Furosemide)	C		
	Headache		Arava (Leflunomide)	C		
	Hypervolaemia		Vioxx (Rofecoxib)	C		
	Liver Function Test		Folacin (Folic Acid,			
	Abnormal		Calcium Phosphate)	C		
	Pharyngolaryngeal Pain		E-Vimin (Tocopheryl			
	Pleural Effusion		Acetate)	C		
	Proteus Infection		Alvedon			
	Renal Impairment		(Paracetamol)	C		
	Sedation		Prednisolon			
	Skin Ulcer		(Prednisolone)	C		
	Urinary Tract Infection		Morfin (Morphine)	C		
	Wound Infection		Prepulsid			
			(Cisapride)	C		
			Zyrlex (Cetirizine)	C		
			Dolcontin (Morphine			
			Sulfate)	C		
			Lanzo (Lansoprazole)	C		
			Stilnoct (Zolpidem			
			Tartrate)	C		
			Loperamid			
			(Loperamide)	C		
			Calcichew-D3			
			(Colecalciferol,			
			Calcium Carbonate)	C		
			Combivent			
			(Ipratropium			
			Bromide, Salbutamol			
			Sulfate)	C		
			Spironolactone	C		

Date:05/08/01ISR Number: 3720629-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100261

Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Aspartate Aminotransferase	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
3200 MG		Increased					
(DAILY), PER		Blood Alkaline					
ORAL		Phosphatase Increased		Vicodin (Paracetamol, Hydrocodone Bitartrate) (Unspecified Pain Pills) (Fluorouracil)	C C C		

Date:05/09/01ISR Number: 3720379-3Report Type:Periodic Company Report #PHEH2000US03955
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Drug Interaction	Health Professional	Tegretol	PS	Novartis Pharmaceuticals Corp	ORAL
QD, ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

<p>400 MG, TID, ORAL QD, ORAL</p>	<p>Neurontin (Gabapentin) Capsule, 400 Mg.</p>	<p>SS</p>	<p>ORAL</p>
	<p>Celebrex (Celecoxib) Capsule</p>	<p>SS</p>	<p>ORAL</p>

Date:05/10/01ISR Number: 3720729-8Report Type:Expedited (15-DaCompany Report #001-0945-M0001145
Age:16 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 12000 MG Other (3000 MG, QID), PER ORAL	Clonic Convulsion Drug Level Above Therapeutic Respiratory Distress Status Epilepticus	Health Professional	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
1200 MG (400 MG, TID)			Neurontin (Gabapentin)	SS		
1 MG (0.5 MG, BID)			Clonazepam	SS		
.5 MG (0.25 MG, BID)			Clonazepam	SS		
3500 MG (EVERY 6 HOURS), PER ORAL			Valproic Acid	SS		ORAL

Date:05/10/01ISR Number: 3721425-3Report Type:Expedited (15-DaCompany Report #047-0945-M0000004
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2400 MG	Blindness Optic Neuritis	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Disability (DAILY), PER Other ORAL			Professional	Rivotril (Clonazepam0 Several Unspecified Medications	C C		

Date:05/10/01ISR Number: 3721458-7Report Type:Expedited (15-DaCompany Report #033-0945-M0100061
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	SEE IMAGE	Alopecia Haematoma	Foreign Health	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
Other		Haemorrhage Intracranial	Professional	Zocor (Simvastatin)	C		

Date:05/10/01ISR Number: 3721512-XReport Type:Expedited (15-DaCompany Report #044-0945-M0000083
Age:6 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Brain Neoplasm Malignant Rhinorrhoea Sarcoma	Foreign Health Professional	Neurontin Carbamazepine	PS C	Pfizer Inc	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/11/01ISR Number: 3721888-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100542

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other 600 MG (DAILY), PER ORAL	Abdominal Distension Anaemia Dyspnoea Eructation Flatulence Gastritis Pleural Effusion Weight Increased	Consumer	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
			Lopressor (Metoprolol Tartrate)	C		
			Norvasc (Amlodipine Besilate)	C		
			Glucotrol (Glipizide)	C		
			Asa (Acetylsalicylic Acid)	C		
			Iron	C		
			Lipitor (Atorvastatin)	C		
			Prevacid (Lansoprazole)	C		

Date:05/11/01ISR Number: 3722533-3Report Type:Expedited (15-DaCompany Report #002-0945-M0100037

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other PER ORAL (SEE IMAGE)	Balance Disorder Blood Pressure Decreased Condition Aggravated Dysarthria Hypotonia Lower Respiratory Tract Infection Prostration Renal Impairment	Foreign Health Professional	Neurontin (Clozapine) (Haloperidol) (Lithium) (Procyclidine)	PS C C C C	Pfizer Inc	ORAL

Sedation

Date:05/14/01ISR Number: 3723775-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100247
 Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Dry Mouth	Health	Neurontin	PS	Parke Davis	
UNKNOWN	900 MG	Feeling Abnormal	Professional			Pharmaceuticals Ltd	
MG, TID),		Movement Disorder					
UNKNOWN		Muscle Twitching					
		Sensory Loss		Multivitamins	C		
		Vision Blurred		Naproxen	C		
				B-Complex	C		

Date:05/14/01ISR Number: 3723808-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100554
 Age:36 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Blood Pressure Increased
Initial or Prolonged	Body Temperature
Other	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
2700 MG	(DAILY), PER	Chest Pain Dizziness Dysphagia	Consumer	Neurontin	PS	Parke Davis Pharmaceuticals Ltd	ORAL
ORAL		Dysphemia Headache					
1800 MG DAILY		Pneumonia Skin Test Positive Tuberculosis Vision Blurred		Neurontin (Gabapentin)	SS		
				Humalog (Insulin Lispro) Pancrease (Pancrelipase) Lorcet (Paracetamol, Hydrocodone Bitartrate) Valium (Diazepam) Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acide, Thiamine Hydrochloride, Vitamin E (Tocopherol)	C C C C C		

Date:05/14/01ISR Number: 3723817-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100553
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - PER ORAL	Initial or Prolonged	Weight Increased	Consumer	Neurontin	PS	Pfizer Inc	ORAL
				Depakote (Valporate Semisodium) Eskalith (Lithium)	C C		

Date:05/14/01ISR Number: 3724116-8Report Type:Expedited (15-DaCompany Report #049-0945-M0100046
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - PER ORAL		Blood Creatinine	Foreign	Neurontin	PS		ORAL
Initial or Prolonged		Increased Blood Urea Increased Glomerular Filtration Rate Decreased Renal Transplant	Health Professional				

Date:05/15/01ISR Number: 3723559-6Report Type:Expedited (15-DaCompany Report #061-0945-M0100019
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - INTRAVENOUS	"SEE IMAGE"	Aspergilloma Dementia	Foreign Consumer	Neurontin Voriconazole	PS SS	Pfizer Inc	
Initial or Prolonged		Hypoxia Nervous System Disorder Pneumonitis	Health Professional	Ambisome (Amphotericine B, Liposome) Effexor (Venlafaxine Hydrochloride) Meropenem (Meropenem)	SS SS C		

Freedom Of Information (FOI) Report

Ciprofloxacin
 (Ciprofloxacin) C
 Zantac (Ranitidine
 Hydrochloride) C
 Fluoxetine
 (Fluoxetine) C
 Fentanyl (Fentanyl) C
 Amitriptyline
 (Amitriptyline) C
 Paracetamol
 (Paracetamol) C
 Amikacin (Amikacin) C
 Clindamycin
 (Clindamycin) C
 Ceftazidime
 (Ceftazidime) C
 Fusidic Acid
 (Fusidic Acid) C

Date:05/15/01ISR Number: 3723561-4Report Type:Expedited (15-DaCompany Report #002-0945-M0100044
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1100 MG Initial or Prolonged (TWICE DAILY) , PER ORAL	Dizziness Mood Altered	Foreign Health Professional	Neurontin	PS	Pfizer Inc	ORAL

Date:05/15/01ISR Number: 3723567-5Report Type:Expedited (15-DaCompany Report #064-0945-M0100001
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death PER ORAL Life-Threatening	Myocardial Infarction Neutropenia Pulmonary Embolism	Foreign Health Professional	Neurontin	PS	Pfizer Inc	ORAL

Date:05/16/01ISR Number: 3724143-0Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Suicide Attempt		Ativan Clonopin Prozac Trazodone Synthroid Gabapentin "Silver Bullet" Otc	PS SS SS SS SS SS SS		

Date:05/17/01ISR Number: 3724359-3Report Type:Direct
Age: Gender: I/FU:I

Company Report #USP 54047

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Neurontin	PS	Pfizer	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/17/01ISR Number: 3724466-5Report Type:Direct
Age:75 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Adenocarcinoma		Neurontin 400 Mgm. Q			
	Pancreatic Carcinoma		8 Hr.	PS		

Date:05/18/01ISR Number: 3724768-2Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Confusional State		Amitriptylline	PS		ORAL
1650 MG PO X						
Hospitalization -	Hallucination					
1						
Initial or Prolonged	Overdose		Bumetanide	C		
Required	Pupillary Reflex Impaired		Loratidine	C		
Intervention to			Pseudoephedrine	C		
Prevent Permanent			Clorazepam	C		
Impairment/Damage			Temazepan	C		
			Daypro	C		
			Doxycycline	C		
			Gabapentin	I		

Date:05/18/01ISR Number: 3725878-6Report Type:Expedited (15-DaCompany Report #049-0945-M0100033
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Other	Dialysis	Foreign	Neurontin	PS	Pfizer Inc	ORAL
2700 MG						
(DAILY), PER	Muscle Spasms	Health				
	Muscle Twitching	Professional				
ORAL						
	Muscular Weakness		Duragesic (Fentanyl)	SS		
TRANSDERMAL						
75	Renal Failure Acute					
(TRANSDERMAL)						
	Tremor		Sandimmune (Ciclosporin)	SS		

Date:05/18/01ISR Number: 3725916-0Report Type:Periodic
Age:58 YR Gender:Female I/FU:I

Company Report #C2001-0126.01

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Clozapine	PS	Mylan Pharmaceuticals Inc	ORAL
100MG Q AM,							
100MG Q 4 PM,							
250MG Q HS							
ORAL							
				Neurontin Parke-Davis	SS	Parke-Davis	ORAL
300MG Q AM,							
800MG Q HS,							
ORAL							
				Dilantin Parke-Davis	SS	Parke-Davis	ORAL
200MG Q AM,							
200MG Q PM,							
ORAL							
				Celecoxib	C		
				Levothyroxine	C		
				Lorazepam	C		
				Zolpidem	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/18/01ISR Number: 3726255-4Report Type:Expedited (15-DaCompany Report #044-0945-M0100141

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 3000 MG Initial or Prolonged (DAILY), PER ORAL	Pyelonephritis Chronic	Foreign Consumer Other	Neurontin (Ketamine) (Tramadol) (Morphine) (Trimethoprim) (Amitriptyline) Losec (Omeprazole)	PS C C C C C C	Pfizer Inc	ORAL

Date:05/21/01ISR Number: 3726922-2Report Type:Expedited (15-DaCompany Report #055-0945-M0100017

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 400 MG Initial or Prolonged (DAILY), PER ORAL	Thrombosis	Foreign Consumer	Neurontin Unspecified Medications	PS C	Pfizer Inc	ORAL

Date:05/21/01ISR Number: 3726923-4Report Type:Expedited (15-DaCompany Report #032-0945-M0100002

Age:49 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 300 MG (DAILY), PER ORAL	Petechiae	Foreign Health Professional	Neurontin	PS	Pfizer Inc	ORAL

Date:05/22/01ISR Number: 3727194-5Report Type:Expedited (15-DaCompany Report #001-0981-M0103661
Age:47 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 10 MG (BID), Initial or Prolonged PER ORAL		Unevaluable Event	Health Professional	Lipitor (Gabapentin)	PS SS	Pfizer Inc	ORAL ORAL
1200 MG (EVERY AM)'600 MG (EVERY NOON);							
1200 MG 40 MG (DAILY);40 MG (BID);40 MG (DAILY); 20 MG (DAILY),				(Geodon)	SS		ORAL
325 MG (DAILY), PER ORAL				(Acetylsalicylic Acid)	SS		ORAL
100 MG (EVERY 4 HOURS PRN), PER ORAL				(Chlorpromazine)	SS		ORAL
2 MG (AT BEDTIME), PER ORAL				(Clonazepam)	SS		ORAL
2 MG (EVERY 6				(Lorazepam)	SS		ORAL

Freedom Of Information (FOI) Report

HOURS); 2 MG
 (EVERY 4 HR
 PRN), PER
 ORAL
 50 MG (Mesoridazine) SS ORAL
 (DAILY);75 MG
 (BID);75 MG
 (TID); 50 MG
 (EVERY 4
 15 MG (AT (Mirtazapine) SS ORAL
 BEDTIME), PER
 ORAL

Date:05/22/01ISR Number: 3727203-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100590
 Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 900 MG (300 Initial or Prolonged MG, TID), UNKNOWN		Anxiety	Literature	Neurontin	PS	Pfizer Inc	
		Bronchospasm	Health				
		Chronic Obstructive Airways Disease Exacerbated Hypercapnia Hypoventilation Insomnia Respiratory Distress Respiratory Failure Respiratory Rate Increased	Professional	Albuterol (Salbutamol) Ipratropium Bromide (Ipratropium Bromide) Clonazepam (Clonazepam) Zolpidem (Zolpidem)	C C C C		

Date:05/22/01ISR Number: 3727204-5Report Type:Expedited (15-DaCompany Report #001-0981-M0103676
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Lipitor	PS	Pfizer Inc	ORAL
20 MG							
Required		Arthralgia					
(DAILY) ,							
Intervention to		Carpal Tunnel Syndrome					
PER ORAL							
Prevent Permanent		Condition Aggravated		Neurotin	SS		ORAL
900 TO							
Impairment/Damage		Disorientation					
1200MG							
(UNKNOWN) ,		Feeling Abnormal					
PER ORAL		Mental Impairment					
UNKNOWN		Neck Pain		Celebrex	SS		ORAL
(UNKNOWN) ,		Oedema Peripheral					
PER ORAL		Pain		(Fentanyl)	C		

Date:05/23/01ISR Number: 3727887-XReport Type:Expedited (15-DaCompany Report #A108646
 Age:25 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Mania	Consumer	Geodon	PS	Pfizer Central	ORAL
Initial or Prolonged		Medication Error	Health			Research	
ORAL							
300.00 MG		Schizophrenia	Professional	Seroquel	SS		ORAL
TOTAL DAILY							
ORAL				Neurontin			
3600.00 MG				(Gabapentin)	SS		
TOTAL TID							
ORAL				Beer	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Depakote C
Clonazepam C

Date:05/23/01ISR Number: 3728004-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100587
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Neurontin	PS	Pfizer Inc	
Other		Coma	Professional	Phenytoin Suspension (Phenytoin Sodium)	SS		
UNKNOWN							
UNKNOWN							

Date:05/24/01ISR Number: 3728409-XReport Type:Direct Company Report #
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin / 300mg			
Required		Sedation		Capsule / Park			
Intervention to		Speech Disorder		Davis, Lot No			
Prevent Permanent				16131va,	PS		ORAL
Impairment/Damage							
600MG BID							
ORAL							

Date:05/24/01ISR Number: 3728888-8Report Type:Direct Company Report #
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin 300mg			
Required		Depressed Level Of		Capsule Park Davis	PS	Park Davis	ORAL
Intervention to		Consciousness					
Prevent Permanent		Speech Disorder					
ORAL							
Impairment/Damage							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional	Ativan	PS	Wyeth Ayerst Laboratories	ORAL
Other		Convulsion					
0.5 MG AS							
NEEDED, ORAL							
DOSE UNKNOWN							
DOSE UNKNOWN							
				Dilantin (Phenytoin Sodium)	SS		
				Neurontin (Gabapentin)	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign	Neurontin	PS	Pfizer Inc	ORAL
Life-Threatening		Dementia					
PER ORAL							
Hospitalization -		Liver Function Test	Consumer	Voriconazole	SS		
INTRAVENOUS	700 MG (BID),						
Initial or Prolonged		Abnormal	Health				
INTRAVENOUS							
				Ambisome (Amphotericine B, Liposome)	SS		
INTRAVENOUS	INTRAVENOUS	Pneumonitis	Professional				
				Effexor (Venlafaxine Hydrochloride)	SS		ORAL
PER ORAL							
				Meropenem (Meropenem)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ciprofloxacin
 (Ciprofloxacin) C
 Zantac (Ranitidine
 Hydrochloride) C
 Fluoxetine
 (Fluoxetine) C
 Fentanyl (Fentanyl) C
 Amitriptyline
 (Amitriptyline) C
 Paracetamol
 (Paracetamol) C
 Amikacin (Amikacin) C
 Clindamycin
 (Clindamycin) C
 Ceftazidime
 (Ceftazidime) C
 Fusidic Acid
 (Fusidic Acid) C

Date:05/25/01ISR Number: 3730605-2Report Type:Expedited (15-DaCompany Report #033-0945-M0100068
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Bone Density Decreased Osteonecrosis	Foreign Consumer Other	Neurontin Didanosine (Didanosine)	PS SS	Pfizer Inc	ORAL
PER ORAL				Efavirenz (Efavirenz)	SS		ORAL
PER ORAL				Abacavir (Abacavir) Bactrim (Sulfamethoxazole, Trimethoprim)	SS SS		ORAL
PER ORAL							

Date:05/25/01ISR Number: 3730608-8Report Type:Expedited (15-DaCompany Report #055-0945-M0100011
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 900 MG Initial or Prolonged (UNKNOWN), Other PER ORAL	Oedema Peripheral Refusal Of Treatment By Patient	Foreign Consumer	Neurontin	PS	Pfizer Inc	ORAL
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Date:05/29/01ISR Number: 3729001-3Report Type:Expedited (15-DaCompany Report #260859
 Age:64 YR Gender:Male I/FU:I

Outcome Dose Hospitalization - Initial or Prolonged	PT Dysphagia	Report Source	Product	Role	Manufacturer	Route
			Klonopin Tablets	PS	Roche	
			Neurontin	SS		
			Risperdal	SS		
			Cogentin	SS		
			Lopressor	C		
			Heparin	C		
			Inderal	C		
			Pepcid	C		
			Asa	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/29/01ISR Number: 3729952-XReport Type:Expedited (15-DaCompany Report #001-0945-M0100502

Age:47 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 600 MG (200 Initial or Prolonged MG, BID) PER ORAL	Asthenia Blood Potassium Decreased Blood Sodium Decreased Convulsion Dizziness Muscle Spasms Peripheral Nerve Injury Weight Increased	Health Professional	Neurontin Depakote (Valproate Semisodium) Mysoline (Primidone) Klonopin (Clonazepam)	PS C C C	Pfizer Inc	ORAL

Date:05/29/01ISR Number: 3730228-5Report Type:Expedited (15-DaCompany Report #044-0945-M0100144

Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300 MG Initial or Prolonged (UNKNOWN) , PER ORAL	Gastrointestinal Obstruction	Foreign Consumer Other	Neurontin Loperamide (Loperamide) Prednisolone (Prednisolone) Codeine Phosphate (Codeine Phosphate) Ranitidine (Ranitidine)	PS C C C C	Pfizer Inc	ORAL

Date:05/29/01ISR Number: 3730390-4Report Type:Expedited (15-DaCompany Report #055-0945-M0100013

Age:62 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Hospitalization - 800 MG Initial or Prolonged (DAILY), PER	Blood Pressure Increased Cough Dyspnoea	Foreign Consumer	Neurontin	PS	Pfizer Inc	ORAL
ORAL			(Insulin)	C		

Date:05/29/01 ISR Number: 3730810-5 Report Type:Expedited (15-DaCompany Report #10838803
Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Agranulocytosis Leukopenia	Foreign Health Professional Other	Zerit	PS	Bristol Myers Squibb Co Pharmaceutical Research Institute	
INTRAVENOUS	50 MILLIGRAM,			Fungizone (Amphotericin B)	SS		
IV				Ancotil (Flucytosine)	SS		
INTRAVENOUS	IV			Fluconazole	SS		
200							
MILLIGRAM,							
2/1 DAY				Gabapentin	SS		
300 MILLIGRAM				Lamivudine + Zidovudine	SS		
				Metoclopramide (Metoclopramide Hcl)	SS		
				Paracetamol	SS		

Freedom Of Information (FOI) Report

Trimethoprim + Sulfamethaxazole (Trimethoprim + Sulfam	SS
Didanosine (Didanosine)	C
Thiamin (Thiamine)	C
Kalii Chloridum (Potassium Chloride)	C
Magnesium (Magnesium)	C
Clonazepam (Clonazepam)	C
Paracetamol + Codeine (Paracetamol + Codeine)	C
Filgrastim (Granulocyte Csf)	C
Nelfinavir (Nelfinavir Mesylate)	C
Guaifenesin (Guaifenesin)	C
3tc (Lamivudine)	C

Date:05/29/01ISR Number: 3730814-2Report Type:Expedited (15-DaCompany Report #10842557
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthralgia Bone Density Decreased Hip Deformity	Foreign Health Professional	Videx	PS	Bristol Myers Squibb Co Pharmaceutical Research Institute	ORAL
ORAL		Hypertriglyceridaemia Lipodystrophy Acquired Osteonecrosis	Other	Zerit (Stavudine) Neurontin (Gabapentin) Ziagen (Abacavir) Bactrim Forte (Trimethoprim+Sulfam ethoxazole) Sustiva (Efavirenz) Lipur (Gemfibrozil) Lipur	SS SS SS SS C C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Disability	Asthenia		Neurontin	PS	Pfizer Inc	ORAL
2100 PER ORAL						
Other	Coordination Abnormal		Wellbutrin			
	Hyporeflexia		(Amfebutamone			
			Hydrochloride)	C		
			Doxepin (Doxepin)	C		
			Tegretol			
			(Carbamazepine)	C		
			Synthroid			
			(Levothyroxine			
			Sodium)	C		
			Premarin (Estrogens			
			Conjugated)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/30/01ISR Number: 3730845-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100593

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2000 MG		Asthenia	Consumer	Neurontin	PS	Pfizer Inc	ORAL
Initial or Prolonged (DAILY), PER Disability ORAL		Difficulty In Walking					
		Fall		Serzone (Nefazodone Hydrochloride)	SS		
				Klonopin (Clonazepam)	C		
				Prilosec (Omeprazole)	C		
				Analgesics	C		

Date:05/30/01ISR Number: 3730855-5Report Type:Expedited (15-DaCompany Report #033-0945-M0100075

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cholecystitis Dyspnoea Fatigue Hepatic Trauma Hypersensitivity Oedema	Foreign Health Professional	Neurontin	PS	Pfizer Inc	

Date:05/30/01ISR Number: 3730856-7Report Type:Expedited (15-DaCompany Report #033-0945-M0100076

Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 200 MG		Blood Alkaline	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Initial or Prolonged (DAILY) PER		Phosphatase Increased	Literature				
Other ORAL		Confusional State	Consumer				
		Disorientation					

Mental Impairment
 Pyrexia
 Rash Maculo-Papular
 Rash Pruritic

Date:05/30/01ISR Number: 3730859-2Report Type:Expedited (15-DaCompany Report #049-0945-M0100051

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haemosiderosis	Foreign	Neurontin	PS	Pfizer Inc	
UNKNOWN		UNK {UNK),					
		Hepatic Cirrhosis	Health				
UNK		Hepatitis	Professional				

Date:05/30/01ISR Number: 3730925-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100592

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ecchymosis	Consumer	Neurontin	PS	Pfizer Inc	ORAL
300 MG		Fall					
(DAILY), PER		Laceration					
ORAL		Pyrexia		Deltasone			
		Sedation		(Prednisone)	C		
		Vision Blurred		Glucotrol			
				(Glipizide)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Avapro (Irbesartan) C
 Hydrochlorothiazide
 (Hydrochlorothiazide
) C

Date:05/31/01ISR Number: 3731374-2Report Type:Expedited (15-DaCompany Report #260859
 Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	6 MG DAILY	Dysphagia Hyporeflexia	Health Professional	Klonopin	PS	Hoffmann La Roche Inc	ORAL
ORAL				Neurontin (Gabapentin)	SS		
	800 MG DAILY			Risperdal (Risperidone)	SS		
	4 MG DAILY			Cogentin (Benztropine Mesylate)	SS		
	2 MG DAILY			Lopressor (Metoprolol Tartrate)	C		
				Heparin (Heparin Sodium)	C		
				Inderal (Propranolol Hydrochloride)	C		
				Pepcid (Famotidine)	C		
				Asa (Aspirin)	C		

Date:06/01/01ISR Number: 3731216-5Report Type:Expedited (15-DaCompany Report #WAES 01027276
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 7 DAY		Alanine Aminotransferase		Vasotec	PS	Merck & Co., Inc	ORAL
Initial or Prolonged		Increased		Vioxx	SS		ORAL

	Aspartate	Gabapentin	SS	ORAL
	Aminotransferase	Metronidazole	SS	ORAL
9	DAY			
	Increased	Furosemide	SS	ORAL
7	DAY			
	Blood Alkaline	Floxacillin Sodium	SS	ORAL
9	DAY			
	Phosphatase Increased	Leflunomide	SS	ORAL
	Blood Bilirubin Increased	Spironolactone	SS	ORAL
	Blood Lactate	Folic Acid	C	
	Dehydrogenase Increased	Cisapride	C	
	Cardiomegaly	[Therapy		
	Culture Urine Positive	Unspecified]	C	
	Depressed Level Of	Albuterol Sulfate		
	Consciousness	And Ipratropium		
	Electroencephalogram	Bromide	C	
	Abnormal	Calcium Carbonate		
	Fatigue	And Cholecalciferol	C	
	Gamma-Glutamyltransferase	Loperamide	C	
	Increased	Zolpidem Tartrate	C	
	Headache	Acetaminophen	C	
	Infected Skin Ulcer	Vitamin E	C	
	Pharyngolaryngeal Pain	Cetirizine		
	Pleural Disorder	Hydrochloride	C	
	Proteus Infection	Lansoprazole	C	
	Renal Failure	Morphine	C	
		Prednisolone	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Morphine C

Date:06/01/01ISR Number: 3731734-XReport Type:Expedited (15-DaCompany Report #B0108767A

Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	1UNIT Twice	Aseptic Necrosis Bone		Ziagen	PS	Glaxo Wellcome	ORAL
	per day	Bone Density Decreased					
	3UNIT per day	Hypertriglyceridaemia		Sustiva	SS		ORAL
	1UNIT Three	Lipodystrophy Acquired		Neurontin	SS		ORAL
	times per day						
	1UNIT per day			Bactrim Forte	SS	Glaxo Wellcome	ORAL
	UNKNOWN			Videx	SS		
	60 DAY			Combivir	C	Glaxo Wellcome	ORAL
				Epivir	C	Glaxo Wellcome	ORAL
				Retrovir	C	Glaxo Wellcome	ORAL
				Crixivan	C		
		738 DAY					
		378 DAY		Invirase	C		
		378 DAY		Viracept	C		
		485 DAY		Zerit	C		
				Depakine Chrono	C		

Date:06/01/01ISR Number: 3732111-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000487

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	BID, PER ORAL	Ecchymosis	Health	Neurontin	PS	Pfizer Inc	ORAL

Gingival Bleeding
Retinal Haemorrhage

Professional

Meclomen (Meclofen
Amate Sodium) C
Premarin (Estrogens
Conjugated) C
Claritin
(Loratadine) C
Aleve (Naproxen
Sodium) C

Date:06/01/01ISR Number: 3732129-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100635
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (DAILY) PER ORAL		Hepatic Steatosis Hepatitis Liver Function Test Abnormal	Health Professional	Neurontin	PS	Pfizer Inc	ORAL
UNKNOWN	UNK, UNK			Methergine (Methylergometrine Maleate)	SS		
UNKNOWN	UNK, UNK			Skelaxin (Metaxalone)	SS		
UNKNOWN	UNK, UNK			Prozac (Fluoxetine Hydrochloride)	SS		

Date:06/01/01ISR Number: 3732388-9Report Type:Expedited (15-DaCompany Report #044-0945-M0100153
Age: Gender:Male I/FU:I

Outcome PT
Hospitalization - Drug Ineffective
Initial or Prolonged Jaundice

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Renal Failure Acute

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
800 MG (BID)		Foreign Health	Neurontin	PS	Pfizer Inc	ORAL
PER ORAL		Professional Company Representative	Carbamazepine (Carbamazepine)	C		

Date:06/01/01ISR Number: 3732618-3Report Type:Expedited (15-DaCompany Report #044-0945-M0100155
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign Health	Neurontin	PS	Pfizer Inc	
Other		Alopecia	Professional	Cardiac Medications	C		

Date:06/04/01ISR Number: 3733093-5Report Type:Expedited (15-DaCompany Report #WAES 01027276
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Potassium Increased Cardiac Failure Congestive	Foreign Company Representative	Vasotec	PS	Merck Research Laboratories Div Merck Co Inc	ORAL
20 MG PO	7 DAY	Cardiomegaly Depressed Level Of		Tab Vioxx (Rofecoxib)	SS		ORAL
25 MG PO		Consciousness		Tab Metronidazole	SS		ORAL
800 MG PO	9 DAY	Electroencephalogram		Tab Furosemide	SS		ORAL
40 MG PO	7 DAY	Abnormal		Cap Gabapentin	SS		ORAL
2400 MG PO		Fatigue Headache		Tab Floxacillin Sodium	SS		ORAL
2200 MG PO	9 DAY						

50 MG PO	Hepatic Failure	Tab Spironolactone	SS	ORAL
20 MG PO	Infected Skin Ulcer	Tab Leflunomide	SS	ORAL
	Metabolic Disorder	(Therapy		
	Pharyngolaryngeal Pain	Unspecified)	C	
	Proteus Infection	Zolpidem Tartrate	C	
	Renal Failure	Vitamin E	C	
	Urinary Tract Infection	Prednisolone	C	
		Morphine	C	
		Loperamide	C	
		Acetaminophen	C	
		Cetirizine		
		Hydrochloride	C	
		Cisapride	C	
		Lansoprazole	C	
		Folic Acid	C	
		Albuterol Sulfate		
		(+) Ipratropium		
		Bromide	C	
		Calcium Carbonate		
		(+) Cholecalciferol	C	

Date:06/04/01ISR Number: 3733305-8Report Type:Expedited (15-DaCompany Report #055-0945-M0100019
Age:72 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 400 MG, Initial or Prolonged (DAILY), PER	Condition Aggravated Varicose Vein	Foreign Consumer	Neurontin	PS	Pfizer Inc	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Unspecified
Antihypertensive
Agent C

Date:06/04/01ISR Number: 3733306-XReport Type:Expedited (15-DaCompany Report #055-0945-M0100016
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Toxicity	Foreign	Neurontin	PS	Pfizer Inc	ORAL
800 MG (BID),		Photophobia	Health				
PER ORAL			Professional				

Date:06/04/01ISR Number: 3733495-7Report Type:Periodic Company Report #A107643
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Health	Geodon	PS	Pfizer Central Research	
40.00 MG		Orthostatic Hypotension	Professional				
TOTAL:DAILY				Neurontin	SS		
				Clonidine	SS		

Date:06/04/01ISR Number: 3733772-XReport Type:Periodic Company Report #A100681
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Consumer	Lithium Carbonate	PS	Pfizer Inc	
		Thinking Abnormal		Gabapentin	SS		
				Divalproex	SS		

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Photophobia	Foreign	Neurontin	PS	Pfizer Inc	ORAL
800 MG (BID),			Health				
PER ORAL			Professional				

Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Hypertriglyceridaemia	Foreign	Ziagen	PS	Glaxo Wellcome Inc	ORAL
1 UNIT /							
TWICE PER DAY							
/ ORAL							
				Efavirenz (Efavirenz)	SS		ORAL
				Gabapentin (Gabapentin)	SS		ORAL
1 UNIT /							
THREE TIMES							
PER DAY /							
ORAL							
				Septra (Sulfamethoxazole/Tr imetho)	SS		ORAL
ORAL				Didanosine			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Didanosine)	SS
Combivir	C
Lamivudine	C
Zidovudine	C
Indinavir Sulfate	C
Saquinavir	C
Nelfinavir Mesylate	C
Stavudine	C
Depakine Chrono	C

Date:06/05/01ISR Number: 3735322-0Report Type:Expedited (15-DaCompany Report #001-0945-M0100327
 Age:68 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetes Mellitus	Consumer	Neurontin	PS	Pfizer Inc	ORAL
600 MG (BID),		Non-Insulin-Dependent					
PER ORAL							

Tenormin (Atenolol)	C
Accupril (Quinapril Hydrochloride)	C
Hydrocortisone (Hydrocortisone)	C
Megace (Megestrol Acetate)	C
Darvocet (Paracetamol, Dextropropoxyphene)	C
Dalmane (Flurazepam Hydrochloride)	C

Date:06/05/01ISR Number: 3735366-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100410
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Depression	Consumer	Neurontin	PS	Pfizer Inc	ORAL
1200 MG							
Initial or Prolonged		Malabsorption	Health				
(DAILY), PER							
Other		Uterine Leiomyoma	Professional				
ORAL							

250 MG

Zoloft (Sertraline) SS

(DAILY)

Vicodin
(Paracetamol,
Hydrocodone
Bitartrate) C

Date:06/05/01ISR Number: 3735548-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100645

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Gastroenteritis	Consumer	Neurontin	PS	Pfizer Inc	ORAL
2400 MG DAILY		Helicobacter					
PER ORAL ;		Loss Of Libido					
1200 MG DAILY		Malaise					
PER ORAL							

Clonazepam
(Clonazepam) C
Serzone (Nefazodone
Hydrochloride) C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/05/01ISR Number: 3735579-6Report Type:Expedited (15-DaCompany Report #PHEH2001US04478

Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Hepatic Steatosis	Health Professional	Methergine	PS	Novartis Pharmaceuticals Corp	
2	MON			Pamelor (Nortriptyline Hydrochloride)	SS		
2	MON			Skelaxin (Metaxalone)	SS		
2	MON			Prozac (Fluoxetine Hydrochloride)	SS		
2	MON			Neurontin (Gabapentin)	SS		

Date:06/06/01ISR Number: 3735426-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100683

Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 200 MG Initial or Prolonged (DAILY), PER Other ORAL		Blood Alkaline Phosphatase Increased Confusional State Culture Urine Positive Dermatitis Exfoliative Gamma-Glutamyltransferase Increased Hypersensitivity Mental Impairment Pyrexia Pyuria Rash Maculo-Papular Rash Pruritic Splenomegaly White Blood Cells Urine	Foreign Literature	Neurontin	PS	Pfizer Inc	ORAL

Positive

Date:06/06/01ISR Number: 3735800-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100482
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 7200 MG Initial or Prolonged (TID), PER		Unevaluable Event	Health Professional	Neurontin	PS	Pfizer Inc	ORAL
ORAL							

Date:06/06/01ISR Number: 3735803-XReport Type:Expedited (15-DaCompany Report #001-0945-M0100657
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other SEE IMAGE		Bone Density Decreased	Consumer	Neurontin	PS	Pfizer Inc	ORAL
				Prozac (Fluoxetine Hydrochloride)	C		
				Unspecified Thyroid Medication	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/07/01ISR Number: 3735750-3Report Type:Expedited (15-DaCompany Report #200112306EU

Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	800 MG/DAY PO 9 DAY	Blood Alkaline Phosphatase Increased	Foreign Other	Noritate	PS	Dermik Laboratories Inc	ORAL
	20 MG/DAY PO	Blood Bilirubin Increased		Leflunomide (Arava)	SS		ORAL
	2.4 G/DAY	Cardiomegaly Chest Pain		Gabapentin (Neurontin)	SS		
	40 MG/DAY PO	Depressed Level Of		Furosemide (Furix)	SS		ORAL
	20 MG/DAY PO 1 WK	Consciousness Ear Pain		Enalapril Maleate (Renitec)	SS		ORAL
	2.2 G/DAY 9 DAY	Fatigue Gamma-Glutamyltransferase Increased		Flucloxacillin Sodium (Heracillin)	SS		
	25 G/DAY PO			Rofecoxib (Vioxx)	SS		ORAL
		Headache		Folic Acid	C		
		Hypervolaemia		Calcium Phosphate			
		Hypocalcaemia		(Folacin)	C		
		Infected Skin Ulcer		Vitamin E	C		
		Liver Disorder		Paracetamol			
		Metabolic Disorder		(Alvedon)	C		
		Pharyngolaryngeal Pain		Prednisone	C		
		Pleural Effusion		Morfin	C		
		Proteus Infection		Cisapride			
		Renal Failure		(Prepulsid)	C		
		Weight Increased		Cetirizine (Zyrlex)	C		
				Morfin	C		
				Lansoprazole (Lanzo)	C		
				Zolpidem Tartrate			
				(Stilnoct)	C		
				Loperamide	C		
				Colecalciferol	C		
				Calcium Carbonate			
				(Calcichew D3)	C		
				Ipratropium Bromide	C		
				Salbutamol Sulfate			
				(Combivent)	C		
				Spirolactone	C		

Date:06/08/01ISR Number: 3736242-8Report Type:Expedited (15-DaCompany Report #A107350
Age:47 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 250.00 MG Required TOTAL:DAILY Intervention to 1200.00 MG Prevent Permanent TOTAL:QID:ORA Impairment/Damage L	Depression Drug Effect Decreased Malabsorption Uterine Leiomyoma	Consumer Health Professional	Zoloft Gabapentin Vicodin	PS SS C	Pfizer Pharmaceuticals Inc	 ORAL

Date:06/11/01ISR Number: 3737156-XReport Type:Expedited (15-DaCompany Report #001-0981-M0103661
Age:47 YR Gender:Unknown I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 10 MG (BID), Initial or Prolonged PER ORAL 1200 MG (EVERY AM); 600 MG (EVERY	No Adverse Drug Effect	Health Professional	Lipitor Gabapentin	PS SS	Pfizer Inc	ORAL ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

NOON); 1200			
MG (EVERY HS.			
40 MG	(Geodon)	SS	ORAL
(DAILY); 40			
MG (BID); 40			
MG (DAILY);			
20 MG (DAILY)			
325 MG	(Acetylsalicylic Acid)	SS	ORAL
(DAILY), PER			
ORAL			
100 MG (EVERY	(Chlorpromazine)	SS	ORAL
4 HOURS PRN),			
PER ORAL			
2 MG (AT	(Clonazepam)	SS	ORAL
BEDTIME), PER			
ORAL			
2 MG (EVERY 6	(Lorazepam)	SS	ORAL
HOURS); 2 MG			
(EVERY 8			
HOURS AS			
NEEDED) PER			
50 MG	(Mesoridazine)	SS	ORAL
(DAILY); 75			

MG (BID); 75
 MG (TID) PER
 ORAL; 50 MG
 15 MG (AT
 BEDTIME), PER
 ORAL
 (Mirtazapine) SS ORAL
 ... C
 .. C
 ... C
 ... C
 ... C
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 ... C
 ... C
 ... C
 C
 ... C

Date:06/11/01ISR Number: 3737229-1Report Type:Expedited (15-DaCompany Report #001-0981-M0103676
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Lipitor	PS	Pfizer Inc	ORAL
10 MG							
Required		Carpal Tunnel Syndrome	Health				
(DAILY), PER							
Intervention to		Disorientation	Professional				
ORAL							
Prevent Permanent		Drug Effect Decreased		Neurontin			
Impairment/Damage		Mental Impairment		(Gabapentin)	SS		ORAL
900 MG (THREE							
TIMES DAILY),							
PER ORAL							
				Celebrex	SS		ORAL
(DAILY), PER							
ORAL							
				Unspecified			
				Narcotics	SS		
				Fentanyl	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/11/01ISR Number: 3737347-8Report Type:Expedited (15-DaCompany Report #055-0945-M0100020

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1200 MG Initial or Prolonged (DAILY), PER		Haemorrhage	Foreign Consumer	Neurontin	PS	Pfizer Inc	ORAL
ORAL							

Date:06/11/01ISR Number: 3737797-XReport Type:Direct

Company Report #

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG BID Initial or Prolonged		Balance Disorder Emotional Disorder Lethargy Speech Disorder Tremor		Gabapentin	PS		

Date:06/11/01ISR Number: 3738019-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100684

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 3600 MG (UNKNOWN),		Accident Concussion	Health Professional	Neurontin	PS	Pfizer Inc	
UNKNOWN							

Date:06/11/01ISR Number: 3738020-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100671

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Blood Urea Increased	Consumer	Neurontin	PS	Pfizer Inc
"SEE IMAGE"					
	Cataract		Tylenol		
	Cataract Operation		(Paracetamol)	C	
	Eye Irritation		Tramadol (Tramadol)	C	
	Fatigue				
	Pain In Extremity				
	Peripheral Coldness				
	Urinary Retention				
	Vision Blurred				

Date:06/11/01ISR Number: 3740732-1Report Type:Periodic Company Report #254373
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Distension	Consumer	Accutane	PS	Hlr Technology	ORAL
ORAL							
		Decreased Appetite		Neurontin			
		Dry Skin		(Gabapentin)	SS		
		Lip Dry		Wellbutrin			
		Mood Swings		(Bupropion			
		Weight Decreased		Hydrochloride)	C		

Date:06/12/01ISR Number: 3738135-9Report Type:Direct Company Report #
 Age:55 YR Gender:Male I/FU:I

Outcome	PT
Other	Blister
	Inflammation

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Oedema Peripheral

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
400 MG TID			Neurontin 200 Mg Parke-Davis	PS	Parke-Davis	ORAL
ORAL			Effexor	C		
			Allegra	C		
			Zocor	C		
			Vioxx	C		
			Doxepin	C		
			Hytrin	C		
			Reglan	C		
			Prevacid	C		
			Chlorpromazine	C		
			Risperdal	C		

Date:06/12/01ISR Number: 3738718-6Report Type:Expedited (15-DaCompany Report #044-0945-M0100161
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - RECTAL 2400 MG, PER Initial or Prolonged RECTAL	Haemolytic Anaemia	Foreign Consumer	Neurontin	PS	Pfizer Inc	
		Other	Sodium Valproate (Valproate Sodium)	C		
			Folic Acid (Folic Acid)	C		

Date:06/12/01ISR Number: 3738736-8Report Type:Expedited (15-DaCompany Report #033-0945-M0100085
Age:38 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Alanine Aminotransferase Increased Asthenia Cholecystitis	Foreign Consumer Other	Neurontin	PS	Pfizer Inc	

Dyspnoea
Hypersensitivity
Oedema
Pneumonia

Date:06/12/01ISR Number: 3738899-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100514
Age:66 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Amnesia
Initial or Prolonged	Cardiac Failure
Other	Congestive
	Cerebrovascular Accident
	Coordination Abnormal
	Dizziness
	Excoriation
	Fall
	Head Injury
	Memory Impairment
	Myocardial Infarction
	Renal Failure
	Syncope

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Freedom Of Information (FOI) Report

Urinary Incontinence

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
900 MG (TID), PER ORAL		Consumer Health Professional	Neurontin Lasix (Furosemide) Thyroid Premarin (Estrogens Conjugated) Allopurinol Flexeril (Cyclobenzaprine Hydrochloride) Prilosec (Omeprazole) Potassium Duragesic Patches	PS C C C C C C C	Pfizer Inc	ORAL

Date:06/12/01ISR Number: 3738900-8Report Type:Expedited (15-DaCompany Report #001-0945-M0100708
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG, DAILY		Anorexia Concussion Diarrhoea Dysmenorrhoea Haemorrhage Nausea Syncope Thirst Visual Disturbance	Consumer	Neurontin	PS	Pfizer Inc	

Date:06/12/01ISR Number: 3738930-6Report Type:Expedited (15-DaCompany Report #055-0945-M0100024
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
Death PER ORAL			Foreign Consumer	Neurontin	PS	Pfizer Inc	ORAL
Date:06/14/01ISR Number: 3740111-7Report Type:Expedited (15-DaCompany Report #044-0945-M0100161 Age: Gender:Female I/FU:F							
Hospitalization - RECTAL	2400 MG	Haemolytic Anaemia	Foreign Consumer	Neurontin	PS	Pfizer Inc	
Initial or Prolonged (UNKNOWN) PER			Other	Sodium Valproate Folic Acid	C C		
RECTAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 17 DAY		Dysphagia		Klonopin Tablets	PS	Roche	
Initial or Prolonged 28 DAY				Neurontin	SS		
28 DAY				Risperdal	SS		
28 DAY				Cogentin	C		

Freedom Of Information (FOI) Report

TAKEN WITH A MEAL.

TAKEN AT BEDTIME.

14 DAY

Lopressor C
 Heparin C
 Inderal C
 Pepcid C
 Asa C

Isordil C
 Flomax C

Colace C
 Lotrimin Cream 1% C

Date:06/18/01ISR Number: 3741508-1Report Type:Expedited (15-DaCompany Report #260859
 Age:64 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2 MG 2 PER		Dysphagia Hyporeflexia	Health Professional	Clonopin	PS	Hoffmann La Roche Inc	ORAL
DAY ORAL				Neurontin (Gabapentin) 400 Mg	SS		ORAL
400 MG 2 PER							
DAY ORAL				Risperdal (Risperdal) 2 Mg	SS		ORAL
2 MG 2 PER							
DAY ORAL				Cogentin (Benztropine Mesylate)	C		
				Lopressor (Metoprolol Tartrate)	C		
				Heparin (Heparin Sodium)	C		

Inderal (Propranolol Hydrochloride) C
 Pepcid (Famotidine) C
 Asa (Aspirin) C
 Isordil (Isosorbide Dinitrate) C
 Flomax (Tamsulosin Hydrochloride) C
 Colace (Docusate Sodium) C
 Lotrimin Cream 1% (Clotrimazole) C

Date:06/18/01ISR Number: 3741692-XReport Type:Expedited (15-DaCompany Report #055-0945-M0100025
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200 MG, PER	Nosocomial Infection	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Life-Threatening		Pneumonia	Consumer				
Hospitalization - Initial or Prolonged							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/18/01ISR Number: 3741700-6Report Type:Expedited (15-DaCompany Report #001-0073-M0100251

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Glucose Increased	Health	Dilantin-125	PS	Parke Davis Div	
		Cerebrovascular Accident	Professional			Warner Lambert Co	
		Convulsion		Neurontin			
		Depression		(Gabapentin)	SS		
		Drug Level Below		Celexa (Citalopram			
		Therapeutic		Hydrobromide)	SS		

Date:06/18/01ISR Number: 3741734-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100716

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Akinesia	Consumer	Neurontin			
		Amnesia		(Gabapentin)	PS	Pfizer Inc	ORAL
300 MG		Dry Mouth					
(DAILY), PER		Fatigue					
ORAL		Movement Disorder		Sinequan (Doxipin			
		Muscle Twitching		Hcl)	C		ORAL
PER ORAL		Sedation		Antivert (Nicotinic			
		Thirst		Acid,Meclozine Hcl)	C		ORAL
PER ORAL		Tremor		Tylenol			
				(Paracetamol)	C		ORAL
PER ORAL							

Date:06/19/01ISR Number: 3742226-6Report Type:Direct

Company Report #

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dysarthria		Seroquel	PS		
Initial or Prolonged		Lethargy		Gabapentin	SS		
Other		Overdose					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 30.00 MG		Drug Ineffective	Consumer	Procardia Xl	PS	Pfizer Laboratories Div Pfizer Inc	ORAL
Prevent Permanent Impairment/Damage RAL		Joint Stiffness					
		Migraine					
900.00 MG		Tongue Disorder		Neurontin	SS		ORAL
TOTAL:DAILY:O							
TOTAL:TID:ORA							

L

Coumadin	C
Corzide	C
Lipitor	C
Folate	C
Valium	C
Aspirin	C
Klor-Con	C
Vitamin C	C
B-Complex	C
Vitamin E	C
Soy Supplement	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/19/01ISR Number: 3742511-8Report Type:Expedited (15-DaCompany Report #001-0945-M0100747

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 900 MG Initial or Prolonged		Angiopathy	Literature	Neurontin	PS	Pfizer Inc	
5 MG (NIGHT), 20 MG (NIGHT),		Intentional Misuse Priapism Suicide Attempt	Health Professional	Olanzapine (Olanzapine)	SS		
				Paroxetine (Paroxetine)	SS		

Date:06/19/01ISR Number: 3742771-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100731

Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN (TID), PER ORAL		Embolism Fatigue Oedema Peripheral Weight Increased	Consumer	Neurontin	PS	Pfizer Inc	ORAL
				Atenolol (Atenolol) Maxzide (Hydrochlorothiazide , Triamterene)	C C		

Date:06/19/01ISR Number: 3742955-4Report Type:Expedited (15-DaCompany Report #055-0945-M0100027

Age:58 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 400 MG (DAILY), PER ORAL		Hypersensitivity Injection Site Extravasation	Foreign Consumer	Neurontin	PS	Pfizer Inc	ORAL

Neuritis

Theophylline C
Bamifylline
Hydrochloride C
Chlorpromazine C
Amitriptyline
Hydrochloride C

Date:06/21/01ISR Number: 3743970-7Report Type:Expedited (15-DaCompany Report #001-0073-M0100251
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Increased	Health	Dilantin-125	PS	Parke Davis Div	
Other		Cerebrovascular Accident	Professional	Neurontin		Warner Lambert Co	
		Convulsion		(Gabapentin)	SS		
		Drug Level Below		Celexa (Citalopram			
		Therapeutic		Hydrobromide)	SS		

Date:06/21/01ISR Number: 3744743-1Report Type:Expedited (15-DaCompany Report #049-0945-M0100057
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adrenal Insufficiency	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Other		Blood Pressure Decreased	Health				
SEE IMAGE		Condition Aggravated	Professional				
		Dizziness					
		Fatigue					
		Gait Disturbance					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/22/01ISR Number: 3745486-0Report Type:Expedited (15-DaCompany Report #001-0945-M0100732

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin	PS	Pfizer Inc	
Other		Surgery		Soma (Carisoprodol)	C		
				Percocet (Oxycodone Hydrochloride, Paracetamol)	C		
				Paxil (Paroxetine Hydrochloride)	C		
				Valium (Diazepam)	C		
				Trazodone (Trazodone)	C		

Date:06/22/01ISR Number: 3745487-2Report Type:Expedited (15-DaCompany Report #055-0945-M0100026

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign	Neurontin	PS	Pfizer Inc	ORAL
Other		Blood Growth Hormone					
PER ORAL		Increased Erythema Multiforme Hypersensitivity	Health Professional				

Date:06/22/01ISR Number: 3745511-7Report Type:Expedited (15-DaCompany Report #001-0945-M0100722

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin	PS	Pfizer Inc	
Hospitalization - (SEE IMAGE)		Anorgasmia					
Initial or Prolonged		Arterial Occlusive Disease Chest Pain		Lopressor (Metoprolol Tartrate)	SS		ORAL
PER ORAL		Disturbance In Attention Hiatus Hernia Mania Memory Impairment Myocardial Infarction					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Anorectal Disorder	Health	Neurontin	PS	Pfizer Inc	
Initial or Prolonged	Colitis Ulcerative	Professional	Celexa (Citalopram			
Other	Dehydration		Hydrobromide)	SS		ORAL
DAILY, PER						
Required	Diarrhoea Haemorrhagic					
ORAL						
Intervention to	Haematocrit Decreased		Risperdal			
Prevent Permanent	Haemoglobin Decreased		(Risperidone)	SS		
Impairment/Damage	Large Intestinal Ulcer		Asacol (Mesalazine)	SS		
	Nausea		Prednisone			
	Pain		(Prednisone)	SS		
	Rectal Haemorrhage		Solu-Medrol			
	Rectal Ulcer		(Methylprednisolone			
	Vomiting		Sodium Succinate)	SS		
			Cyclosporine	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/25/01ISR Number: 3747316-XReport Type:Expedited (15-DaCompany Report #001-0945-M0100729

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Fatigue Medication Error Overdose	Consumer	Neurontin	PS	Pfizer Inc	

Date:06/25/01ISR Number: 3747333-XReport Type:Expedited (15-DaCompany Report #001-0945-M0100761

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Circulatory Collapse	Consumer	Neurontin	PS	Pfizer Inc	
2100 MG (QID)		Head Injury Headache Loss Of Consciousness Rash Pruritic Suicide Attempt		Prevacid (Lansoprazole)	C		

Date:06/26/01ISR Number: 3748127-1Report Type:Expedited (15-DaCompany Report #031-0945-M0100017

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Foreign	Neurontin	PS	Pfizer Inc	ORAL
PER ORAL		Loss Of Consciousness	Health	Baclofen (Baclofen)	SS		ORAL
PER ORAL			Professional				

Date:06/26/01ISR Number: 3748130-1Report Type:Expedited (15-DaCompany Report #055-0945-M0100029

Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Foreign	Neurontin	PS	Pfizer Inc	ORAL
300 MG							

		Pain		Consumer			
(DAILY), PER							
ORAL							
Date:06/26/01ISR Number: 3748131-3Report Type:Expedited (15-DaCompany Report #044-0945-M0100175							
Age:	Gender:Male	I/FU:I					
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Chromaturia	Foreign	Neurontin	PS	Pfizer Inc	ORAL
300 MG		Jaundice	Consumer				
(DAILY), PER		Oedema Peripheral	Other				
ORAL		Sedation		Oxycodone			
				(Oxycodone)	C		
				Omeprazole			
				(Omeprazole)	C		
				Meloxicam			
				(Meloxicam)	C		
				Amitriptyline			
				(Amitriptyline)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/26/01ISR Number: 3748234-3Report Type:Expedited (15-DaCompany Report #049-0945-M0100033

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Foreign	Neurontin	PS	Pfizer Inc	ORAL
2400 MG; 2700		Dialysis	Health				
MG (DAILY),		Muscle Spasms	Professional				
PER ORAL		Muscle Twitching	Other	Duragesic (Fentanyl)	SS		
TRANSDERMAL	75 TRANSD	Muscular Weakness		Sandimmune			
		Overdose		(Ciclosporin)	SS		
		Renal Failure Acute					
		Tremor					

Date:06/26/01ISR Number: 3748248-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100606

Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mouth Ulceration	Consumer	Neurontin	PS	Pfizer Inc	ORAL
600 MG (300							
MG, BID), PER							
ORAL							

Ambien (Zolpidem	
Tartrate)	C
Atenolol (Atenolol)	C
Ativan (Lorazepam)	C
Atrovent	
(Ipratropium	
Bromide)	C
Cardizem Cd	
(Diltiazem	
Hydrochloride)	C
Claritin	
(Loratadine)	C
Celexa (Citalopram	
Hydrobromide)	C
Celebrex (Celecoxib)	C
Diamox	

(Acetazolamide)	C
Nitroglycerin Cr (Glyceryl Trinitrate)	C
Plavix (Clopidogrel)	C
Prempro (Medroxyprogesterone Acetate, Estrogens Conjugated)	C
Prevacid (Lansoprazole)	C
Requip	C
Wellbutrin Sr (Amfebutamone Hydrochloride)	C
Zocor (Simvastatin)	C
Talacen (Paracetamol, Pentazocine Hydrochloride)	C
Aspirin (Acetylsalicylic Acid)	C
Diovan (Valsartan)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/26/01ISR Number: 3748463-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100754

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hangover Suicide Attempt	Consumer	Neurontin	PS	Pfizer Inc	

Date:06/28/01ISR Number: 3749482-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100758

Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Systemic Lupus	Health	Neurontin	PS	Pfizer Inc	ORAL
1500 MG		Erythematosis	Professional				
(BID), PER							

ORAL

Klonopin
(Clonazepam) C

Date:06/28/01ISR Number: 3749850-5Report Type:Expedited (15-DaCompany Report #001-0981-M0104435

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 10 MG (DAILY)		Drug Dependence	Health	Lipitor	PS	Pfizer Inc	
Initial or Prolonged Disability 2400 MG		Drug Withdrawal Syndrome Myalgia	Professional	Gabapentin (Gabapentin)	SS		

(DAILY)

(Diltiazem
Hydrochloride) C

Date:06/28/01ISR Number: 3749855-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100769

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia Disorientation Fall Tremor	Consumer	Neurontin	PS	Pfizer Inc	

Date:06/29/01ISR Number: 3750630-5Report Type:Expedited (15-DaCompany Report #033-0945-M0100075
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - SEE IMAGE Initial or Prolonged Other		Asthenia Cholecystitis Dyspnoea Encephalitis Fatigue Hepatocellular Damage Hypersensitivity Oedema	Foreign Health Professional	Neurontin	PS	Pfizer Inc	

Date:06/29/01ISR Number: 3750679-2Report Type:Expedited (15-DaCompany Report #001-0981-M0103661
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 10 MG (BID), Initial or Prolonged PER ORAL SEE IMAGE,		Unevaluable Event	Health Professional	Lipitor Gabapentin	PS SS	Pfizer Inc	ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

PER ORAL				
SEE IMAGE		Geodon	SS	ORAL
325 MG		Acetylsalicylic Acid	SS	ORAL
(DAILY), PER				
ORAL		Chlorpromazine	SS	ORAL
100 MG (EVERY				
4 HOURS PRN),				
PER ORAL		Clonazepam	SS	ORAL
2 MG (AT				
BEDTIME), PER				
ORAL		Lorazepam	SS	ORAL
2 MG (EVERY 6				
HOURS), PER				
ORAL, 2 MG				
(EVERY 8				
HOURS AS		Mesoridazine	SS	
SEE IMAGE				

Date:06/29/01ISR Number: 3750712-8Report Type:Expedited (15-DaCompany Report #2012999
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	20 MG BID PO	Death	Health	Oxycontin Cr	PS	Purdue Pharma Lp	ORAL
			Professional	Oxyir Capsules (Oxycodone Hydrochloride)	SS		ORAL
	MG PRN PO						

50 MG HS PO

Elavil (Amitriptylline) SS ORAL

24 MG QD PO

Neurontin (Gabapentin) SS ORAL

Soma (Carisoprodol) SS

Xanax (Alprazolam) SS

Date:07/02/01ISR Number: 3750941-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100779

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG (BID)		Cerebrovascular Accident	Health	Neurontin	PS	Pfizer Inc	
		Dysarthria Speech Disorder	Professional	Klonopin (Clonazepam)	C		
				Methylphenidate	C		
				Wellbutrin (Bupropion)	C		

Date:07/02/01ISR Number: 3750959-0Report Type:Expedited (15-DaCompany Report #001-0945-M0100808

Age:52 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Blood Potassium Decreased Depressed Level Of Consciousness Drug Level Above Therapeutic Eye Disorder Feeling Abnormal Hypoaesthesia Insomnia Paraesthesia Paralysis

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Freedom Of Information (FOI) Report

		Sedation Speech Disorder	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Neurontin	PS	Pfizer Inc	ORAL
100 MG UNK							
PER ORAL							
UNKNOWN	UNK, UNK, UNK			Dilantin (Phenytoin Sodium)	SS		
				Phenobartbital	C		
				Synthroid (Levothyroxine Sodium)	C		
				Inderal (Propranolol Hydrochloride)	C		
				Prilosec (Omeprazole)	C		
				Albuterol (Salbutamol)	C		
				Zyrtec (Cetirizine Hydrochloride)	C		
				Singulair (Montelukast)	C		
				Trimethoprim	C		
				Allopurinol	C		
				Viokase (Pancrelipase)	C		
				Phazyme (Pancreatin, Dimeticone, Activated, Pepsin, Diastase)	C		
				Climara (Estradiol)	C		
				Unspecified Vitamins	C		
				Hydrochlorothiazide	C		
				Voixx (Rofecoxib)	C		
				Colace (Docusate Sodium)	C		
				Peri-Colace (Docusate Sodium, Casanthranol)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Foreign	Neurontin	PS	Pfizer Inc	ORAL
1400 MG							
Other		Drug Interaction	Health				
(DAILY), PER							
ORAL		Viral Infection	Professional				
				Morphine (Morphine)	SS		
INTRAVENOUS	INTRAVENOUS			Carbamazepine			
				(Carbamazepine)	SS		ORAL
PER ORAL							

Date:07/02/01ISR Number: 3751123-1Report Type:Expedited (15-DaCompany Report #044-0945-M0100172
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Burning Sensation	Foreign	Neurontin	PS	Pfizer Inc	ORAL
1500 MG, PER							
Initial or Prolonged		Muscle Spasms	Health				
ORAL			Professional	Nortriptyline			
				(Nortriptyline)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/02/01ISR Number: 3751289-3Report Type:Expedited (15-DaCompany Report #048-0945-M0100003

Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1500 MG Initial or Prolonged (TID), PER ORAL		Psychotic Disorder Thinking Abnormal	Foreign Health Professional	Neurontin Carbamazepine	PS SS	Pfizer Inc	ORAL

Date:07/02/01ISR Number: 3751295-9Report Type:Expedited (15-DaCompany Report #002-0945-M0100047

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma Drug Interaction Drug Level Above Therapeutic Overdose Respiratory Depression	Foreign Health Professional	Neurontin Oxycodone Clonazepam	PS SS SS	Pfizer Inc	

Date:07/02/01ISR Number: 3751310-2Report Type:Expedited (15-DaCompany Report #002-0945-M0100045

Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death PER ORAL		Drug Interaction	Foreign Health Professional	Neurontin Zyprexia (Olanzapine) Hydromorphone	PS SS SS	Pfizer Inc	ORAL

Date:07/02/01ISR Number: 3751433-8Report Type:Expedited (15-DaCompany Report #001-0945-M0100592

Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 300 MG	Arthralgia	Consumer	Neurontin	PS	Pfizer Inc	ORAL
(DAILY), PER	Asthenia					
ORAL	Ecchymosis					
	Fall		Deltasone			
	Laceration		(Prednisone)	C		
	Myalgia		Glucotrol			
	Pain In Extremity		(Glipizide)	C		
	Paraesthesia		Avapro (Irbesartan)	C		
	Pyrexia		Hydrochlorthiazide			
	Sedation		(Hydrochlorothiazide			
	Tremor)	C		
	Vision Blurred					

Date:07/02/01ISR Number: 3751451-XReport Type:Expedited (15-DaCompany Report #001-0945-M0100784
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - SEE IMAGE	Coma	Consumer	Neurontin	PS	Pfizer Inc	
Initial or Prolonged	Communication Disorder Peripheral Vascular Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/02/01ISR Number: 3751749-5Report Type:Expedited (15-DaCompany Report #044-0945-M0100153

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 800 MG (BID), PER ORAL	Jaundice Sepsis	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Carbamazepine	PS C	Pfizer Inc	ORAL

Date:07/02/01ISR Number: 3751812-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100621

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Blood Albumin Increased Blood Creatinine Increased Burning Sensation Csf Protein Increased Nerve Conduction Studies Abnormal Paraparesis Radiculopathy	Literature Consumer	Neurontin Amitriptyline (Amitriptyline) Cyclosporin A (Ciclosporin) Prednisone Furosemide Famotidine Glibenclamide Zolpidem	PS SS SS C C C C C	Pfizer Inc	

Date:07/03/01ISR Number: 3752455-3Report Type:Expedited (15-DaCompany Report #01P-167-0108018-00

Age:32 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Disability 2.6 GM, 1 IN 1 D, PER ORAL; 2 GM, 1	Coombs Positive Haemolytic Anaemia	Foreign Health Professional Other	Depakene	PS	Abbott Laboratories Pharmaceutical Products Div	ORAL

IN 1 D, PER

ORAL

Gabapentin

SS

ORAL

2400 MG, 1 IN

1 D, PER ORAL

Folic Acid

C

Date:07/05/01ISR Number: 3752526-1Report Type:Direct

Company Report #

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly 900MG PO Q 8H Required Intervention to Prevent Permanent Impairment/Damage	Amniocentesis Abnormal Caesarean Section Complications Of Maternal Exposure To Therapeutic Drugs Movement Disorder Neonatal Disorder Premature Baby Renal Disorder		Neurontin 300mg Po	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/06/01ISR Number: 3753570-0Report Type:Direct
 Age:59 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Arrhythmia		Seroquel 300 Mg Bid			
Hospitalization -	300 MG BID PO 14 MON	Cataract		Po	PS		ORAL
Initial or Prolonged		Diabetes Mellitus		Neurontin 300mg Qid			
Required		Difficulty In Walking		Po	SS		ORAL
300MG QID PO 16 MON							
Intervention to		Faecal Incontinence					
Prevent Permanent		Hypertension					
Impairment/Damage		Obesity					
		Urinary Incontinence					
		Weight Increased					

Date:07/06/01ISR Number: 3753572-4Report Type:Direct
 Age:43 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hypoglycaemia		Neurontin 50 Mg	PS		ORAL
ONE PILL 4 X							
A DAY ORAL		Hypoglycaemic Coma					

Date:07/06/01ISR Number: 3753774-7Report Type:Expedited (15-DaCompany Report #001-0945-M0100793
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abnormal Behaviour	Health	Neurontin	PS	Pfizer Inc	ORAL
7200 MG							
(DAILY), PER		Convulsion	Professional				
ORAL		Drug Interaction					
		Hostility		Ultram (Tramadol Hydrochloride)	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 9000 MG		Blood Pressure Increased	Consumer	Neurontin	PS	Pfizer Inc	
Initial or Prolonged PER ORAL		Depression		Valium (Diazepam)	SS		ORAL
		Drug Dependence		Extra Strength			
25-30 (HS),		Dysphagia		Tylenol Pm	SS		ORAL
PER ORAL		Feeling Jittery					
		Gait Disturbance		Unspecified			
		Nausea		Antidepressants	C		
		Overdose					
		Sexual Assault Victim					
		Suicidal Ideation					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG		Hepatic Cirrhosis	Foreign	Neurontin	PS	Pfizer Inc	
		Hepatitis B	Health Professional	Neurontin (Gabapentin)	SS		
900 MG				Amitriptyline (Amitriptyline)	SS		
				Magnesium Sulfate (Magnesium Sulfate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/06/01ISR Number: 3754520-3Report Type:Expedited (15-DaCompany Report #044-0945-M0100069

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Meningioma	Foreign	Neurontin	PS	Pfizer Inc	
Other		Post Procedural Complication	Health Professional	Valproate (Valproic Acid)	C		

Date:07/09/01ISR Number: 3754389-7Report Type:Direct

Company Report #

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG 2 TID		Dyspnoea		Gabapentin	PS		
Initial or Prolonged 300 MG 3 TID		Neutropenia		Gabapentin	SS		
		Night Sweats Pyrexia					

Date:07/09/01ISR Number: 3755095-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100810

Age:12 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (DAILY)		Abnormal Behaviour	Consumer	Neurontin	PS	Pfizer Inc	
		Back Disorder					
		Bipolar Disorder		Paxil (Paroxetine Hydrochloride)	SS		
		Disturbance In Social Behaviour		Albuterol (Salbutamol)	C		
		Dysphagia					
		Intervertebral Disc Protrusion					
		Pollakiuria					

Date:07/09/01ISR Number: 3756246-9Report Type:Expedited (15-DaCompany Report #002-0945-M0100083

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Retinitis Pigmentosa	Foreign Health Professional	Neurontin Phenytoin (Phenytoin) Hydrochlorothiazide (Hydrochlorothiazide) Pravastatin (Pravastatin)	PS SS C C	Pfizer Inc	

Date:07/10/01ISR Number: 3756036-7Report Type:Direct Company Report #
Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG, 100MG		Dermatitis		Gabapentin	PS		ORAL

HS ORAL

Date:07/10/01ISR Number: 3756215-9Report Type:Expedited (15-DaCompany Report #044-0945-M0100023
Age: Gender:Female I/FU:I

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Vision Blurred	Foreign Health

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

			Professional Company Representative	Product	Role	Manufacturer	Route
Dose	Duration			Neurontin	PS	Pfizer Inc	ORAL
1800 MG							
(DAILY), PER							
ORAL							

Date:07/10/01ISR Number: 3756292-5Report Type:Expedited (15-DaCompany Report #A108646
Age:25 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Geodon	PS	Pfizer Central Research	ORAL
Hospitalization - Initial or Prolonged PO		Abnormal Behaviour Drug Ineffective	Consumer Health				
300.00 MG		Mania	Professional	Seroquel	SS		ORAL
TOTAL:DAILY:0		Medication Error					
RAL		Schizophrenia		Neurontin (Gabapentin)	SS		
3600.00 MG							
TOTAL:TID				Beer	SS		ORAL
ORAL				Depakote Clonazepam	C C		

Date:07/11/01ISR Number: 3756721-7Report Type:Expedited (15-DaCompany Report #200113149DE
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Arava	PS	Aventis Pharmaceuticals Inc	ORAL
Other		Acute Psychosis Condition Aggravated	Foreign Health				
10 MG/DAY PO	4 DAY						

	Drug Interaction	Professional	Carbamazepine		
	Grand Mal Convulsion	Other	(Tegretal - Slow Release)	SS	ORAL
1600 MG/DAY					
PO					
			Gabapentin		
			(Neurontin)	SS	ORAL
3200 MG/DAY					
PO					
			Morphine Sulfate		
			(Mst)	C	
			Ibuprofen (Ibu)	C	
			Prednisone	C	

Date:07/12/01ISR Number: 3757011-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100769
Age:75 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Anxiety	Consumer	Neurontin	PS	Pfizer Inc	
Initial or Prolonged	Asthenia					
	Disorientation					
	Ecchymosis					
	Fall					
	Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/12/01ISR Number: 3757054-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100819

Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Arthralgia Pulmonary Oedema	Consumer Health Professional	Neurontin	PS	Pfizer Inc	

Date:07/12/01ISR Number: 3757062-4Report Type:Direct

Company Report #

Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG 1T BID		Dizziness		Neurontin (Gabapentin) (300mg)	PS		
				Flonase	C		
				Atenolol	C		
				Ranitidine	C		
				Albuterol	C		
				Zocor	C		
				Flomax	C		
				Roxicet	C		
				Hctz	C		
				Cardura	C		

Date:07/12/01ISR Number: 3757120-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100708

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG, DAILY		Anorexia Concussion Diarrhoea Dysmenorrhoea Epilepsy Fall Haemorrhage Medication Error Nausea	Consumer Health Professional	Neurontin	PS	Pfizer Inc	

Thirst
Vitreous Floaters

Date:07/12/01ISR Number: 3757124-1Report Type:Direct
Age:73 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 100MG PO HS Initial or Prolonged 2' X 1 WEEK	Urticaria		Neurontin	PS		ORAL

PRIOR TO

ADMISSION

Date:07/12/01ISR Number: 3757186-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100826
Age:46 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Other	Abdominal Pain Alopecia Decreased Appetite Diabetes Mellitus Diabetic Neuropathy

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective Excoriation Fall	Consumer	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL
1200 MG (600 MG, BID), PER ORAL		Herpes Zoster Infection Injection Site Thrombosis Liver Function Test Abnormal Nausea					
		Oral Intake Reduced Paralysis Sedation Sepsis Swelling		Dilantin (Phenytoin Sodium) Glucophage (Metformin Hydrochloride)	SS SS		ORAL
PER ORAL		Thyroid Disorder Vomiting		Atenolol (Atenolol) Xanax (Alprazolam) Oxycontin (Oxycodone Hydrochloride)	C C C		

Date:07/12/01ISR Number: 3757417-8Report Type:Expedited (15-DaCompany Report #032-0945-M0100003
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening FOUR TIMES A Hospitalization - DAY, PER ORAL Initial or Prolonged Other		Increased Bronchial Secretion Upper Respiratory Tract Infection	Foreign Health Professional Company Representative	Neurontin Carbamazepine Lysomycil Ranitidine Hydrochloride Zuclopendixol Domperidone Sodium Bicarbonate, Potassium Chloride, Sodium Chloride, Macrogol)	PS C C C C C C	Pfizer Inc	ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300 MG	Arthralgia	Consumer	Neurontin	PS	Pfizer Inc	ORAL
Initial or Prolonged (DAILY), PER	Asthenia					
Other ORAL	Ecchymosis					
	Fall		Deltasone			
	Laceration		(Prednisone)	C		
	Myalgia		Glucotrol			
	Pain		(Glipizide)	C		
	Pain In Extremity		Avapro (Irbesartan)	C		
	Paraesthesia		Hydrochlothiazide	C		
	Pyrexia					
	Sedation					
	Tremor					
	Vision Blurred					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/13/01ISR Number: 3758837-8Report Type:Expedited (15-DaCompany Report #033-0945-M0100065

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour Aggression Cognitive Disorder Condition Aggravated Confusional State Convulsion Electrocardiogram Abnormal Epilepsy Irritability	Foreign Health Professional	Neurontin (Donepezil Hydrochloride) (Hydroxyzine Hydrochloride)	PS C C	Pfizer Inc	

Date:07/13/01ISR Number: 3758924-4Report Type:Expedited (15-DaCompany Report #044-0945-M0100184

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (UNKNOWN), UNKNOWN		Dizziness Loss Of Consciousness Pain	Foreign Consumer Other	Neurontin Tramadol (Tramadol) Paracetamol (Paracetamol) Rofecoxib (Rofecoxib) Amitriptyline (Amitriptyline)	PS C C C C	Pfizer Inc	

Date:07/19/01ISR Number: 3761857-0Report Type:Expedited (15-DaCompany Report #002-0945-M0100056

Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other PER ORAL		Facial Palsy Headache	Foreign Consumer	Neurontin	PS	Pfizer Inc	ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 300 MG	Confusional State	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Initial or Prolonged (DAILY), PER	Dermatitis	Health				
ORAL	Oedema	Professional				
	Rash Erythematous	Other	Tibolone	C		
			Folic Acid	C		
			Imdur (Isosorbide Mononitrate)	C		
			Losec (Omeprazole)	C		
			Folic Acid	C		
			Imdur (Isosorbide Mononitrate)	C		
			Losec (Omeprazole)	C		
			Zocor (Simvastatin)	C		
			Asa (Acetylsalicylic Acid)	C		
			Methotrextrate	C		
			Sulphasalazine	C		
			Meloxicam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydroxychloroquine C
 Atenolol C
 Paroven (Troloxerutin) C

Date:07/19/01ISR Number: 3762319-7Report Type:Expedited (15-DaCompany Report #044-0945-M0100158
 Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fungal Infection	Foreign	Neurontin			
		Hepatic Cirrhosis	Health	(Gabapentin)	PS	Pfizer Inc	
600 MG							
(DAILY),900		Hepatitis B	Professional				
			Other				
MG DAILY				Amitriptyline	SS		ORAL
50- 100MG							
(ONCE DAILY)							
, PER ORAL				Diclofenac	C		
				Magnesium Sulfate	C		
				Morphine	C		

Date:07/19/01ISR Number: 3762375-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100622
 Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cardiac Disorder	Consumer	Neurontin	PS	Pfizer Inc	ORAL
300 MG (TID),							
Initial or Prolonged		Fall	Health				
PER ORAL							
		Lower Limb Fracture	Professional	Toprol Xl			
				(Metoprolol			
				Succinate)	C		
				Zestril (Lisinopril)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia	Consumer	Neurontin	PS	Pfizer Inc	ORAL
200 MG BID		Convulsion					
PER ORAL		Difficulty In Walking					
		Gait Disturbance					
		Movement Disorder					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Mouth	Consumer	Tylenol	PS	Mcneil Consumer Products Co Div	
500 MG, OD,		Headache				Mcneilab Inc	ORAL
		Nausea					
PO	YR			Neurontin	SS		ORAL
300 MG,							
ONCE, PO							
1 DOSE							
				Valium	C		
				Tagamet	C		
				Roxicet	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/20/01ISR Number: 3762896-6Report Type:Expedited (15-DaCompany Report #049-0945-M0100064

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 600 MG, PER		Dyskinesia	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Initial or Prolonged ORAL		Muscle Twitching	Health Professional	Methadone (Methadone) Diazepam (Diazepam) Aponal (Doxepin Hydrochloride)	C C C		

Date:07/20/01ISR Number: 3762908-XReport Type:Expedited (15-DaCompany Report #001-0073-M0100251

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Glucose Increased Cerebrovascular Accident	Health Professional	Dilantin-125	PS	Parke Davis Div Warner Lambert Co	
		Convulsion Depression Drug Level Above Therapeutic Drug Level Below Therapeutic		Neurontin (Gabapentin) Celexa (Citalopram Hydrobromide)	SS SS		

Date:07/20/01ISR Number: 3762943-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100592

Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG, Initial or Prolonged (DAILY), PER		Amnesia	Consumer	Neurontin	PS	Pfizer Inc	ORAL
Other ORAL		Arthralgia Asthenia Disorientation Drug Hypersensitivity Ecchymosis		Deltasone (Prednisone) Glucotrol	C		

Electrocardiogram St	(Glipizide)	C
Segment Depression	Avapro (Irbesartan)	C
Fall	Hydrodiuril	
Fear	(Hydrochlorothiazide	
Headache)	C
Laceration	Oxycodone	
Myalgia	(Oxycodone)	C
Nausea		
Pain In Extremity		
Pallor		
Paraesthesia		
Pyrexia		
Sedation		
Tremor		
Visual Disturbance		

Date:07/23/01ISR Number: 3763579-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100876

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Priapism	Health Professional	Neurontin	PS	Pfizer Inc	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/23/01ISR Number: 3763584-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100881

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Cardiac Disorder	Consumer	Neurontin	PS	Pfizer Inc	
900 MG		Cleft Lip		Klonopin			
		Cleft Palate		(Clonazepam)	SS		
2 MG		Complications Of Maternal Exposure To Therapeutic		Tegretol			
				(Carbamazepine)	SS		
800 MG		Drugs		Topamax (Topiramate)	SS		
200 MG		Neonatal Disorder Pregnancy					

Date:07/23/01ISR Number: 3763620-3Report Type:Expedited (15-DaCompany Report #034-0945-M0100007

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Optic Atrophy	Foreign	Neurontin	PS	Pfizer Inc	ORAL
100 MG TID,		Papilloedema	Health				
Initial or Prolonged			Professional	Baclofen (Baclofen)	SS		ORAL
PER ORAL				Paroxetine			
Other				(Paroxetine)	SS		ORAL
10 MG TID,							
PER ORAL				Omeprazol			
20 MG (20 MG,				(Omeprazole)	SS		ORAL
DAILY), PER							
ORAL							
20 MG (20 MG,							
DAILY), PER							
ORAL							

100 MG (100

MG, DAILY),

PER ORAL

Levothyroxine
(Levothyroxine)

SS

ORAL

Date:07/23/01ISR Number: 3763751-8Report Type:Expedited (15-DaCompany Report #001-0945-M0100866

Age:80 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG		Blood Bicarbonate Decreased	Literature Health	Neurontin Celebrex (Celecoxib)	PS SS	Pfizer Inc	
		Blood Potassium Increased Haemodialysis Pulmonary Oedema Renal Failure Acute	Professional	Isosorbide Dinitrate (Isosorbide Dinitrate) Insulin (Insulin) Norvasc (Amlodipine Besilate) Coumadin (Warfarin Sodium) Desyrel (Trazodone Hydrochloride) Atenolol (Atenolol) Erythropoietin (Erythropoietin) Lansoprazole (Lansoprazole)	SS SS SS SS SS SS SS SS SS SS		

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Metoprolol C
Morphine C

Date:07/25/01ISR Number: 3764360-7Report Type:Expedited (15-DaCompany Report #A0154816A
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea	Consumer	Imitrex	PS	Glaxo Wellcome	
SUBCUTANEOUS		YR					
		Eye Haemorrhage		Imitrex	SS	Glaxo Wellcome	ORAL
YR							
		Meningioma		Neurontin	SS		
2700MG Per							
day	6	MON					
		Migraine					
		Vomiting					

Date:07/26/01ISR Number: 3765795-9Report Type:Direct Company Report #
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Neutrophil Count		Clozaril	PS		ORAL
625MG/D PO							
Initial or Prolonged		Decreased		Neurontin?	SS		
		Psychotic Disorder		Synthroid	C		
		White Blood Cell Count		Lotensin	C		
		Decreased		Ditropan	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Neurontin C

Date:07/26/01ISR Number: 3766326-XReport Type:Expedited (15-DaCompany Report #049-0945-M0100058
 Age:79 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 200 MG	Constipation	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Other (DAILY); 500	Cough	Health				
MG, PER ORAL	Decreased Appetite	Professional				
	Dysphagia		Magnesium Verla (Magnesium			
	Face Oedema		Glutamate)	C		
	Fatigue		Plavix (Clopidogrel)	C		
	Joint Swelling		Dytide			
	Localised Oedema		(Benzthiazide,			
	Oedema		Triamterene)	C		
	Oedema Peripheral		Cetebe (Ascorbic			
	Oliguria		Acid)	C		
	Sedation		Laxoberal (Sodium			
	Urinary Retention		Picosulfate)	C		
			Pantozol			
			(Pantoprazole			
			Sodium)	C		
			Ergenyl Chrono			
			(Valproic Acid,			
			Valproate Sodium)	C		
			Dantamacrin			
			(Dantrolene Sodium)	C		
			Novadral Retard			
			(Norfenefrine			
			Hydrochloride)	C		

Date:07/26/01ISR Number: 3766327-1Report Type:Expedited (15-DaCompany Report #031-0945-M0100017
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other PER ORAL	Drug Interaction	Foreign	Neurontin	PS	Pfizer Inc	ORAL

PER ORAL	Loss Of Consciousness	Health	Baclofen	SS	ORAL
		Professional	Insuline (Insulin Human)	C	
			Dantroleen	C	
			Morfine Sulfate	C	
			Paracetamol	C	
			Metoprolol	C	
			Cabasalate Calcium	C	
			Flucloxacilline	C	
			Augmentin (Clavulanate Potassium, Amoxicillin Trihydrate)	C	
			Nadroparine	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/26/01ISR Number: 3766337-4Report Type:Expedited (15-DaCompany Report #002-0945-M0100008

Age:76 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Abdominal Pain	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Hospitalization - 300 MG (TID), PER ORAL	Coronary Artery Disease	Health				
Initial or Prolonged	Myocardial Infarction	Professional	Tranylcypromine	C		
	Syncope	Company	Lorazepam	C		
		Representative	Pinaverium	C		

Date:07/26/01ISR Number: 3766401-XReport Type:Expedited (15-DaCompany Report #A117293

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged 100.00 MG	Drug Effect Decreased Drug Interaction	Consumer	Viagra	PS	Pfizer Agricultural Div	ORAL
TOTAL:PRN:ORA	Inhibition					
L	Headache					
ORAL	Nasal Congestion		Gabapentin	SS		ORAL
	Pain		Cardizem Cd	C		
	Parathyroid Disorder		Detrol	C		
			Aspirin	C		
			Prinivil	C		

Date:07/26/01ISR Number: 3766625-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100716

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other 1200 MG	Amnesia	Consumer	Neurontin	PS	Pfizer Inc	ORAL
(BID), PER	Coma	Health				
ORAL	Dry Mouth	Professional				

Dysarthria
 Loss Of Consciousness
 Muscle Twitching
 Sedation
 Thirst
 Tremor

Sinequan (Doxepin
 Hydrochloride) C
 Antivert (Nicotinic
 Acid, Meclozine
 Hydrochloride) C
 Tylenol
 (Paracetamol) C
 Zanaflex
 (Tizanidine) C
 Ultram (Tramadol
 Hydrochloride) C

Date:07/27/01ISR Number: 3766680-9Report Type:Expedited (15-DaCompany Report #002-0945-M0100083
 Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Retinitis Pigmentosa	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Dose			Health				
1600 MG			Professional				
Other							
(BID), PER							
ORAL							
				Phenytoin (Phenytoin)	SS		ORAL
300 MG (TID),							
PER ORAL							
				Hydrochlorothiazide (Hydrochlorotihiazid e)	C		
				Pravastatin (Pravastatin)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/27/01ISR Number: 3766697-4Report Type:Expedited (15-DaCompany Report #051-0073-M0100003

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300 MG	Coordination Abnormal	Foreign	Dilantin-125	PS		ORAL
Initial or Prolonged (UNKNOWN),	Dizziness	Health				
PER ORAL	Drug Level Above	Professional				
300 MG	Therapeutic		Neurontin	SS	Pfizer Inc	ORAL
(UNKNOWN) PER	Gait Disturbance					
ORAL	Nystagmus					

Date:07/27/01ISR Number: 3767126-7Report Type:Expedited (15-DaCompany Report #A0154816A

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other SUBCUTANEOUS	Diarrhoea	Consumer	Imitrex	PS	Glaxo Wellcome Inc	
DURATION SUBCUTANEOUS	SEE TEXT/ Drug Effect Decreased					
ORAL	Eye Haemorrhage Meningioma Vomiting		Imitrex Tablet (Sumatriptan Succinate)	SS		ORAL
2700 MG/PER DAY			Gabapentin (Formulation Unknown) (Gabapentin)	SS		

Date:07/27/01ISR Number: 3769687-0Report Type:Periodic

Age:57 YR Gender:Female I/FU:I

Company Report #001-0945-M0100555

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tooth Discolouration	Health Professional	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL

900 MG (TID),

PER ORAL

Fosinopril Sodium	C
Estropipate	C
Colestipol Hydrochloride)	C

Date:07/27/01ISR Number: 3769688-2Report Type:Periodic Company Report #001-0945-M0100567
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetes Mellitus Overdose	Consumer	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL

3200 MG, PER

ORAL

Date:07/27/01ISR Number: 3769689-4Report Type:Periodic Company Report #001-0945-M0100607
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Sedation	Consumer	Neurontin	PS	Parke Davis Pharmaceutical Research Div Warner Lambert Co	ORAL

PER ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/30/01ISR Number: 3767821-XReport Type:Expedited (15-DaCompany Report #055-0945-M0100017

Age:36 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 400 MG	Bone Pain	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Initial or Prolonged (DAILY), PER	Condition Aggravated	Consumer				
ORAL	Intervertebral Disc Disorder Oedema Peripheral Pain In Extremity Thrombosis		Unspecified Medications	C		

Date:07/30/01ISR Number: 3767823-3Report Type:Expedited (15-DaCompany Report #031-0945-M0100007

Age: Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 900 MG (TID),	Condition Aggravated	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Initial or Prolonged PER ORAL	Dyspnoea	Consumer				
	Lung Disorder	Health Professional	(Other Pulmonal Medication)	C		

Date:07/30/01ISR Number: 3767825-7Report Type:Expedited (15-DaCompany Report #049-0945-M0100065

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization -	Condition Aggravated	Foreign	Neurontin	PS	Pfizer Inc	
Initial or Prolonged Other	Deafness	Consumer	Celebrex (Celecoxib)	C		

Date:07/30/01ISR Number: 3768873-3Report Type:Expedited (15-DaCompany Report #A117697

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 5.00 MG TOTAL		Anxiety	Health	Zyrtec	PS	Pfizer Inc	
Intervention to DAILY		Blood Cholesterol	Professional				
Prevent Permanent Impairment/Damage		Increased Fatigue		Gabapentin Atorvastatin	SS SS		
10.00 MG		Gastrooesophageal Reflux					
TOTAL DAILY		Disease Weight Increased		Prozac Levo Thyroxin Methylphenidate Estratest Omeprazole Alprazolam Loratadine Lisinopril	SS C C C C C C C		

Date:07/30/01ISR Number: 3768879-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100898
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - PER ORAL		Drug Ineffective	Consumer	Neurontin	PS	Pfizer Inc	ORAL
Initial or Prolonged		Drug Interaction Headache		Viagra (Sildenafil Citrate)	SS		ORAL
100 MG (AS NEEDED), PER ORAL		Nasal Congestion Parathyroid Disorder		Cardizem Cd(Diltiazem			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C
 Detrol(Tolterodine
 Tartrate) C
 Aspirin
 (Acetylsalicylic
 Acid) C
 Prinivil
 (Lisinopril) C

Date:07/31/01ISR Number: 3777690-XReport Type:Periodic
 Age:42 YR Gender:Female I/FU:I

Company Report #2001053072US

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG, PRN, ORAL		Cold Sweat Drug Interaction	Consumer	Celebrex	PS	Gd Searle And Co	ORAL
400 MG, TID, ORAL		Flushing Hyperhidrosis Nausea		Neurontin (Gabapentin)	SS		ORAL
		Oedema Peripheral Rash Erythematous		Vitamin B6	C		

Date:07/31/01ISR Number: 3777708-4Report Type:Periodic
 Age:56 YR Gender:Female I/FU:I

Company Report #2001055409US

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG, BID, ORAL		Arterial Disorder Difficulty In Walking	Consumer	Celebrex	PS	Gd Searle And Co	ORAL
600 MG, BID			Professional	Neurontin (Gabapentin)	SS		
				Skelatin (Metaxalone)	C		
				Unspecified Hormone	C		

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG, QD, ORAL		Amnesia Disorientation	Consumer Health	Celebrex	PS	Gd Searle And Co	ORAL
900 MG, TID, ORAL		Feeling Abnormal Mental Impairment	Professional	Neurontin (Gabapentin)	SS		ORAL
10 MG, QD, ORAL		Oedema Peripheral Pain In Extremity		Lipitor (Atorvastatin)	SS		ORAL
				Narcotic Pain Medication (Unspecified) Duragesic	SS C		

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG /D PO		Oedema Peripheral		Neurontin	PS		ORAL
				Wellbutrin Ativan	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/01/01ISR Number: 3768503-0Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State		Neurontin	PS		ORAL
100MG PO TID		Disorientation		Risperdal	SS		ORAL
0.5MG PO BID		Dysphagia					
		Lethargy					

Date:08/01/01ISR Number: 3769116-7Report Type:Expedited (15-DaCompany Report #001-0945-M0100731
 Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Embolism	Consumer	Neurontin	PS	Pfizer Inc	ORAL
(TID), PER		Fatigue					
ORAL		Oedema Peripheral		Atenolol (Atenolol)	C		
		Sedation		Maxzide			
		Weight Increased		(Hydrochlorothiazide , Triamterene)	C		

Date:08/01/01ISR Number: 3769256-2Report Type:Expedited (15-DaCompany Report #046-0945-M0100016
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Activated Partial	Foreign	Neurontin	PS	Pfizer Inc	ORAL
100 MG		Thromboplastin Time	Health				
(DAILY), PER		Shortened	Professional				
ORAL; 900 MG		Drug Interaction					
(TID), PER				Warfarin (Warfarin Sodium)	SS		
ORAL							

Date:08/02/01ISR Number: 3770326-3Report Type:Expedited (15-DaCompany Report #044-0945-M0100193

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Inappropriate	Foreign	Neurontin	PS	Pfizer Inc	ORAL
PER ORAL		Antidiuretic Hormone Secretion	Health Professional				

Date:08/02/01ISR Number: 3770327-5Report Type:Expedited (15-DaCompany Report #044-0945-M0100194

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Hypercapnia	Foreign	Neurontin	PS	Pfizer Inc	ORAL
PER ORAL		Respiratory Failure	Health Professional Company Representative				

Date:08/02/01ISR Number: 3770401-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100897

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatic Encephalopathy Neurological Symptom	Health Professional	Neurontin (Gabapentin)	PS	Pfizer Inc	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/03/01ISR Number: 3770803-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100487
 Age:33 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 9000 MG	Balance Disorder	Consumer	Neurontin	PS	Pfizer Inc	
Initial or Prolonged PER ORAL	Dysphagia	Health	Valium (Daizepam)	SS		ORAL
50 GM (HS), PER ORAL	Hypertension Laceration Nausea	Professional	Extra Strength Tylenol Pm	SS		ORAL
	Overdose Suicidal Ideation Tremor		Unspecified Antidepressants	C		

Date:08/03/01ISR Number: 3771070-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100907
 Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 1500 MG	Bipolar Disorder	Consumer	Neurontin	PS	Pfizer Inc	ORAL
Initial or Prolonged (QID), PER ORAL	Diarrhoea Drug Interaction					
	Gastrointestinal Disorder Irritability Malaise Nausea Pain		Effexor (Venlafaxine Hydrochloride) Seroquel (Quetiapine) Depakote (Valproate Semisodium)	SS SS SS		

Date:08/03/01ISR Number: 3771100-4Report Type:Expedited (15-DaCompany Report #044-0945-M0100192
 Age:64 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other PER ORAL	Liver Function Test	Foreign	Neurontin	PS	Pfizer Inc	ORAL

Abnormal

Health
Professional

Unspecified Oncology
Treatment C

Date:08/06/01ISR Number: 3771482-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100915

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 1200 MG Initial or Prolonged	Chest Discomfort Chest X-Ray Abnormal Face Oedema Oedema Peripheral Pulmonary Hypertension	Consumer	Neurontin Lasix (Furosemide) Glucophage (Metformin Hydrochloride) Insulin (Insulin) Digoxin (Digoxin) Meclizine (Meclozine) Coumadin (Warfarin Sodium) Colace (Docusate Sodium)	PS C C C C C C C	Pfizer Inc	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/06/01ISR Number: 3771483-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100779

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident	Health	Neurontin	PS	Pfizer Inc	
1200 MG		Dysarthria	Professional				
(DAILY)		Speech Disorder		Klonopin (Clonazepam)	C		
				Methylphenidate	C		
				Wellbutrin (Bupropion)	C		

Date:08/06/01ISR Number: 3771923-1Report Type:Expedited (15-DaCompany Report #055-0945-M0100024

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cerebrovascular Accident	Foreign	Neurontin	PS	Pfizer Inc	ORAL
UNKNOWN, PER		Dysphagia	Consumer				
Hospitalization -		Metastasis					
ORAL							
Initial or Prolonged							

Date:08/06/01ISR Number: 3771940-1Report Type:Expedited (15-DaCompany Report #061-0945-M0100031

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiomyopathy	Foreign	Neurontin	PS	Pfizer Inc	ORAL
900 MG (TREE			Health				
TIMES DAILY),			Professional				
PER ORAL				Opioids	C		

Date:08/06/01ISR Number: 3772028-6Report Type:Expedited (15-DaCompany Report #055-0945-M0100011

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia	Foreign	Neurontin	PS	Pfizer Inc	ORAL
900 MG, PER							
Hospitalization -		Gastrointestinal Disorder	Consumer				
ORAL							
Initial or Prolonged		Metastasis					
Other		Oedema Peripheral					

Date:08/06/01ISR Number: 3772034-1Report Type:Expedited (15-DaCompany Report #049-0945-M0100047
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Creatinine	Foreign	Neurontin	PS	Pfizer Inc	ORAL
600 MG (TWICE							
DAILY), PER		Increased	Health				
ORAL		Blood Urea Increased	Professional				
		Dialysis		(Heparin-Fraction,			
		Glomerular Filtration		Sodium Salt)	C		
		Rate Decreased		Prednisone			
				(Prednisone)	C		
				Furorese			
				(Furosemide)	C		
				Erypo (Epoetin Alfa)	C		
				Tenormin (Atenolol)	C		
				Eryfer (Sodium			
				Bicarbonate, Ferrous			
				Sulfate, Ascorbic			
				Acid)	C		
				Prograf (Tacrolimus)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Magnesium
 (Magnesium) C
 Nephrotrans (Sodium
 Bicarbonate) C
 Rocaltrol
 (Calcitriol) C
 Asa (Acetylsalicylic
 Acid) C
 Benzbromarone
 (Benzbromarone) C
 Marcumar
 (Phenprocoumon) C
 Antra (Omeprazole) C
 Xanef (Enalapril
 Meleate) C
 Allopurinol
 (Allopurinol) C

Date:08/06/01ISR Number: 3772068-7Report Type:Expedited (15-DaCompany Report #033-0945-M0100104
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1800 MG (TID)	Amyotrophy	Foreign	Neurontin	PS	Pfizer Inc	
Initial or Prolonged Other	Balance Disorder Encephalopathy Gait Disturbance	Consumer	Lioresal (Baclofen) Rivotril (Clonazepam)	C C		

Date:08/06/01ISR Number: 3772071-7Report Type:Expedited (15-DaCompany Report #064-0945-M0100006
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other PER ORAL	Nephritis Interstitial	Foreign	Neurontin	PS	Pfizer Inc	ORAL
		Health Professional				

Date:08/06/01ISR Number: 3772336-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100758
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Systemic Lupus	Health	Neurontin	PS	Pfizer Inc	ORAL
1500 MG		Erythematosis	Professional				
(DAILY), PER							
ORAL							

Klonopin (Clonazepam)	C					
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Date:08/06/01ISR Number: 3772337-0Report Type:Expedited (15-DaCompany Report #001-0945-M0100849
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin	PS	Pfizer Inc	ORAL
200 MG		Difficulty In Walking					
(DAILY), PER		Extrapyrmidal Disorder					
ORAL		Gait Disturbance					
		Speech Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/06/01ISR Number: 3772338-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100916

Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Withdrawal Syndrome	Consumer	Neurontin	PS	Pfizer Inc	
2400 MG							
(THREE TIMES							
DAILY)							

Methadone
(Methadone) SS

Date:08/07/01ISR Number: 3771413-6Report Type:Direct

Company Report #

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Jaw Disorder		Neurontin/ 400 Mg/			
Initial or Prolonged		Tremor		Park Davis	PS	Park Davis	ORAL
800 MG/ TID/							
ORAL							

Catapres C
Lovenox C
Sonata C
Aspirin C
Potassium C
Motrin C
Paxil C
Oxycontin C
Lopressor C
Ativan C

Date:08/08/01ISR Number: 3772910-XReport Type:Expedited (15-DaCompany Report #049-0945-M0100046

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Creatinine	Foreign	Neurontin	PS	Pfizer Inc	ORAL
300 MG PER							

Initial or Prolonged
ORAL

Increased

Health

Blood Urea Increased
Glomerular Filtration
Rate Decreased
Oliguria
Renal Transplant

Professional

Ciclosporin
(Ciclosporin) C
Molsidomine
(Molsidomine) C
Tilidine (Tilidine) C
Naloxone (Naloxone) C
Prednisone
(Prednisone) C
Diltiazem
(Diltiazem) C
Ass (Acetylsalicylic
Acid) C
Calcium Brause C

Date:08/09/01ISR Number: 3774157-XReport Type:Expedited (15-DaCompany Report #064-0073-M0100001

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Nephritis Interstitial	Foreign Health Professional	Dilantin-125 Gabapentin (Gabapentin) Carbamazepine (Carbamazepine) Clobazam (Clobazam)	PS SS SS SS	Parke Davis Div Warner Lambert Co	

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Freedom Of Information (FOI) Report

Date:08/09/01ISR Number: 3774159-3Report Type:Expedited (15-DaCompany Report #049-0945-M0000028

Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	1200 MG	Cataract	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Other (THREE TIMES DAILY) , PER ORAL		Coma	Health				
		Drug Interaction	Professional				
		Feeling Drunk					
		Increased Appetite		Katadolon			
3 MG, PER , ORAL		Pleural Infection		(Flupirtine Maleate)	SS		ORAL
		Thrombosis					
		Tremor		(Nexium Mups)	C		
				(Prednisolone)	C		
				(Psyllium Hydrophilic Mucilloid)	C		
				(Colecalciferol, Calcium Carbonate)	C		
				Pancreatin)	C		
				Thyronajod (Levothyroxine Sodium)	C		
				Migraine Medication	C		
				Vioxx (Rofecoxib)	C		
				Karil "Norvatis"	C		
				Zantac (Ranitidine Hydrochloride)	C		
				Ergenyl "Labaz (Valproate Sodium)	C		
				Cyproterone Acetate (Cyproterone Acetate)	C		
				Laxoberal (Sodium Picosulfate)	C		
				Valproic Acid (Valproic Acid)	C		
				(Ass 100)	C		

Date:08/09/01ISR Number: 3774277-XReport Type:Expedited (15-DaCompany Report #001-0945-M0100227
Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haematuria Proteinuria	Health Professional	Neurontin (Gabapentin)	PS	Pfizer Inc	ORAL

3600 MG
(TID), PER

ORAL

Depakote (Valproate
Semisodium) C
Seroquel
(Quetiapine) C

Date:08/09/01ISR Number: 3774952-7Report Type:Expedited (15-DaCompany Report #001-0945-M0100592
Age:75 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Amnesia Arthralgia
Other	Asthenia Disorientation Electrocardiogram St

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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		Segment Depression					
		Fall					
		Fear					
		Headache	Report Source				
300 MG		Myalgia	Consumer	Neurontin	PS	Pfizer Inc	ORAL
(DAILY), PER		Nausea					
ORAL		Pallor					
		Paraesthesia		Deltasone			
		Sedation		(Prednisone)	C		
		Tremor		Glucotrol			
		Vision Blurred		(Glipizide)	C		
				Avapro (Irbesartan)	C		
				Hydrodiuril			
				(Hydrochlorothiazide			
)	C		
				Oxycodone	C		

Date:08/09/01ISR Number: 3775201-6Report Type:Expedited (15-DaCompany Report #044-0945-M0100199
 Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Balance Disorder	Foreign	Neurontin	PS	Pfizer Inc	ORAL
900 MG (TID),		Hyperglycaemia	Consumer				
Initial or Prolonged		Hypoglycaemia		Clopidogrel	C		
PER ORAL		Thirst		Ramipril	C		
Other				Furosemide	C		
				Digoxin	C		
				Amitriptyline	C		
				Loprazolam	C		

Date:08/09/01ISR Number: 3775240-5Report Type:Expedited (15-DaCompany Report #049-0945-M0100051
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chronic Hepatitis	Foreign	Neurontin	PS	Pfizer Inc	ORAL
1200 MG, PER							

ORAL	Haemosiderosis	Health		
	Hepatic Cirrhosis	Professional	(Phenprocoumon)	C
			(Metformin Hydrochloride)	C
			(Levothyroxine Sodium)	C

Date:08/09/01ISR Number: 3775330-7Report Type:Expedited (15-DaCompany Report #18000-025
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchopneumonia Coma Completed Suicide Overdose Pulmonary Congestion Pulmonary Oedema	Health Professional Other	Roxicodone Tablets (Oxycodone Hci Tablets Usp), 5mg Roxane Laboratorie Inc.	PS	Roxane Laboratories Inc.	
UNKNOWN		Syncope Toxicologic Test Abnormal		Oxycontin (Controlled Release Oxycodone Hcl) 40mg	SS	Purdue Pharma L.P.	
UNKNOWN				Carisoprodol	SS		
UNKNOWN				Ranitidine	SS		
UNKNOWN				Gabapentin	SS		
UNKNOWN				Propoxyphene	SS		

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UNKNOWN	Meprobamate	SS
UNKNOWN	Valproic Acid	SS
UNKNOWN	Caffeine Anhydride	SS
	Hydrocodone/Apap	C
	Prozac (Fluoxetine)	C
	Depakote (Divalproex Sodium)	C
	Skelaxin	C
	Zyprexa	C
	Remeron (Mirtazapine).	C

Date:08/13/01ISR Number: 3776081-5Report Type:Expedited (15-DaCompany Report #2000COU0810
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 5-10 MG QD PO		Fall Haematochezia	Consumer	Coumadin	PS	Dupont Merck Pharmaceutical Co	ORAL
		International Normalised Ratio Increased Spinal Fracture		Neurontin (Gabapentin) (Alendronate Sodium)	SS SS		ORAL
70 [MG] WEEKLY PO				Imdur (Isosorbide Mononitrate) Norvasc (Amlodipine Besylate) (Multivitamin) Percocet (Oxycodone/Acetamino phen) Thiamin (Thiamin Hydrochloride) (Magnesium Oxide)	C C C C C		

Date:08/13/01ISR Number: 3776161-4Report Type:Expedited (15-DaCompany Report #001-0945-M0100944
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG		Grand Mal Convulsion	Consumer	Neurontin	PS	Pfizer Inc	
(THREE TIMES A DAY)		Medication Error					

(Lorazepam)	C
Oxycodone Hydrochloride, Paracetaol)	C
(Zolpidem Tartrate)	C
(Fentanyl Patches)	C

Date:08/13/01ISR Number: 3776180-8Report Type:Expedited (15-DaCompany Report #055-0945-M0100019
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 400 MG		Condition Aggravated	Foreign	Neurontin	PS	Pfizer Inc	ORAL
Initial or Prolonged (DAILY), PER Disability ORAL		Decreased Appetite	Consumer				
		Insomnia					
		Varicose Vein		Unspecified Antihypertensive Agent	C		

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Date:08/13/01ISR Number: 3776209-7Report Type:Expedited (15-DaCompany Report #061-0945-M0100031

Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Cardiomyopathy	Foreign	Neurontin	PS	Pfizer Inc	ORAL
1600 MG		Ejection Fraction	Health				
Other		Abnormal	Professional				
(QID), PER		Mitral Valve Incompetence		Amitriptyline			
ORAL		White Blood Cell Count		(Amitriptyline)	C		
		Decreased		Asa (Acetylsalicylic			
				Acid)	C		
				Imuran			
				(Azathioprine)	C		
				Lasix (Furosemide)	C		
				Magensium Aspartate			
				(Magnesium Apartate)	C		
				Methadone Syrup			
				(Methadone0	C		
				Pethidine			
				(Pethidine)	C		
				Phenergan			
				(Promethazine			
				Hydrochloride)	C		
				Premarin (Estrogens			
				Conjugated)	C		
				Prepulsid			
				(Cisapride)	C		
				Temazepam			
				(Temazepam0	C		
				Zofran (Ondansetron			
				Hydrochloride)	C		

Date:08/15/01ISR Number: 3778324-0Report Type:Expedited (15-DaCompany Report #055-0945-M0100020

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Haemorrhage	Foreign	Neurontin	PS	Pfizer Inc	ORAL
1200 MG			Consumer				
Hospitalization -							
(DAILY), PER							

Initial or Prolonged
ORAL

Date:08/15/01ISR Number: 3778339-2Report Type:Expedited (15-DaCompany Report #044-0945-M0100204
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatitis Cholestatic	Foreign	Neurontin	PS	Pfizer Inc	ORAL
900 MG (300			Health				
MG, TID), PER			Professional				
ORAL							

Date:08/15/01ISR Number: 3778456-7Report Type:Direct Company Report #
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased		Neurontin / Gabapentin	PS		ORAL
900 MGM TID							
P.O.							

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Freedom Of Information (FOI) Report

Date:08/15/01ISR Number: 3778573-1Report Type:Expedited (15-DaCompany Report #002-0945-M0100103

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Foreign	Neurontin	PS	Pfizer Inc	ORAL
SEE IMAGE		Erection Increased	Consumer	Novo-Clobazam	C		
		Sedation		(Cefixime)	C		

Date:08/15/01ISR Number: 3778620-7Report Type:Expedited (15-DaCompany Report #044-0945-M0100203

Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Failure	Foreign	Neurontin	PS	Pfizer Inc	ORAL
1200 MG		Congestive	Consumer				
(DAILY), PER		Weight Increased					
ORAL				Digoxin	C		
				Warfarin	C		
				Insulin	C		
				Co-Proxamil			
				(Paracetamil,			
				Dextropropoxyphene			
				Hydrochloride)	C		

Date:08/16/01ISR Number: 3778681-5Report Type:Expedited (15-DaCompany Report #001-0945-M0101000

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Health	Neurontin	PS	Pfizer Inc	
		Myalgia	Professional				
		Pyrexia					
		Red Blood Cell					
		Sedimentation Rate					
		Increased					

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG, PER Initial or Prolonged ORAL		Hypomania	Foreign Health Professional	Neurontin	PS	Pfizer Inc	ORAL

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 400 MG		Cerebrovascular Accident	Consumer	Neurontin	PS	Pfizer Inc	
		Drug Ineffective		Adalat (Nifedipine)	C		
				Insulin (Insulin)	C		
				Lipitor (Atorvastatin)	C		
				Lasix (Furosemide)	C		
				Ambien (Zolpidem Tartrate)	C		

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Freedom Of Information (FOI) Report

Date:08/17/01ISR Number: 3779790-7Report Type:Direct
 Age:82 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300MG TID	Anaemia		Neurontin 300mg	PS		ORAL
Hospitalization - ORAL		Neutropenia					
Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Pancytopenia Thrombocytopenia					

Date:08/20/01ISR Number: 3780111-4Report Type:Direct
 Age: Gender: I/FU:I

Company Report #USP 54213

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Neurontin	PS	Pfizer	

Date:08/20/01ISR Number: 3780149-7Report Type:Direct
 Age: Gender: I/FU:I

Company Report #USP 54223

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Neurontin	PS	Pfizer	

Date:08/20/01ISR Number: 3780158-8Report Type:Direct
 Age: Gender: I/FU:I

Company Report #USP 54227

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Neurontin	PS	Pfizer	

Date:08/20/01ISR Number: 3780410-6Report Type:Expedited (15-DaCompany Report #049-0945-M0100047
 Age:32 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 600 MG (TWICE Initial or Prolonged DAILY) , PER Other ORAL	Blood Creatine Increased	Foreign	Neurontin	PS	Pfizer Inc	ORAL
	Blood Urea Increased	Health				
	Cyst	Professional				
	Dialysis Glomerular Filtration Rate Decreased		(Heparin-Fraction, Sodium Salt)	C		
			Prednisolone (Prednisolone)	C		
			Furorese (Furosemide)	C		
			Erypo (Epoetin Alfa)	C		
			Tenormin (Atenolol)	C		
			Eryfer (Sodium Bicarbonate, Ferrous Sulfate, Ascorbic Acid)	C		
			Prograf (Tacrolimus)	C		
			Magnesium (Magnesium)	C		
			Nephrotrans (Sodium Bicarbonate)	C		
			Rocaltrol (Calcitriol)	C		
			Asa (Acetylsalicylic Acid)	C		

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Benzbromarone
 (Benzbromarone) C
 Marcumar
 (Phenprocoumon) C
 Antra (Omeprazole) C
 Xanef (Enalapril
 Maleate) C
 Allopurinol
 (Allopurinol) C

Date:08/20/01ISR Number: 3780470-2Report Type:Direct
 Age:51 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abnormal Dreams Feeling Jittery		Gabapentin 100 Mg Parke-Davis	PS	Parke-Davis	ORAL
200 MG TID		Hallucination					
ORAL		Night Sweats		Gabapentin Nortriptyline Diazepam Morpine Felodipine Lisinopril Citalopram Oxycodone	C C C C C C C C		

Date:08/20/01ISR Number: 3780738-XReport Type:Expedited (15-DaCompany Report #001-0945-M0100993
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Hepatic Failure Ventricular Tachycardia	Health Professional Company Representative	Neurontin (Gabapentin)	PS	600 Mg (Daily),	

Date:08/20/01ISR Number: 3780782-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100996
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tongue Disorder	Health	Neurontin	PS	Pfizer Inc	ORAL
200 MG (BID),			Professional				
PER ORAL							

Date:08/20/01ISR Number: 3780798-6Report Type:Expedited (15-DaCompany Report #2001068228AU
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Oedema Peripheral	Foreign	Celebrex	PS	Gd Searle And Co	ORAL
200 MG, QD,		Pain In Extremity	Health				
Initial or Prolonged			Professional	Gabapentin(Gabapenti			
ORAL			Other	n)	SS		ORAL
1200 MG, QD,							
ORAL				Sinemet	C		

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Date:08/20/01ISR Number: 3780845-1Report Type:Expedited (15-DaCompany Report #033-0945-M0100107

Age:52 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 1800 MG Hospitalization - (DAILY), Initial or Prolonged UNKNOWN	Coma	Foreign Health Professional	Neurontin (Diazepam) (Clorazepate Dipotassium) (Diltiazem Hydrochloride) Codeine Phosphate Paracetamol (Ketoprofen)	PS C C C C C	Pfizer Inc	

Date:08/20/01ISR Number: 3780923-7Report Type:Expedited (15-DaCompany Report #044-0945-M0100194

Age: Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability	Hypercapnia Respiratory Failure	Foreign Health Professional Company Representative	Neurontin	PS	Pfizer Inc	

Date:08/23/01ISR Number: 3782420-1Report Type:Expedited (15-DaCompany Report #001-0945-M0100606

Age:77 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (300 MG, BID), PER ORAL	Mouth Ulceration	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Ambien (Zolpidem Tartrate)	C
Atenolol (Atenolol)	C
Ativan (Lorazepam)	C
Atrovent (Ipratropium Bromide)	C
Cardizem Cd (Diltiazem Hydrochloride)	C
Claritin (Loratadine)	C
Celexa (Citalopram Hydrobromide)	C
Celebrex (Celecoxib)	C
Diamox (Acetazolamide)	C
Nitroglycerin Cr (Glyceryl Trinitrate)	C
Plavix (Clopidogrel)	C
Prempro (Medroxyprogesterone Acetate, Estrogens Conjugated)	C
Prevacid	

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(Lansoprazole) C
 Requip C
 Wellbutrin Sr
 (Amfebutamone
 Hydrochloride) C
 Zocor (Simvastatin) C
 Talacen
 (Paracetamol,
 Pentazocine
 Hydrochloride) C
 Aspirin
 (Acetylsalicylic
 Acid) C
 Diovan (Valsartan) C

Date:08/23/01ISR Number: 3782740-0Report Type:Expedited (15-DaCompany Report #049-0945-M0100078
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State	Foreign	Neurontin			
		Metastases To Central	Health	(Gabapentin)	PS		ORAL
PER ORAL		Nervous System	Professional	(Fentanyl)	C		
		Prothrombin Time Abnormal					

Date:08/23/01ISR Number: 3782743-6Report Type:Expedited (15-DaCompany Report #031-0945-M0100020
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		International Normalised	Foreign	Gabapentin			
		Ratio Increased	Consumer	(Gabapentin)	PS		ORAL
300 MG (THREE		Prothrombin Time					
TIMES A DAY),		Prolonged					
PER ORAL				(Phenprocoumon)	SS		ORAL
3 MG, PER							
ORAL				(Quinapril)	C		
				(Glimepiride)	C		

(Ipratropium) C
(Bromhexine) C
(Doxycycline) C
(Atorvastatin) C

Date:08/23/01ISR Number: 3782778-3Report Type:Expedited (15-DaCompany Report #033-0366-M0100003

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Proteinuria	Foreign Consumer	Zarontin (Ethosuximide) Neurontin (Gapapentin)	PS SS		

2400 MG
(THREE TIMES
A DAY)

Date:08/24/01ISR Number: 3783096-XReport Type:Expedited (15-DaCompany Report #055-0945-M0100027

Age: Gender:Unknown I/FU:F

Outcome
Hospitalization -
Initial or Prolonged

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Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG	(DAILY), PER	Hypersensitivity	Foreign	Gabapentin	PS		ORAL
ORAL		Neuritis	Consumer				
		Thermal Burn					
				Theophylline (Theophylline)	C		
				Bamifylline Hydrochloride (Bamifylline Hydrochloride)	C		
				Chlorpromazine (Chlorpromazine)	C		
				Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		

Date:08/24/01ISR Number: 3783200-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100592
Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG	Arthralgia Asthenia	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other (DAILY, PER		Cardiac Disorder					
ORAL		Disorientation					
		Dizziness		Deltasone (Prednisone)	C		
		Ecchymosis		Glucotrol (Glipizide)	C		
		Electrocardiogram St Segment Depression		Avapro (Irbesartan)	C		
		Fall		Hydrodiuril (Hydrochlorothiazide			
		Fear)	C		
		Feeling Abnormal		Oxycodone			
		Head Injury		(Oxycodone)	C		
		Headache					
		Myalgia					

Nausea
 Pain
 Pallor
 Paraesthesia
 Pyrexia
 Sedation
 Tremor
 Visual Disturbance

Date:08/27/01ISR Number: 3783245-3Report Type:Expedited (15-DaCompany Report #2001-08-0004

Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hyperkalaemia Hypoxia Mental Impairment	Health Professional Company	Thalomid (Thalidomide 50 Mg) Capsules	PS		ORAL
100 MG QD		Pneumonia	Representative				
ORAL	11 DAY	Renal Failure Acute		Aredia Neurontin	SS SS		ORAL
300 MG TID							
ORAL				Naprosyn Tablets	SS		ORAL
PRN ORAL				Paxil	C		

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Vitamins Nos C
 Hydrocodone Compound C
 Methadone C
 Latanoprost C

Date:08/27/01ISR Number: 3783247-7Report Type:Expedited (15-DaCompany Report #2001-07-0134
 Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anaemia Arthritis Asthenia	Health Professional Company	Thalomid (Thalidomide 50 Mg) Capsules	PS		ORAL
400 MG QD		Constipation	Representative				
ORAL		Dehydration		Aredia	SS		
INTRAVENOUS	90 MG	Diarrhoea					
INTRAVENOUS		Flushing					
NOS		Hypocalcaemia		Neurontin	SS		
300 MG BID		Hypoproteinaemia		Arava	SS		
		Leukopenia		Bicnu	C		
		Nausea		Dexamethasone	C		
		Nephrotic Syndrome		Cytosan	C		
		Oedema		Biaxin	C		
		Oedema Peripheral					
		Orthostatic Hypotension					
		Renal Failure Acute					
		Small Intestinal					
		Obstruction					
		Stomatitis					
		Syncope					
		Thrombocytopenia					
		Tremor					

Date:08/27/01ISR Number: 3783364-1Report Type:Expedited (15-DaCompany Report #A114661
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Electrocardiogram T Wave Inversion Overdose Tachycardia	Health Professional	Ziprasidone Po Gabapentin	PS SS		

Date:08/27/01
 Age: Gender:Male I/FU:I
 ISR Number: 3783493-2
 Report Type:Expedited (15-DaCompany Report #001-0945-M0101017

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 3200 MG (TID), 400 MG (BID)		Chest Pain Drug Level Below Therapeutic Hypertension Pain Pyrexia Red Blood Cell Sedimentation Rate Increased Tachycardia	Health Professional	Neurontin (Gabapentin) Dilantin (Phenytoin Sodium)	PS SS		

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Date:08/27/01ISR Number: 3783495-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100900

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2700 MG (3 TIMESA DAY), PER ORAL		Convulsion Depressed Level Of Consciousness Drug Level Above Therapeutic Hypoxia Lethargy Physical Examination Abnormal	Foreign Health Professional	Neurontin (Gabapentin) Carnitor (Levocaristine) Toprol Xl (Metoprolol Succinate) Phenobarbital (Phenobarbital) Diovan Hct (Valsartan) Zantac (Ranitidine Hydrochloride0 Depakote (Valproate Semisodium) Albuterol (Salbutamol) Reglan (Metoclopramide)	PS C C C C C C C		ORAL

Date:08/27/01ISR Number: 3783500-7Report Type:Expedited (15-DaCompany Report #A117697

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 5.00 MG Intervention to TOTAL:DAILY Prevent Permanent Impairment/Damage 10.00 MG TOTAL:DAILY		Anxiety Blood Cholesterol Increased Fatigue Gastrooesophageal Reflux Disease	Health Professional	Zyrtec Tablets Gabapentin Atorvastatin Prozac	PS SS SS SS		

Hypertension
Weight Increased

Levo Thyroxin C
Methylphenidate C
Estratest C
Omeprazole C
Alprazolam C
Loratadine C
Lisinopril C

Date:08/27/01ISR Number: 3783536-6Report Type:Expedited (15-DaCompany Report #001-0945-M0101003

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (DAILY), PER ORAL		Arthritis Swelling Vein Disorder	Consumer	Neurontin (Gabapentin) Lasix (Furosemide)	PS C		ORAL

PO

(Oxycodone Hydrochloride)	SS	ORAL
Soma	SS	
Neurontin (Gabapentin)	SS	
Antidepressant (Unspecified)	SS	

Date:08/28/01ISR Number: 3784108-XReport Type:Expedited (15-DaCompany Report #FRA002698

Age:86 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG BID PO		Hyponatraemia	Foreign	Rythmol "Biosedra"	PS		ORAL
1 DF OD OTHER			Health	Nitridem Tts	SS		
1 DF OD PO			Professional	Coversyl	SS		ORAL
1 DF BID PO			Other	Neurontin	SS		ORAL
1 DF OD PO				Deroxat	SS		ORAL
1 DF QD PO				Aldalix	SS		ORAL

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Date:08/29/01ISR Number: 3785021-4Report Type:Expedited (15-DaCompany Report #HQ5232127AUG2001

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Level Above Therapeutic Drug Toxicity	Consumer	Effexor (Venlafaxine Hydrochloride, Tablet)	PS		ORAL
ORAL				Neurontin (Gabapentin,)	SS		

Date:08/29/01ISR Number: 3785081-0Report Type:Expedited (15-DaCompany Report #2012999

Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accident	Health	Oxycontin Cr			
		Cholecystectomy	Professional	Tablets, 20 Mg			
		Drug Toxicity	Other	(Oxycodone Hydrochloride)	PS		ORAL
20 MG BID PO		Fracture Malunion					
		Hepatic Steatosis		Oxyir Capsules			
		Hepatomegaly		(Oxycodone Hydrochloride)	SS		ORAL
MG PRN PO		Nephrosclerosis					
		Obesity		Elavil			
50 MG HS PO		Oedema		(Amitriptylline)	SS		ORAL
		Pulmonary Congestion		Neurontin			
24 MG QD PO		Pyelonephritis Chronic		(Gabapentin)	SS		ORAL
				Soma (Carisprodol)	SS		
				Xanax (Alprazolam)	SS		
				Hydrocodone			
				Bitartrate	SS		
				Ranitidine	SS		
				Caffeine	SS		
				Nicotine	SS		
				Trazodone	SS		
				Acetaminophen	SS		
				Nortriptyline	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation	Health	Ziprasidone Po	PS		ORAL
80.00 MG		Atrioventricular Block	Professional				
TOTAL; BID;		Gastritis					
ORAL		Haematemesis		Gabapentin	SS		ORAL
1200.00 MG		Insomnia					
TOTAL; BID;		Nausea					
ORAL		Sinus Tachycardia		Risperdal	C		
		Vomiting		Trazadone	C		
				Ativan	C		
				Cogentin	C		
				Multivitamin	C		

Outcome	PT
Hospitalization -	Coordination Abnormal
Initial or Prolonged	Difficulty In Walking
	Drug Ineffective

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG	(DAILY),	Hypoaesthesia Movement Disorder Pain	Consumer	Neurontin (Gabapentin)	PS		
		Sedation Sensation Of Heaviness		(Leflonomide)	C		
		Vision Blurred		(Pantoprazole)	C		
		Visual Disturbance		(Prasterone)	C		
				(Calcium)	C		
				(Prednisone)	C		
				(Phenylbutazone)	C		
				(Morphine)	C		
				(Infliximab)	C		
				(Fentanyl)	C		
				(Risedronic Acid)	C		
				(Erocalciferol)	C		

Date:08/29/01ISR Number: 3785385-1Report Type:Expedited (15-DaCompany Report #001-0945-M0101034
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG		Blood Pressure Increased Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other (QID), PER		Heart Rate Increased		(Pirbuterol Acetate)	SS		
ORAL							
RESPIRATORY							
(INHALATION)	INHALATION						

Date:08/29/01ISR Number: 3785387-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100916
Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Abdominal Pain Upper	Consumer	Neurontin	
2400 MG	Drug Interaction	Health	(Gabapentin)	PS
(THREE TIMES	Inhibition	Professional		
DAILY)	Drug Withdrawal Syndrome			
15 MG (THREE			(Methadone)	SS
TIMES DAILY)			Ibuprofen	C

Date:08/29/01ISR Number: 3785605-3Report Type:Expedited (15-DaCompany Report #046-0945-M0100016
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Interaction	Foreign	Neurontin			
SEE IMAGE		International Normalised	Health	(Gabapentin)	PS		ORAL
35 MG (TWICE		Ratio Increased	Professional	Warfarin (Warfarin			
A WEEK), PER		Pain In Extremity	Company	Sodium)	SS		ORAL
ORAL		Prothrombin Level	Representative				
		Decreased		(Paracetamol)	C		
				(Sodium Bicarbonate,			
				Potassium Chloride,			
				Sodium Chloride,			
				Macrogol)	C		
				(Morphine Sulfate)	C		
				(Sodium Picosulfate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/29/01ISR Number: 3785830-1Report Type:Expedited (15-DaCompany Report #049-0945-M0100076

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Circulatory Collapse	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG, PER		Depressed Level Of	Professional				
ORAL		Consciousness		Malarone (Atovaquone)	SS		ORAL
(DAILY), PER		Diarrhoea					
ORAL		Dissociation					
		Drug Interaction		(Ethinylestradiol			
		Loss Of Consciousness		Chlormadinone			
		Nausea		Acetate)	C		
		Sleep Disorder					

Date:08/30/01ISR Number: 3785246-8Report Type:Direct

Company Report #

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Chest Discomfort		Levoxyl 100 Mcg./0.1			
2 TABS PER		Chest Pain		Mg Walgreens	PS	Walgreens	ORAL
DAY ORAL		Dyspnoea					
		Medication Error		Neurontin 400 Mg.			
1 CAPSU PER		Palpitations		Walgreens	SS	Walgreens	ORAL
DAY ORAL							

Date:08/30/01ISR Number: 3785935-5Report Type:Direct

Company Report #

Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dyspnoea		Neurontin 300mg	PS		
600 MG TID							

Initial or Prolonged	Eosinophilia	Pepcid	C
Required	Hypoxia	Zosyn	C
Intervention to	Pulmonary Oedema	Morphine	C
Prevent Permanent	Pyrexia	Propofol	C
Impairment/Damage	Respiratory Distress	Lovenox	C
	Sinus Tachycardia	Tylenol	C
		Prevacid	C
		Elavil	C
		Lasix	C
		Levaquin	C
		Solu-Medrol	C
		Carapres-Tts	C
		Benadryl	C
		Phenergan	C

Date:08/31/01ISR Number: 3786123-9Report Type:Expedited (15-DaCompany Report #266750
Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Confusional State		Naprosyn	PS	Roche	
Initial or Prolonged	Movement Disorder		Thalomid	SS		
11 DAY						
	Pneumonia		Aredia	SS		
	Renal Failure		Neurontin	SS		
			Paxil	C		
			Vitamin Nos	C		
			Hydrocodone	C		
			Methadone	C		
			Latanoprost	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

274 DAY Alkeran C
 274 DAY Prednisone C

Date:08/31/01ISR Number: 3786793-5Report Type:Expedited (15-DaCompany Report #2001069647FR
 Age:86 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Gait Disturbance Hyponatraemia	Foreign Health Professional	Aldalix (Spironolactone, Furosemide) Tablet	PS		ORAL
1 UG, QD, ORAL			Other				
TRANSDERMAL	1, QD,			Nitriderm Tts (Glyceryl Trinitrate)	SS		
TRANSDERMAL							
150 MG, BID, ORAL				Rythmol "Knoll" (Propafenone Hydrochloride)	SS		ORAL
1 QD				Coversyl (Perindopril)	SS		
2 BID				Neurontin (Gabapentin)	SS		
20 MG, ORAL				Deroxat (Paroxetine Hydrochloride)	SS		ORAL

Date:08/31/01ISR Number: 3787103-XReport Type:Periodic Company Report #A114887
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other	Accidental Overdose	Health	Ziprasidone Po	PS
	Dermatitis	Professional	Neurontin (Gabapentin)	SS
			Depakote (Sodium Valproate)	SS

Date:08/31/01ISR Number: 3787300-3Report Type:Expedited (15-DaCompany Report #049-0945-M0100057
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Adrenal Insufficiency Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
PER ORAL;	1600						
MG(UNK);	1800	Dizziness	Professional				
MG(UNK)		Fatigue					
		Gait Disturbance		..	C		
		Hypotension		..	C		
		Stress		, , ,	C		

Date:08/31/01ISR Number: 3787780-3Report Type:Periodic Company Report #A117663
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Libido Increased	Consumer	Ziprasidone Po	PS		ORAL
ORAL				Gabapentin	SS		
600.00	MG						
TOTAL:	DAILY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/01ISR Number: 3787805-5Report Type:Periodic
Age:39 YR Gender:Male I/FU:I

Company Report #A118730

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness Dyspnoea	Consumer	Ziprasidone Po	PS	Pfizer Regulatory Safety	ORAL
80.00 MG		Hypertension					
TOTAL: BID:ORA		Hypotension					
L		Rhinitis		Neurontin	SS		ORAL
240.00 MG		Tachycardia					
TOTAL: TID:ORA							
L							

Date:08/31/01ISR Number: 3787816-XReport Type:Periodic
Age:48 YR Gender:Female I/FU:I

Company Report #A119101

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Confusional State		Ziprasidone Po	PS	Pfizer Regulatory Safety	ORAL
80.00 MG		Depersonalisation					
TOTAL: BID:ORA		Muscle Twitching					
L				Neurontin	SS		ORAL
1800.00 MG							
TOTAL: DAILY:O							
RAL				Effexor Hormone (Unspecified)	C C		

Date:08/31/01ISR Number: 3788145-0Report Type:Expedited (15-DaCompany Report #001-0945-M0101041
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Electrocardiogram T Wave Inversion	Health Professional	Neurontin (Gabapentin)	PS		

Date:08/31/01ISR Number: 3788155-3Report Type:Expedited (15-DaCompany Report #001-0945-M0101044
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 100 MG (DAILY), PER ORAL		Decreased Appetite Ear Infection Tinnitus Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:09/05/01ISR Number: 3788568-XReport Type:Expedited (15-DaCompany Report #PHRM2001FR01899
Age:86 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged TRANSDERMAL		Depression Fall Hyperkalaemia Hyponatraemia Subdural Haematoma	Foreign Health Professional Other	Nitriderm Tts (Glyceryl Trinitrate) Trans-Therapeutic-Sy stem	PS		
150 MG, BID, ORAL				Rythmol "Knoll" (Propafenone Hydrochloride) Tablet	SS		ORAL
				Coversyl			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

1 DF, QD, ORAL	(Perindopril) Tablet	SS	ORAL
2 DF, BID, ORAL	Neurontin (Gabapentin)	SS	ORAL
1 DF, QD, ORAL	Deroxat (Paroxetine Hydrochloride)	SS	ORAL
1 DF, QD, ORAL	Aldalix (Furosemide) Capsule	SS	ORAL
ORAL	Athymil (Mianserin Hydrochloride) Tablet	SS	ORAL

Date:09/05/01ISR Number: 3788762-8Report Type:Expedited (15-DaCompany Report #266750
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1 PER PRN Initial or Prolonged ORAL		Blood Albumin Decreased	Health	Naprosyn (Naproxen)	PS	Hoffmann-La Roche	ORAL
		Blood Calcium Decreased	Professional				
100 MG DAILY ORAL	11 DAY	Blood Potassium Decreased Confusional State	Other	Thalomid (Thalidomide) 50 Mg	SS	Hoffmann-La Roche	ORAL
		Coordination Abnormal					
		Dehydration		Aredia (Pamidronate Disodium)	SS		
		Mental Disorder		Neurontin			
		Movement Disorder		(Gabapentin)	SS		ORAL
300 MG 3 PER DAY ORAL		Pneumonia					
		Renal Failure Acute					
				Paxil (Paroxetine)	C		

Vitamin Nos C
 Hydrocodone
 (Hydrocodone
 Bitartrate) C
 Methadone (Methadone
 Hydrochloride) C
 Latanoprost C
 Alkeran (Melphalan) C
 Predisone C

Date:09/06/01ISR Number: 3788845-2Report Type:Expedited (15-DaCompany Report #PHEH2001US07090
 Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Albumin Decreased Blood Calcium Decreased Clumsiness Confusional State Coordination Abnormal Dehydration	Health Professional	Aredia (Disodium Pamidronate) Solution For Infusion Thalidomide (Thalidomide)	PS SS		
100 MG, QD	11	DAY		Naprosyn (Naproxen) Neurontin (Gabapentin)	SS SS		ORAL
300 MG, TID, ORAL		Dyspnoea Exacerbated Fall Hyperkalaemia Hypokalaemia Hypoxia Movement Disorder Pneumonia Renal Failure Acute		Paxil Hydrocodone (Hydrocodone) Methadone (Methadone) Latanoprost	C C C		

Freedom Of Information (FOI) Report

(Latanoprost) C
 Vitamins Nos
 (Vitamins Nos) C
 Alkeran (Melphalan) C
 Prednisone C

Date:09/06/01ISR Number: 3788982-2Report Type:Expedited (15-DaCompany Report #044-0945-M0100214
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG	Aspartate Aminotransferase	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Disability (DAILY), PER		Increased	Professional				
ORAL		Atrial Flutter					
		Blood Alkaline Phosphatase Increased		(Lansoprazole) (Amitriptyline)	C C		
		Dermatitis Atopic					

Date:09/06/01ISR Number: 3789051-8Report Type:Expedited (15-DaCompany Report #001-0945-M0101033
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG (TID)	Anaemia Arthralgia	Health Professional	Neurontin (Gabapentin)	PS		
Other		Asthenia Blood Albumin Decreased		Thalidomide (Thalidomide)	SS		ORAL
100 MG		Blood Calcium Decreased					
(DAILY), PER		Blood Potassium Decreased					
ORAL		Clumsiness Coordination Abnormal Fall		Aredia (Pamidronate Disodium) Naprosyn (Naproxen)	SS SS		ORAL
(PRN), PER		Groin Pain					
ORAL		Hypoaesthesia		Calcium Carbonate	C		

Medication Error	Lopid (Gemfibrozil)	C
Mental Impairment	Advair	C
Movement Disorder	Pyridoxine	
Neutropenia	Hydrochloride	C
Pain In Extremity	Hydrocodone	C
Pharyngolaryngeal Pain	Lipitor	
Pneumonia	(Atorvastatin)	C
Renal Failure	Omeprazole	C
Spinal Column Stenosis	Paroxetine	
Tenderness	Hydrochloride	C
White Blood Cell Count	Latanoprost	C
Decreased	Promethazine	
	Hydrochloride	C
	Methadone	C
	Vitamins	C

Date:09/06/01ISR Number: 3789092-0Report Type:Expedited (15-DaCompany Report #001-0945-M0101043
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Health	Neurontin			
Other		Abnormal Behaviour	Professional	(Gabapentin)	PS		ORAL
PER ORAL		Dysarthria		Diazepam	C		
		Facial Palsy		Zolpidem Tartrate	C		
		Staring					
		Weight Increased					

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation	Health	Ziprasidone Po	PS		ORAL
80.00 MG		Amenorrhoea	Professional				
TOTAL: BID:		Cardiac Disorder					
ORAL		Gastritis		Gabapentin	SS		ORAL
1200.00 MG		Haematemesis					
TOTAL: BID:		Hyperprolactinaemia					
ORAL		Sinus Tachycardia		Risperdal	C		
		Vomiting		Trazadone	C		
				Ativan	C		
				Cogentin	C		
				Multivitamin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/11/01ISR Number: 3790902-1Report Type:Expedited (15-DaCompany Report #PHEH2001US06513

Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Albuminuria Anaemia Asthenia	Health Professional	Aredia (Disodium Pamidronate) Solution For Infusion	PS		
INTRAVENOUS	90 MG,	QMO, Blood Urea Increased					
INTRAVENOUS		Constipation Dehydration		Thalidomide (Thalidomide)	SS		
200 MG, QHS		Diarrhoea Dizziness		Neurontin (Gabapentin)	SS		
300 MG, BID		Flushing		Arava (Leflunomide)	SS		
254 DAY		Hypocalcaemia Hypoproteinaemia Leukopenia Mucosal Inflammation Nausea Nephrotic Syndrome Oedema Oedema Peripheral Orthostatic Hypotension Pain Proteinuria Renal Failure Acute Syncope Thrombocytopenia		Carmustine (Carmustine) Cytosan (Cyclophosphamide) Decadron Vincristine (Vincristine) Adriamycin (Doxorubicin) Prevacid (Lansoprazole) Synthroid Ambien Prednisone Epogen (Epoetin Alfa) Biaxin (Clarithromycin)	C C C C C C C C C C C C C C		

Date:09/11/01ISR Number: 3791351-2Report Type:Expedited (15-DaCompany Report #033-0945M0100108

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Fall	Foreign	Neurontin			

Initial or Prolonged (BID), PER Other ORAL	Gait Spastic Hyponatraemia	Health Professional	(Gabapentin)	PS	ORAL
DAILY, PER ORAL			Nitroderm (Glyceryl Trinitrate)	SS	ORAL
(BID), PER ORAL			Rythmol (Propafenone)	SS	ORAL
(DAILY), PER ORAL			Coversyl (Perindopril)	SS	ORAL
PER ORAL			Deroxat (Paroxetine Hydrochloride)	SS	ORAL
(DAILY), PER ORAL			Aldalix (Furosemide, Spironolactone)	SS	ORAL
			Zoloft (Sertraline Hydrochloride)	C	
			Athymil (Mianserin Hydrochloride)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/11/01ISR Number: 3791561-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101060

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Alanine Aminotransferase	Consumer	Neurontin			
Initial or Prolonged	Increased		(Gabapentin)	PS		
Other	Anxiety		Dilantin (Phenytoin			
	Aspartate		Sodium)	SS		
	Aminotransferase					
	Increased					
	Blood Alkaline					
	Phosphatase Increased					
	Blood Cholesterol					
	Increased					
	Blood Triglycerides					
	Increased					
	Depression					
	Dyspnoea					
	Facial Nerve Disorder					
	Facial Pain					
	Facial Palsy					
	Lethargy					
	Liver Function Test					
	Abnormal					
	Muscle Spasms					
	Nausea					
	Pain In Jaw					
	Periodontitis					
	Splenomegaly					
	Suicidal Ideation					
	Tearfulness					
	Tooth Abscess					
	Trigeminal Neuralgia					
	Urticaria					
	Vertigo					
	Weight Increased					

Date:09/12/01ISR Number: 3791741-8Report Type:Direct

Company Report #

Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Overdose		Gabapentin (300 Mg)	PS		ORAL
300 MG PO QD						

Initial or Prolonged
X 7 D THEN

BID

Nitroglycerine S.L.	C
Tylenol #3	C
Albuterol Inhaler	C
Lopressor	C
Maalox	C
Asa Nifedipine	C
Prevacid	C

Date:09/12/01ISR Number: 3791790-XReport Type:Direct
Age:28 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dystonia		Geodon	PS		
60 MG		Oral Discomfort		Zoloft	SS		
BID		Oral Pain		Neurontin	SS		
BID							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/12/01ISR Number: 3792196-XReport Type:Expedited (15-DaCompany Report #064-0945-M0100008

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Complications Of Maternal Exposure To Therapeutic	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
PER ORAL		Drugs Foetal Disorder	Professional Company Representative				

Date:09/12/01ISR Number: 3792198-3Report Type:Expedited (15-DaCompany Report #064-0945-M0100007

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Complications Of Maternal Exposure To Therapeutic	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
PER ORAL		Drugs Foetal Disorder	Professional Company Representative				

Date:09/12/01ISR Number: 3792301-5Report Type:Expedited (15-DaCompany Report #2014149

Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Gastrointestinal Haemorrhage	Health Professional Company Representative	Oxycontin Cr Tablets, 10 Mg (Oxycodone Hydrochloride)	PS		ORAL
MG UNKNOWN PO				Neurontin (Gabapentin)	SS		

Date:09/14/01ISR Number: 3792823-7Report Type:Expedited (15-DaCompany Report #B0119027A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	500MG Twice Disability per day	Abnormal Behaviour Aggression		Lamotrigine Valproate Sodium	PS SS	Glaxo Wellcome	ORAL
Other	20MG per day	Agitation		Clobazam	SS		
800MG Three times per day		Condition Aggravated		Gabapentin	SS		
2.5UNIT per day		Depression		Hydrocortisone	SS	Glaxo Wellcome	ORAL
		Fall		Thyroxine Sodium	SS	Glaxo Wellcome	ORAL
		Insomnia					
		Motor Dysfunction					
		Obsessive-Compulsive Disorder					

Date:09/14/01ISR Number: 3793104-8Report Type:Periodic Company Report #PERCOCET2000-00428
Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aspiration	Health	Percocet	PS	Endo	ORAL
Life-Threatening		Overdose	Professional	Duragesic 25mcg/H	SS	Janssen	
TRANSDERMAL	75 MCG/H ONCE			Fentanyl Injection 0.05mg/Ml			
TRANSD				Janssen	SS	Janssen	
INTRAVENOUS	ONCE IV			Seroquel	SS		ORAL
PO							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

PO	Revia Dupont	SS	Dupont	ORAL
PO	Neurontin	SS		ORAL
PO	Trazodone	SS		ORAL
PO	Depakote	SS		ORAL
PO	Amitriptyline	SS		ORAL

Date:09/14/01ISR Number: 3793636-2Report Type:Expedited (15-DaCompany Report #001-0945-M0101061
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthralgia	Consumer	Neurontin			
Other		Asthenia		(Gabapentin)	PS		ORAL
300 MG		Condition Aggravated					
(DAILY), PER		Pain In Extremity					
ORAL							

Date:09/14/01ISR Number: 3793796-3Report Type:Expedited (15-DaCompany Report #2001071793FR
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia	Foreign Health	Ternomin(Atenolol) Tablet, 100mg	PS		ORAL
100 MG, QD,		Herpes Simplex	Professional				
ORAL	58 DAY	Pyrexia	Other	Neurontin(Gabapentin)	SS		ORAL
ORAL	58 DAY	Viral Infection		Solupred(Prednisolon e Sodium Sulfobenzoate) 30 Mg	SS		ORAL
30 MG, QD,							

ORAL				Prograf (Tacrolimus)	SS		ORAL
ORAL				Flagentyl (Secnidazole) 500mg	SS		ORAL
500 MG, QD,							
ORAL				Triflucan (Fluconazole) 400 Mg	SS		
INTRAVENOUS	400 MG, QD,						
IV							

Date: 09/14/01
 ISR Number: 3793800-2
 Report Type: Expedited (15-DaCompany Report #033-0945-M0100113)
 Age: 47 YR Gender: Female I/FU: I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 800 MG PER Other ORAL	Alanine Aminotransferase Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
PER ORAL	Aspartate Aminotransferase Increased	Professional	Paracetamol, Opium, Caffeine	SS		
40 MG PER ORAL	Blood Bilirubin Increased Conversion Disorder Hepatocellular Damage International Normalised		Fluoxetine (Fluoxetine Hydrochloride)	SS		
1200 MG PER ORAL	Ratio Increased Malaise		Ibuprofen (Ibuprofen)	SS		ORAL
150 MG PER ORAL	Multiple Sclerosis Neuralgia		(Tetrazepam)	SS		ORAL
150 MG PER	Prothrombin Time Prolonged Sjogren'S Syndrome Vomiting		Plaquenil (Hydroxychloroquine Sulfate)	SS		ORAL

Freedom Of Information (FOI) Report

ORAL

(Azathioprine) C
 (Levothyroxine Sodium) C
 (Omeprazole) C
 (Alprazolam) C
 (Anethole Trithione) C
 (Tamsulosin) C
 (Clorazepate Dipotassium) C
 (Sucrose, Casein, Starch Hydrolysed, Yeast Dried, Lactoprotein, Fat Emulsions, Vitamins (Pseudoephedrine) C

Date:09/14/01ISR Number: 3794083-XReport Type:Expedited (15-DaCompany Report #002-0945-990030
 Age:10 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Disorder	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Hospitalization - 1400 MG		Cardio-Respiratory Arrest					
Initial or Prolonged (DAILY), PER		Drug Interaction	Professional				
Other ORAL		Hypotension					
		Liver Disorder		Morphine (Morphine)	SS		
INTRAVENOUS	INTRAVENOUS	Pain		Carbamazepine (Carbamazepine)	SS		ORAL
200 MG (BID),		Respiratory Depression					
PER ORAL		Tachycardia					
		Viral Infection		Amitriptyline (Amitriptyline)	SS		ORAL
75 MG							
(NIGHTLY),							
PER ORAL							

Outcome PT
Hospitalization - Alanine Aminotransferase
Initial or Prolonged Increased
Aspartate
Aminotransferase
Increased
Blood Alkaline
Phosphatase Increased
Blood Bilirubin Increased
Blood Lactate
Dehydrogenase Increased
Cardiomegaly
Chest Pain
Culture Urine Positive
Depressed Level Of
Consciousness
Ear Pain
Fatigue
Headache
Hypervolaemia
Hypocalcaemia
Infected Skin Ulcer
Liver Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Metabolic Disorder Pharyngolaryngeal Pain Pleural Effusion	Report Source	Product	Role	Manufacturer	Route
9	DAY	Proteus Infection		Floxacillin Sodium	PS		ORAL
		Renal Failure Acute		Leflunomide	SS		ORAL
7	DAY			Vioxx	SS	Merck & Co., Inc	ORAL
				Vasotec	SS		ORAL
7	DAY			Gabapentin	SS		ORAL
				Furosemide	SS		ORAL
9	DAY			Metronidazole	SS		ORAL
				Spironolactone	SS		ORAL
				Folic Acid	C		
				Cisapride	C		
				Albuterol Sulfate And Ipratropium Bromide	C		
				Calcium Carbonate And Cholecalciferol	C		
				Loperamide	C		
				Zolpidem Tartrate	C		
				Acetaminophen	C		
				Morphine	C		
				Cetirizine Hydrochloride	C		
				Lansoprazole	C		
				Morphine	C		
				Prednisolone	C		
				Vitamin E	C		

Date:09/17/01ISR Number: 3794194-9Report Type:Expedited (15-DaCompany Report #001-0945-M0101074
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Confusional State	Consumer	Neurontin (Gabapentin)	PS		ORAL
3600 MG		Convulsion					
(DAILY), PER							

ORAL

Difficulty In Walking

Dizziness
Paraesthesia
Speech Disorder

Thalidomide
(Thalidomide) SS
Vancomycin C
Dexamethasone C
Rabeprazole Sodium C
Warfarin Sodium C

Date:09/17/01ISR Number: 3794247-5Report Type:Expedited (15-DaCompany Report #001-0945-M0101072

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL Other		Concussion Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Fall Intervertebral Disc Disorder Stress Suspiciousness		Zoloft (Sertraline Hydrochloride) Klonopin (Clonazepam) Soma (Carisoprodol) Daypro (Oxaprozin) Zyprexa (Olanzapine)	C C C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/17/01ISR Number: 3794490-5Report Type:Expedited (15-DaCompany Report #047-0945-M0000003

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Cleft Lip And Palate Complications Of Maternal	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
2400 MG		Exposure To Therapeutic	Professional				
(DAILY), PER		Drugs					
ORAL		Hypotonia		Frisium (Clobazam)	SS		ORAL
40 MG, PER				(Folic Acid)	C		
ORAL				Vitamin D (Ergocalciferol)	C		
				(Clonazepam)	C		

Date:09/17/01ISR Number: 3794561-3Report Type:Expedited (15-DaCompany Report #WAES 01027276

Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Alanine Aminotransferase Increased	Foreign Other	Tab Vasotec (Enalapril Maleate)	PS	Merck Sharp & Dohme	ORAL
20 MG/DAILY		Aspartate					
PO	7 DAY	Aminotransferase Increased		Tab Vioxx (Rofecoxib)	SS		ORAL
25 MG/DAILY		Blood Alkaline					
PO		Phosphatase Increased		Cap Gabapentin	SS		ORAL
2400 MG/DAILY		Blood Bilirubin Increased					
PO		Blood Creatinine		Tab Metronidazole	SS		ORAL
800 MG/DAILY		Increased					
PO	9 DAY						

40 MG/DAILY		Blood Lactate	Tab Furosemide	SS	
		Dehydrogenase Increased			
PO	7	DAY			
		Blood Potassium Increased	Tab Floxacillin		
		C-Reactive Protein	Sodium	SS	ORAL
2200 MG/DAILY		Increased			
PO	9	DAY			
		Cardiomegaly	Tab Leflunomide	SS	ORAL
20 MG/DAILY		Chest Pain			
PO		Condition Aggravated	Tab Spironolactone	C	ORAL
50 MG/DAILY		Depressed Level Of			
PO		Consciousness	Acetaminophen	C	
		Ear Pain	Albuterol Sulfate		
		Electroencephalogram	(+) Ipratropium		
		Abnormal	Bromid	C	
		Fatigue	Calcium Carbonate		
		Headache	(+) Cholecalciferol	C	
		Hepatic Enzyme Increased	Cetirizine		
		Hypervolaemia	Hydrochloride	C	
		Hypocalcaemia	Cisapride	C	
		Liver Disorder	Folic Acid	C	
		Liver Function Test	Lansoprazole	C	
		Abnormal	Loperamide	C	
		Metabolic Disorder	Morphine	C	
		Pain	Morphine	C	
		Pharyngolaryngeal Pain	Prednisolone	C	
		Pleural Effusion	Vitamin E	C	
		Proteus Infection	Zolpidem Tartrate	C	
		Renal Disorder			
		Renal Failure			
		Urinary Tract Infection			
		Weight Decreased			
		Weight Increased			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/17/01ISR Number: 3794820-4Report Type:Expedited (15-DaCompany Report #061-0945-M0100035

Age:82 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Oedema Peripheral	Foreign	Neurontin			
Hospitalization -		Sepsis	Consumer	(Gabapentin)	PS		ORAL
1200 MG			Health				
Initial or Prolonged			Professional				
(DAILY), PER							
Other							
ORAL				Celecoxib	SS		ORAL
200 MG							
(DAILY), PER							
ORAL				Levodopa Carbidopa	C		
				Unspecified			
				Medications	C		
				Fursemide	C		
				Potassium Chloride	C		
				Atorvastatin	C		
				Karvezide	C		
				Ranitidine			
				Hydrochloride	C		
				Clopidogrel	C		
				Doxepin			
				Hydrochloride	C		

Date:09/17/01ISR Number: 3794822-8Report Type:Expedited (15-DaCompany Report #A121061

Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Asthenia	Foreign	Diffucan Injection	PS		
INTRAVENOUS	400.00	MG	Health				
Initial or Prolonged		Dermatitis	Professional				
TOTAL:DAILY:I							
		Herpes Simplex					
INTRAVENOUS			Other	Gabapentine	SS		ORAL
ORAL		Pyrexia					

30.00 MG
 Predisolone SS ORAL
 TOTAL:DAILY:0
 RAL
 DAILY:ORAL
 Tacrolimus SS ORAL
 Secnidazole C
 Atenolol C

Date:09/18/01ISR Number: 3794983-0Report Type:Direct Company Report #
 Age:54 YR Gender:Male I/FU:I
 Outcome PT Report Source Product Role Manufacturer Route
 Dose Duration
 Other Nightmare Gabapentin PS
 300 G HS

Date:09/18/01ISR Number: 3794985-4Report Type:Direct Company Report #
 Age:56 YR Gender:Male I/FU:I
 Outcome PT Report Source Product Role Manufacturer Route
 Dose Duration
 Other Dermatitis Gabapentin PS ORAL
 600 MG BID

ORAL
 Verapamil Hcl C
 Terazosin Hcl C
 Alendronate C
 Lansoprazole C
 Latanoprost C
 Calcium C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/19/01ISR Number: 3796179-5Report Type:Expedited (15-DaCompany Report #061-0945-M0100041

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2700 MG (THREE TIMES A DAY), PER ORAL	Circulatory Collapse Condition Aggravated Dizziness	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:09/19/01ISR Number: 3796192-8Report Type:Expedited (15-DaCompany Report #044-0981-M0100739

Age:50 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Hepatitis Jaundice Cholestatic	Foreign	Atorvastatin (Atorvastatin) Gabapentin (Gabapentin)	PS SS		

Date:09/19/01ISR Number: 3796202-8Report Type:Expedited (15-DaCompany Report #055-0945-M0100011

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Hospitalization - 900 MG PER Initial or Prolonged ORAL Other	Cardiac Disorder Oedema Peripheral Poor Peripheral Circulation	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
			Unspecified Drug For Circulation Disorder	C		
			Unspecified Drug For Prostate Disorder	C		
			Unspecified Drug For A Stomach Disorder	C		
			Unspecified Drug For Hypertension	C		

Date:09/19/01ISR Number: 3796238-7Report Type:Expedited (15-DaCompany Report #055-0945-M0100024

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia	Foreign	Gabapentin			
Hospitalization - PER ORAL		Cardiac Disorder	Consumer	(Gabapentin)	PS		ORAL
Initial or Prolonged Other		Cerebrovascular Accident Dysphagia Poor Peripheral Circulation		Unspecified Drug For Circulation Disorder Unspecified Drug For Prostate Disorder Unspecified Drug For Stomach Disorder Unspecified Drug For Hypertension	C C C C		

Date:09/20/01ISR Number: 3795853-4Report Type:Direct

Company Report #

Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 900 MG TID PO Intervention to Prevent Permanent Impairment/Damage		Rash Papular		Gabapentin	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/20/01ISR Number: 3796552-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100815

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Glucose Increased	Health	Neurontin			
		Blood Potassium Decreased	Professional	(Gabapentin)	PS		ORAL
300 MG (TID),		Condition Aggravated					
PER ORAL		Decreased Activity		Celecoxib	SS		ORAL
200 MG		International Normalised					
(DAILY), PER		Ratio Decreased					
ORAL		Joint Stiffness		Levofloxacin	SS		ORAL
500 MG		Maculopathy					
(DAILY), PER		Neuropathy Peripheral					
ORAL		Oedema Peripheral		Clonazepam	C		
		Pain In Extremity		Estrogens Conjugated	C		
		Paraesthesia		Furosemide	C		
		Sedation		Digoxin	C		
		Tarsal Tunnel Syndrome		Warfarin Sodium	C		
		Vision Blurred		Cyanocobalamin	C		
		Visual Acuity Reduced					
		Vitreous Detachment					

Date:09/20/01ISR Number: 3796673-7Report Type:Expedited (15-DaCompany Report #001-0981-M0107389

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cardiac Disorder	Consumer	Atorvastatin			
Initial or Prolonged		Neuropathy Peripheral		(Atorvastatin)	PS		ORAL
40 MG							
Other							
(DAILY), PER							
ORAL				Gabapentin			
				(Gabapentin)	SS		

(Unspecified
Medications For
Heart Disease) C

Date:09/20/01ISR Number: 3796676-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100761
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2100 MG (QID),		Circulatory Collapse Headache Loss Of Consciousness Rash Erythematous Rash Pruritic Suicide Attempt	Consumer Health Professional	Neurontin (Gabapentin) Prevacid (Lansaprozole)	PS C		

Date:09/20/01ISR Number: 3796987-0Report Type:Expedited (15-DaCompany Report #001-0945-M0101044
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 100 MG (DAILY), PER ORAL		Condition Aggravated Decreased Appetite Labyrinthitis Otitis Externa Tinnitus Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/24/01ISR Number: 3797795-7Report Type:Expedited (15-DaCompany Report #001-0945-M0101088

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Convulsion Tremor	Health Professional	Neurontin (Gabapentin)	PS		

Date:09/24/01ISR Number: 3797855-0Report Type:Expedited (15-DaCompany Report #001-0945-M0101094

Age:13 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Hospitalization - 900 MG (TID) Initial or Prolonged INTRAVENOUS 368 MG (BID), Other INTRAVENOUS	Abnormal Behaviour Acute Sinusitis Atrophy Blood Culture Positive	Consumer	Neurontin (Gabapentin) Voriconazole	PS SS		
96 CC (HOURLY) 40 MG (BID)	Cardiac Arrest Convulsion Disorientation Drug Interaction Fall Fungal Abscess Central Nervous System Injury Metabolic Encephalopathy Sedation Urinary Incontinence		Cyclosporin (Ciclosporin) Oxycontin (Oxycodone Hydrochloride)	SS SS		

Date:09/24/01ISR Number: 3797954-3Report Type:Expedited (15-DaCompany Report #001-0945-M0101101

Age:29 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 4800 MG (1600	Blood Pressure Increased Confusional State	Literature Health	Gabapentin (Gabapentin)	PS		

Other
MG, TID),

Heart Rate Increased

Professional

Medication Error
Respiratory Rate
Increased

Date:09/24/01ISR Number: 3797963-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101044

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 100 MG (DAILY), PER ORAL	Condition Aggravated Decreased Appetite Ear Infection Medication Error Paranoia Psychotic Disorder Tinnitus Weight Decreased	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:09/24/01ISR Number: 3797965-8Report Type:Expedited (15-DaCompany Report #001-0945-M0101102

Age:36 YR Gender:Male I/FU:I

Outcome	PT
Other	Anxiety Bipolar Disorder Blood Pressure Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Clonic Convulsion Confusional State Disturbance In Attention	Report Source	Product	Role	Manufacturer	Route
3600 MG (1200 MG, TID)		Heart Rate Increased	Literature	Gabapentin	PS		
		Hyperhidrosis	Health				
		Medication Error Palpitations Respiratory Rate Increased Tremor	Professional				

Date:09/24/01ISR Number: 3797971-3Report Type:Expedited (15-DaCompany Report #001-0945-M0101092
Age: Gender:Male I/FU:I

Outcome Dose Other (TID), PER ORAL	Duration	PT Decreased Appetite Drug Ineffective Feeling Cold Haemoglobin Decreased Hyperhidrosis Malaise Memory Impairment Pneumonia Sedation Staphylococcal Infection Weight Decreased	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Levaquin (Levofloxacin)	SS		
				Oxycontin (Oxycodone Hydrochloride)	SS		
				Vicodin (Paracetamol, Hydrocodone Bitartrate)	SS		
				Flexeril (Cyclobenzaprine Hydrochloride)	C		
				Flexeril (Cyclobenzaprine Hydrochloride)	C		

Date:09/24/01ISR Number: 3797988-9Report Type:Expedited (15-DaCompany Report #001-0945-M0101017
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3200 MG (TID) Other		Chest Pain Hypertension	Health Professional	Neurontin (Gabapentin)	PS		
400 MG (BID)		Pain Pyrexia Red Blood Cell Sedimentation Rate Increased Tachycardia		Dilantin (Phenytoin Sodium)	SS		

Date:09/24/01ISR Number: 3798273-1Report Type:Expedited (15-DaCompany Report #061-0945-M0100042
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - PER ORAL Initial or Prolonged		Blood Bicarbonate Decreased Blood Lactic Acid Increased Renal Impairment	Foreign Health Professional	Gabapentin	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/24/01ISR Number: 3798274-3Report Type:Expedited (15-DaCompany Report #055-0945-M0100024

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiovascular Disorder	Foreign	Gabapentin			
Hospitalization - PER ORAL		Cerebrovascular Accident	Consumer	(Gabapentin)	PS		ORAL
Initial or Prolonged		Dysphagia		Unspecified Drug For Circulation Disorder	C		
Other		Gastrointestinal Disorder Metastasis		Unspecified Drug For Prostate Disorder	C		
				Unspecified Drug For Stomach Disorder	C		
				Unspecified Drug For Hypertension	C		

Date:09/24/01ISR Number: 3798275-5Report Type:Expedited (15-DaCompany Report #055-0945-M0100011

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiovascular Disorder	Foreign	Gabapentin			
Hospitalization - 900 MG, PER ORAL		Gastrointestinal Disorder	Consumer	(Gabapentin)	PS		ORAL
Initial or Prolonged		Metastasis					
Other		Oedema Peripheral Refusal Of Treatment By Patient		Unspecified Drug For Circulation Disorder	C		
				Unspecified Drug For Prostate Cancer	C		
				Unspecified Drug For Stomach Disorder	C		
				Unspecified Drug For Hypertension	C		

Date:09/24/01ISR Number: 3798284-6Report Type:Expedited (15-DaCompany Report #420-0945-M0100002

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Foreign	Gabapentin	PS		ORAL
900 MG (300							

Life-Threatening
MG, TID), PER
Hospitalization -
ORAL
Initial or Prolonged

Health
Professional
Company
Representative

Date:09/24/01ISR Number: 3801585-6Report Type:Expedited (15-DaCompany Report #420-0945-M0100002
Age:66 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Foreign	Gabapentin	PS		ORAL
900 MG (300 Life-Threatening MG, TID), PER Hospitalization - ORAL Initial or Prolonged			Health Professional Company Representative				

Date:09/25/01ISR Number: 3799985-6Report Type:Direct Company Report #
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Confusional State		Gabapentin (300mg) (Parke-Davis)	PS	Parke-Davis	ORAL
300MG 1T QD PO							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/25/01ISR Number: 3801203-7Report Type:Expedited (15-DaCompany Report #044-0945-M0100204
 Age:50 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 Other MG, TID), PER ORAL	Hepatitis Jaundice Cholestatic	Foreign Health Professional	Neurontin(Gabapentin)	PS		ORAL
			(Insulin Human, Insulin Human Injection, Isophane) (Metformin) (Amitriptyline) (Dihydrocodeine) (Ramipril)	C C C C C		

Date:09/26/01ISR Number: 3798912-5Report Type:Direct Company Report #
 Age:37 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG BID PO 5 MG BID PO 3 MON	Fear Sedation		Gabapentin (300 Mg) Haloperidol (5 Mg)	PS SS		ORAL ORAL
			Benzotropine Depakote Haldol Decanoate	C C C		

Date:09/26/01ISR Number: 3799085-5Report Type:Expedited (15-DaCompany Report #001-0945-M0101104
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 3 GM (DAILY), PER ORAL	Drug Withdrawal Syndrome Dyskinesia Intestinal Obstruction	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Muscle Rigidity

Glibenclamide

C

Date:09/26/01ISR Number: 3800301-1Report Type:Direct
Age:49 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 1800 MG TID Intervention to ORAL Prevent Permanent Impairment/Damage		Urinary Incontinence		Gabapentin	PS		ORAL

Date:09/27/01ISR Number: 3800211-XReport Type:Direct
Age:51 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dermatitis		Gabapentin P-D	PS		

Date:09/27/01ISR Number: 3800230-3Report Type:Direct
Age:51 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 1 TID PO		Asthenia		Gabapentin 300mg P-D	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/27/01ISR Number: 3800231-5Report Type:Direct
 Age:73 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Oedema Peripheral		Gabapentin 400mg P-D	PS		ORAL
1 QHS PO							

Date:09/28/01ISR Number: 3801902-7Report Type:Direct
 Age:63 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diarrhoea		Gabapentin 300 Mg Pd	PS		
2 QD HS							
Periorbital Oedema							

Date:09/28/01ISR Number: 3802363-4Report Type:Expedited (15-DaCompany Report #386-0945-M0100001
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Hospitalization - Initial or Prolonged 300 MG (100 MG,TID), PER							
Sedation							
Professional							
ORAL							
					Fluticasone Propionate	C	
					Fenoterol Hydrobromide, Ipratropium Bromide	C	
					Warfarin	C	
					Metildigoxin	C	
					Alprazolam	C	
					Ranitidine	C	
					Theophylline	C	

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		International Normalised	Foreign	Gabapentin	PS		ORAL
300 MG (THREE		Ratio Increased	Consumer				
TIMES A DAY),		Prothrombin Time					
PER ORAL		Prolonged		Phenprocoumon	SS		ORAL
3MG, PER							
ORAL				Alcohol (Ethanol)	SS		ORAL
5 GLASSES							
(DAILY), PER							
ORAL				Quinapril	C		
				Glimepiride	C		
				Ipratropium	C		
				Bromhexine	C		
				Doxycycline	C		
				Atorvastatin	C		

Age:23 YR Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Mydriasis Respiratory Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Sedation Sensory Disturbance Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
PATIENT TOOK 50 X 25 MG TABLETS			Foreign Health Professional	Topamax (25 Mg Tablet) (Topiramate)	PS		
				Tranxilium (Clorazepate Dipotassium)	SS		
				Neurontin (Gabapentin)	SS		

Date:09/28/01ISR Number: 3802672-9Report Type:Periodic Company Report #001-0073-M0100031
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (QHS),		Convulsion	Consumer	Dilantin (Phenytoin Sodium)	PS		
				Neurontin (Gabapentin)	SS		

Date:09/28/01ISR Number: 3802709-7Report Type:Periodic Company Report #001-0073-M0100002
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG DAILY Other ; 600 MG DAILY 2700 MG (900		Cerebrovascular Accident Convulsion Difficulty In Walking Drug Level Increased Movement Disorder Sedation	Consumer	Dilantin (Phenytoin Sodium)	PS		
				Neurontin (Gabapentin)	SS		

MG TID) ;

1800-2700 MG

(600-900 MG

TID)

Percodan (Acetylsalicylic Acid, Caffeine, Phenacetin, Oxycodone	SS
Flexeril (Cyclobenzaprine Hydrochloride)	C
(Atenolol)	C
(Morphine)	C
Gemfibrozil Tablets	C

Date:09/28/01ISR Number: 3802884-4Report Type:Periodic
Age:65 YR Gender:Male I/FU:I

Company Report #2000UW04635

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Bradycardia	Health	Seroquel "Zeneca"	PS	Zeneca	ORAL
25 MG BID PO		Hypothermia	Professional	Olanzapine	SS		
				Gabapentin	SS		
				Reglan	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/01/01ISR Number: 3801992-1Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300 MG BID	1 WK			Gabapentin	PS		
Required							
Intervention to				Propoxyphene	SS		
Prevent Permanent							
Impairment/Damage							

Date:10/01/01ISR Number: 3802417-2Report Type:Expedited (15-DaCompany Report #001-0945-M0101157
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
Initial or Prolonged							
800 MG (BID)			Consumer	Neurontin (Gabapentin)	PS		

Date:10/01/01ISR Number: 3802444-5Report Type:Expedited (15-DaCompany Report #001-0945-M0101034
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
Initial or Prolonged							
1200 MG			Consumer	Neurontin (Gabapentin)	PS		ORAL
Other							
(QID), PER			Professional				
ORAL							
				Maxair (Pirbuterol Acetate)	SS		
INHALANT							

Date:10/01/01ISR Number: 3802448-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100944
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Grand Mal Convulsion	Consumer Health Professional	Neurontin (Gabapentin)	PS		
1200 MG							
(THREE TIMES A DAY)							

Lorazepam	C	
Paracetamol, Oxycodone Hydrochloride	C	
Zolpidem Tartrate	C	
Fentanyl Patches	C	

Date:10/01/01ISR Number: 3802458-5Report Type:Expedited (15-DaCompany Report #001-0945-M0101118
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 OR 400 MG, DAILY, PER ORAL		Confusional State Increased Appetite Memory Impairment Pain Vision Blurred Weight Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Cyclobenzaprine Hydrochloride Dextropropoxyphene Hydrochloride	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/01/01ISR Number: 3803140-0Report Type:Expedited (15-DaCompany Report #386-0945-M0100002

Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), Other PER ORAL		Sedation	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:10/02/01ISR Number: 3803009-1Report Type:Direct

Company Report #

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG HS ORAL		Tremor		Gabapentin	PS		ORAL

Date:10/02/01ISR Number: 3813508-4Report Type:Direct

Company Report #

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 1800 MG/DAY Intervention to Prevent Permanent Impairment/Damage		Choreoathetosis		Gabapentin	PS		

Date:10/03/01ISR Number: 3804428-XReport Type:Expedited (15-DaCompany Report #001-0945-M0101150

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG , PER		Bundle Branch Block Right Cerebral Infarction	Consumer	Neurontin (Gabapentin)	PS		ORAL

Other ORAL	Convulsion					
	Drug Level Below Therapeutic			Dilantin (Phentytoin Sodium)	SS	ORAL
PER ORAL	Partial Seizures			Folic Acid Calcitonin Salmon	C C	

Date:10/03/01ISR Number: 3804430-8Report Type:Expedited (15-DaCompany Report #001-0945-M0101138
Age: Gender:Female I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG, PER Other ORAL	Condition Aggravated Joint Stiffness	Consumer	Neurontin (Gabapentin)	PS		ORAL
PER ORAL	Movement Disorder Parkinsonian Gait Partial Seizures		Dilantin (Phenytoin Sodium)	SS		ORAL
			Folic Acid Calcitonin, Salmon	C C		

Date:10/03/01ISR Number: 3804445-XReport Type:Expedited (15-DaCompany Report #001-0945-M0101144
Age: Gender:Female I/FU:I

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (TWICE Other DAILY), PER ORAL	Hypertension	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Zyrtec (Cetirizine)			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C

Date:10/03/01ISR Number: 3804447-3Report Type:Expedited (15-DaCompany Report #001-0945-M0101139

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG Other (TWICE DAILY)		Convulsion Loss Of Consciousness Speech Disorder	Consumer Company Representative	Neurontin (Gabapentin) Topamax (Topiramate)	PS C		

Date:10/03/01ISR Number: 3805987-3Report Type:Expedited (15-DaCompany Report #061-0945-M0100044

Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (DAILY), PER ORAL		Crying Mood Altered Pain	Foreign Consumer	Neurontin (Gabapentin) (Morphine) (Pindolol)	PS C C		ORAL

Date:10/03/01ISR Number: 3805988-5Report Type:Expedited (15-DaCompany Report #002-0945-M0100118

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other PER ORAL		Blood Creatine Phosphokinase Increased	Foreign Health Professional	Gabapentin (Gabapentin) Haloperidol (Haloperidol) (Unspecified Medications)	PS SS C		ORAL

Date:10/03/01ISR Number: 3806803-6Report Type:Expedited (15-DaCompany Report #064-0945-M0100008
Age:84 DY Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly 900 MG PER ORAL	Duration Abortion Induced Complications Of Maternal Exposure To Therapeutic	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
	Drugs Exomphalos Limb Reduction Defect Multiple Congenital Abnormalities	Company Representative	(Acetylsalicylic Acid)	C		

Date:10/03/01ISR Number: 3806804-8Report Type:Expedited (15-DaCompany Report #064-0945-M0100007
Age:33 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG PER Other ORAL	Duration Abortion Induced Complications Of Maternal Exposure To Therapeutic	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
	Drugs Exomphalos Limb Reduction Defect Multiple Congenital Abnormalities	Company Representative	(Acetylsalicylic Acid)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/03/01ISR Number: 3806806-1Report Type:Expedited (15-DaCompany Report #049-0945-M0100092

Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bone Disorder	Foreign	Neurontin	PS		ORAL
PER ORAL		Fibrosis Hereditary Spherocytosis Pathological Fracture Tibia Fracture	Health Professional				

Date:10/03/01ISR Number: 3807896-2Report Type:Expedited (15-DaCompany Report #001-0945-M0101122

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Ivth Nerve Paralysis	Health Professional	Neurontin (Gabapentin)	PS		
Other							

Date:10/05/01ISR Number: 3806826-7Report Type:Expedited (15-DaCompany Report #001-0945-M0101153

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Epiglottitis Faecal Incontinence	Consumer	Neurontin (Gabapentin)	PS		
Other		Pneumonia					

Date:10/05/01ISR Number: 3806923-6Report Type:Expedited (15-DaCompany Report #044-0945-M0100231

Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase Increased	Foreign Health	Neurontin(Gabapentin)	PS		ORAL
PER ORAL		Aspartate Aminotransferase Increased	Professional				

Blood Alkaline
Phosphatase Increased
Blood Bilirubin Increased
Gamma-Glutamyltransferase
Increased

Date:10/05/01ISR Number: 3807032-2Report Type:Expedited (15-DaCompany Report #001-0945-M0101163
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation Chills	Consumer	Neurontin (Gabapentin)	PS		
300 MG (ONE DOSE)		Difficulty In Walking Motor Dysfunction Renal Colic Tremor Vomiting		Imipramine Hydrochloride Alprazolam	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/05/01ISR Number: 3807721-XReport Type:Expedited (15-DaCompany Report #001-0073-M0100434

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Fatigue	Consumer	Dilantin (Phenytoin			
Initial or Prolonged	Overdose		Sodium)	PS		
Other			Neurontin			
			(Gabapentin)	SS		

Date:10/08/01ISR Number: 3806523-8Report Type:Expedited (15-DaCompany Report #EMADSS2001005733

Age:23 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Mydriasis	Foreign	Topamax (25 Mg			
Hospitalization -	Overdose	Health	Tablet) (Topiramate)	PS		
PATIENT TOOK						
Initial or Prolonged	Respiratory Disorder	Professional				
50 X 25 MG						
TABLETS	Sedation					
	Suicide Attempt		Tranxilium			
			(Clorazepate			
			Dipotassium)	SS		
			Neurontin			
			(Gabapentin)	SS		

Date:10/08/01ISR Number: 3807648-3Report Type:Expedited (15-DaCompany Report #NSADSS2001029547

Age:21 YR Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death	Completed Suicide	Literature	Haldol (Haloperidol)	PS		ORAL
ORAL		Health	Benztropine			
		Professional	(Benztropine)	SS		
			Gabapentin			
			(Gabapentin)	SS		

Date:10/09/01ISR Number: 3806082-XReport Type:Direct
Age:36 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysarthria		Neurontin (400mg)	PS		ORAL
400MG 1T PO		Tongue Disorder					
BID		Tongue Oedema		Clonidine	C		
		Visual Disturbance		Zyprexa	C		
				Protonix	C		

Date:10/10/01ISR Number: 3807498-8Report Type:Expedited (15-DaCompany Report #061-0945-M0100047
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Aggression	Foreign	Neurontin			
Other		Brain Neoplasm	Health	(Gabapentin)	PS		ORAL
PER ORAL		Drug Effect Decreased	Professional	(Carbamazepine)	C		
			Company	(Valproate Sodium)	C		
			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/10/01ISR Number: 3807875-5Report Type:Expedited (15-DaCompany Report #033-0945-M0100113

Age:47 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 800 MG PER	Alanine Aminotransferase Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other ORAL	Aspartate	Professional				
PER ORAL	Aminotransferase Increased	Other	Paracetamol, Opium, Caffeine	SS		ORAL
40 MG PER	Blood Bilirubin Increased Coagulation Factor V Level Increased		Fluoxetine (Fluoxetine Hydrochloride)	SS		ORAL
ORAL	Conversion Disorder					
1200 MG PER	Hepatic Infarction Hepatocellular Damage		Ibuprofen (Ibuprofen)	SS		ORAL
ORAL	Malaise					
150 MG PER	Neuralgia		Tetrazepam	SS		ORAL
ORAL	Overdose					
PER ORAL	Prothrombin Time Prolonged Urinary Tract Disorder		Plaquenil (Hydroxychloroquine Sulfate)	SS		ORAL
	Vomiting		Azathioprine Levothyroxine Sodium Omeprazole Alprazolam Anethole Trithione Tamsulosin Clorazepate Dipotassium Sucrose, Casein, Starch Hydrolysed, Yeast Dried, Lactoprotein, Fat Emulsions, Vitamins Pseudoephedrine	C C C C C C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation	Health	Ziprasidone Po	PS		ORAL
80.00 MG		Drooling	Professional				
TOTAL: BID:		Electrocardiogram Qt					
ORAL		Prolonged		Gabapentin	SS		ORAL
1200 MG		Gastritis					
TOTAL: BID:		Haematemesis					
ORAL		Insomnia		Risperdal	C		
		Nausea		Trazadone	C		
		Respiratory Arrest		Ativan	C		
		Vomiting		Cogentin	C		
				Multivitamin	C		
				Haldol	C		
				Tylenol	C		
				Protonix	C		
				Restoril	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/10/01ISR Number: 3809341-XReport Type:Expedited (15-DaCompany Report #001-0945-M0100849
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG		Difficulty In Walking					
(DAILY), PER		Extrapyramidal Disorder					
ORAL		Gait Disturbance Muscle Disorder					

Date:10/10/01ISR Number: 3809613-9Report Type:Expedited (15-DaCompany Report #001-0945-M0101181
Age:79 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - 600 MG BID		Blood Glucose Decreased Bone Pain	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Initial or Prolonged PER ORAL		Coma					
Other 25 MG		Ecchymosis		Vioxx (Rofecoxib)	SS		
		Fall		(Methadone)	C		
		Inflammation		Fosamax (Alendronate Sodium)	C		
		Loss Of Consciousness		Nasal Calcitonin (Calcitonin)	C		

Date:10/10/01ISR Number: 3809684-XReport Type:Expedited (15-DaCompany Report #001-0945-M0101064
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia Dizziness	Health Professional	Neurontin (Gabapentin)	PS		ORAL
100 MG, PER		Dyspnoea					
ORAL							

Heart Rate Increased

Meridia (Sibutramine Hydrochloride) SS

ORAL

10 MG

(DAILY), PER

ORAL

Lorcet (Paracetamol, Hydrocodone Bitartrate) SS

ORAL

10 MG, PER

ORAL

Date:10/10/01ISR Number: 3809685-1Report Type:Expedited (15-DaCompany Report #001-0945-M0101175

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cellulitis	Health Professional	Neurontin (Gabapentin)	PS		ORAL

2700 MG

(TID), PER

ORAL

Corgard (Nadolol) C
Lipitor (Atorvastatin) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/10/01ISR Number: 3810470-5Report Type:Expedited (15-DaCompany Report #049-0945-M0100095

Age:65 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 1800 MG Initial or Prolonged (DAILY), PER ORAL	Grand Mal Convulsion	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			(Clonazepam)	C		

Date:10/11/01ISR Number: 3807710-5Report Type:Direct

Company Report #

Age:72 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 300 MG PO BID Intervention to Prevent Permanent Impairment/Damage	Muscle Twitching Tremor		Neurontin	PS		ORAL

Date:10/12/01ISR Number: 3807779-8Report Type:Expedited (15-DaCompany Report #A0154816A

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other SUBCUTANEOUS 100MG Per day 2700MG Per day	Diarrhoea 6MG Per day YR Eye Disorder Food Poisoning Hypersensitivity Vomiting		Imitrex Imitrex Neurontin	PS SS SS	Glaxo Wellcome Glaxo Wellcome	ORAL

Date:10/12/01ISR Number: 3808531-XReport Type:Expedited (15-DaCompany Report #055-0945-M0100049

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Foreign	Gabapentin	PS		ORAL
PER ORAL			Consumer				

Date:10/12/01ISR Number: 3808532-1Report Type:Expedited (15-DaCompany Report #044-0945-M0100236

Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Drug Interaction	Foreign	Gabapentin	PS		ORAL
Other		International Normalised	Health	(Gabapentin)			
400 MG, PER		Ratio Increased	Professional				
ORAL				(Warfarin)	C		
				(Digoxin)	C		
				Furosemide			
				(Furosemide)	C		
				(Lisinopril)	C		

Date:10/12/01ISR Number: 3808856-8Report Type:Expedited (15-DaCompany Report #049-0945-M0100097

Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Arrhythmia	Foreign	Neurontin	PS		ORAL
600 MG			Health	(Gabapentin)			
(DAILY), PER			Professional				

ORAL

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Freedom Of Information (FOI) Report

Atehexal (Atenolol) C
 Lorzaar (Losartan Potassium) C
 (Acetylsalicylic Acid) C
 Bezafibrat (Bezafibrate) C
 Humalog Mix (Insulin Lispro) C
 Ossofortin (Colecalciferol, Calcium Gluconate, Calcium Phosphate) C
 Presomen (Estrogens Conjugated) C
 (Magnesium) C

Date:10/12/01ISR Number: 3808861-1Report Type:Expedited (15-DaCompany Report #044-0945-M0100235
 Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG, PER ORAL		Convulsion	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				(Cefradine) (Oxycodone Hydrochloride) (Tramadol) (Nizatidine) (Sodium Bicarbonate, Potassium Chloride, Sodium Chloride, Macrogol)	C C C C C		

Date:10/12/01ISR Number: 3808932-XReport Type:Expedited (15-DaCompany Report #386-0945-M0100002
 Age:81 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Confusional State	Foreign	Neurontin			

Initial or Prolonged 300 MG (TID), Other PER ORAL	Coordination Abnormal Difficulty In Walking Dysphagia Gait Disturbance Movement Disorder Sedation Speech Disorder	Health Professional	(Gabapentin)	PS	ORAL
			Sinvacor (Simvastatin) Berodual (Fenoterol Hydrobromide, Ipratropium Bromide) Aspirin (Acetylsalicylic Acid)	C C C	

Date:10/12/01ISR Number: 3809070-2Report Type:Expedited (15-DaCompany Report #033-0945-M0100122
Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Life-Threatening	Coma	Foreign
Hospitalization -	Epilepsy	Health
Initial or Prolonged		Professional Company

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
		Neurontin (Gabapentin)	PS		

Date:10/12/01ISR Number: 3809854-0Report Type:Expedited (15-DaCompany Report #001-0945-M0101194
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accident	Health	Neurontin			
Hospitalization - Initial or Prolonged		Calcinosis	Professional	(Gabapentin)	PS		
Other		Cerebrovascular Accident		Sinequan (Doxepin Hydrochloride)	SS		
		Coma		Roxanol (Morphine Sulfate)	SS		
		Coronary Artery Disease		Percocet			
		Drug Toxicity		(Paracetamol, Oxycodone			
		Emphysema		Hydrochloride)	SS		
		Hypertension		Theodur			
		Myocardial Infarction		(Theophylline)	SS		
		Overdose		Imdur (Isosorbide Mononitrate)	SS		
		Thermal Burn		Lasix (Furosemide)	SS		
				Skelaxin			
				(Metaxalone)	SS		
				Relafen (Nabumetone)	SS		
				Aldomet (Methyldopa)	SS		
				Prednisone			
				(Prednisone)	SS		
				Humulin	SS		
				Ativan (Lorazepam)	SS		
				(Diltiazem)	C		

Date:10/12/01ISR Number: 3809856-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101197
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Ineffective	Consumer	Neurontin			

PER ORAL	Finger Deformity	(Gabapentin)	PS	ORAL
	Tendon Rupture	(Paracetamol, Hydrocodone Bitartrate)	C	
		(Codeine Phosphate, Paracetamol)	C	

Date:10/12/01ISR Number: 3809860-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100810
 Age:12 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bipolar Disorder	Consumer	Neurontin			
		Dysphagia	Health	(Gabapentin)	PS		
900 MG (TID)							
		Mania	Professional	Paxil (Paroxetine Hydrochloride)	SS		
20 MG (DAILY)							
				Prozac (Fluoxetine Hydrochloride)	SS		
				Albuterol			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Salbutamol) C

Date:10/12/01ISR Number: 3809873-4Report Type:Expedited (15-DaCompany Report #2000COU1548
Age:59 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged PO	Gastrooesophageal Reflux Disease International Normalised Ratio Decreased Prothrombin Time Shortened Transient Ischaemic Attack Trigeminal Neuralgia	Consumer	Coumadin (Crystalline Warfarin Sodium) Neurontin (Gabapentin) Tegretol (Carbamazepine) Dilantin (Phenytoin Sodium) Ni (Baclofen) Vioxx (Rofecoxib) Lanoxin (Digoxin) Ni Verapamil Lipitor (Atorvastatin Calcium) Ni Other (S) - Unspecified	PS SS SS SS SS SS SS SS SS		ORAL

Date:10/12/01ISR Number: 3810672-8Report Type:Expedited (15-DaCompany Report #001-0945-M0101183
Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG PER Other ORAL	Amnesia Blood Pressure Increased Dry Mouth Dysarthria Dysuria Fatigue Miosis Nausea	Consumer	Neurontin (Gabapentin) Vioxx (Rofecoxib) Zanaflex (Tizanidine)	PS SS SS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Fall Gait Disturbance Hyperkalaemia	Foreign Health Professional	Aldalix (Spironolactone, Furosemide) Tablet	PS		ORAL
1 DF, QD, ORAL		Hyponatraemia	Other				
TRANSDERMAL	1, QD,			Nitriderm Tts(Glyceryl Trinitrate)	SS		
TRANSDERMAL							
150 MG, BID, ORAL				Rythmol "Knoll"(Propafenone Hydrochloride)	SS	Knoll	ORAL
1 QD				Coversyl(Perindopril)	SS		
2 BID				Neurontin(Gabapentin)	SS		
				Deroxat(Paroxetine)			

Freedom Of Information (FOI) Report

20 MG, ORAL				Hydrochloride)	SS		ORAL
				Acticarbine (Charcoal, Activated)	C		
				Dafalgan	C		

Date:10/16/01ISR Number: 3810187-7Report Type:Expedited (15-DaCompany Report #044-0945-M0100235
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Bite Coma	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
300 MG, PER ORAL		Convulsion Eye Rolling Muscle Twitching Salivary Hypersecretion	Professional	(Cefradine) (Oxycodone Hydrochloride) (Tramadol) (Nizatidine) (Sodium Bicarbonate, Potassium Chloride, Sodium Chloride, Macrogol)	C C C C C		

Date:10/16/01ISR Number: 3810201-9Report Type:Expedited (15-DaCompany Report #049-0945-M0100098
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dyspnoea Myasthenia Gravis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (BID), PER ORAL			Professional	Immunosuppressive Agents	C		

Date:10/16/01ISR Number: 3810205-6Report Type:Expedited (15-DaCompany Report #002-0945-M0100123
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fall	Foreign	Gabapentin			
1800 MG		Hypoaesthesia	Consumer	(Gabapentin)	PS		ORAL
(DAILYI), PER		Muscle Rigidity					
ORAL		Pain					
		Spinal Fracture		(Amitriptyline)	C		
		Urinary Retention		(Lorazepam)	C		
				(Fluoxetine)	C		
				Asa/Oxycodone	C		

Date:10/16/01ISR Number: 3810243-3Report Type:Expedited (15-DaCompany Report #PERI00201004166
Age:86 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Fall	Foreign	Coversyl			
Initial or Prolonged		Hyponatraemia	Health	(Perindopril)	PS		ORAL
2 DF QD PO			Professional	Aldalix	SS		ORAL
1 DF QD PO			Other	Deroxat (Paroxetine			
DAILY PO				Hydrochloride)	SS		ORAL
				Nitriderm Tts			
				(Glyceryl			
				Trinitrate)	SS		ORAL
1 DF PO							

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Freedom Of Information (FOI) Report

1 DF PO				(Propafenone)	SS		ORAL
4 DF PO				Neurontin (Gabapentin)	SS		ORAL
				Dafalgan (Paracetamol)	C		

Date:10/16/01ISR Number: 3810542-5Report Type:Expedited (15-DaCompany Report #001-0945-M0101201
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract Macular Oedema	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG, PER		Nausea					
ORAL		Photosensitivity Reaction Visual Disturbance		Thyroid Vioxx (Rofecoxib)	C C		

Date:10/16/01ISR Number: 3810550-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101189
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300		Epistaxis					
MG, BID) PER		Hyperhidrosis					
ORAL		Impetigo Skin Infection Staphylococcal Infection Stevens-Johnson Syndrome					

Date:10/16/01ISR Number: 3810551-6Report Type:Expedited (15-DaCompany Report #001-0945-M0101179
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Glucose Decreased Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 MG (300 MG, QID) PER ORAL		Hypoglycaemia Unawareness					
SUBCUTANEOUS	SUBCUTANEOUS			Insulin (Insulin)	SS		
				(Mirtazpine)	C		
				(Nicotinic Acid, Meclozine Hydrochloride)	C		
				(Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Retinol, Riboflavin, (Steroid Injections)	C C		

Date:10/16/01ISR Number: 3810951-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101094
Age:13 YR Gender:Female I/FU:F

Outcome	PT
Death	Acute Respiratory
Hospitalization - Initial or Prolonged	Distress Syndrome
Other	Agitation Aspiration

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Brain Abscess Cardiac Arrest Convulsion			
900 MG (TID)		Consumer	Neurontin (Gabapentin)	PS		
40 MG (BID)			Difficulty In Walking Disorientation			
			Drug Interaction Fall			
			Oxycontin (Oxycodone Hydrochloride)	SS		
			Fungal Infection	C		
			Hypercarnia	C		
			Lactic Acidosis			
			Limb Injury	C		
			Metabolic Encephalopathy			
			Oxygen Saturation Decreased	C		
			Pneumonia Cytomegaloviral	C		
			Urinary Incontinence	C		
			Viral Infection	C		
			Voriconazole (Ciclosporin)	C		
			(Sertraline Hydrochloride)	C		
			(Methylprednisolone Sodium Succinate)	C		
			Solutions For Parenteral Nutrition	C		
			(Immunoglobulins)	C		
			(Granulocyte Colony Stimulating Factor)	C		
			(Pentamidine)	C		
			(Aciclovir)	C		
			(Metronidazole)	C		
			(Amphotericine B, Liposome)	C		
			(Ursodeoxycholic Acid)	C		
			(Methadone)	C		
			(Azithromycin)	C		
			(Lorazepam)	C		
			(Meropenem)	C		
			(Ciprofloxacin Hydrochloride)	C		
			(Fentanyl)	C		
			(Omeprazole)	C		
			(Vitamins Nos)	C		
			(Magnesium Oxide)	C		
			(Calcium Carbonate)	C		

Date:10/18/01ISR Number: 3811376-8Report Type:Expedited (15-DaCompany Report #044-0945-M0100231
Age:64 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization - Initial or Prolonged 900 MG (300 Other MG, TID), PER ORAL	Bile Duct Obstruction Blood Alkaline Phosphatase Increased Neoplasm Pancreatic Disorder	Foreign Health Professional	Neurontin (Gabapentin)	PS	ORAL
			Clodronate Disodium Omeprazole Paracetamol Erythromycin Fluconazole Lactulose	C C C C C C	

Date:10/18/01ISR Number: 3811377-XReport Type:Expedited (15-DaCompany Report #033-0945-M0100107
Age:52 YR Gender:Female I/FU:F

Outcome PT
Life-Threatening Coma
Medication Error
Sedation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vertigo

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
900 MG (TID), PER ORAL		Foreign Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
			Diazepam	C		
			Clorazepate			
			Dipotassium Diltiazem	C		
			Hydrochloride	C		
			Efferalgan (Codeine Phosphate, Paracetamol)	C		
			Ketoprofen	C		

Date:10/18/01
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (200 MG, TID), PER ORAL		Chills Decreased Appetite Drug Ineffective Feeling Cold	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Haemoglobin Decreased		Levaquin (Levofloxacin)	SS		
		Hyperhidrosis		Oxycontin (Oxycodone Hydrochloride)	SS		
		Memory Impairment		Vicodin (Paracetamol, Hydrocodone Bitartrate)	SS		
		Nausea		Flexeril (Cyclobenzaprine Hydrochloride)	C		
		Pneumonia					
		Sedation					
		Staphylococcal Infection					
		Weight Decreased					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Angiopathy Diarrhoea Eye Disorder	Consumer	Imitrex Injection (Sumatriptan Succinate)	PS		
SUBCUTANEOUS	6 MG/	PER Meningioma					
DAY/		Migraine					
SUBCUTANEOUS		Multiple Allergies Vomiting		Imitrex Tablet (Sumatriptan Succinate)	SS		ORAL
100 MG/	PER						
DAY/	ORAL						
				Gabapentin (Formulation Unknown) (Gabapentin)	SS		
2700 MG/	PER						
DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/18/01ISR Number: 3811520-2Report Type:Expedited (15-DaCompany Report #044-0945-M0100239

Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Drug Interaction	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		International Normalised Ratio Increased	Professional				
400 MG (DAILY), PER ORAL				(Warfarin)	C		
				(Digoxin)	C		
				(Furosemide)	C		
				(Lisinopril)	C		

Date:10/18/01ISR Number: 3811789-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101206

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous Pregnancy	Consumer	Neurontin (Gabapentin)	PS		

Date:10/18/01ISR Number: 3811809-7Report Type:Expedited (15-DaCompany Report #001-0945-M0101043

Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abdominal Distension Dysarthria	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG (TID), PER ORAL		Facial Palsy					
		Staring		Diazepam	C		
		Weight Increased		Zolpidem Tartrate	C		
				Venlafazine Hydrochloride	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Amenorrhoea	Consumer	Neurontin			
Initial or Prolonged	Arthralgia		(Gabapentin)	PS		ORAL
300 MG TID						
Other	Bipolar Disorder					
PER ORAL						
	Burning Sensation		(Oxycodone			
	Condition Aggravated		Hydrochloride)	C		
	Constipation					
	Fear					
	Hypoaesthesia					
	Nervous System Disorder					
	Neuralgia					
	Oedema					
	Pyrexia					
	Shock					
	Vaginal Candidiasis					

Outcome	PT	Report Source
Hospitalization -	Fall	Foreign
Initial or Prolonged	Hyperkalaemia	Health
	Hyponatraemia	Professional

Freedom Of Information (FOI) Report

Other

Dose	Duration	Product	Role	Manufacturer	Route
TRANSDERMAL	TRANSDERMAL	Nitriderm Tts(Glyceryl Trinitrate) Trans-Therapeutic-Sy stem	PS		
150 MG, BID, ORAL		Rythmol "Knoll"(Propafenone Hydrochloride)Tablet	SS		ORAL
1 DF, QD, ORAL		Conversyl(Perindopri l)Tablet	SS		ORAL
2 DF, BID, ORAL		Neurontin(Gabapentin)	SS		ORAL
ORAL		Deroxat(Paroxetine Hydrochloride)	SS		ORAL
1 DF, QD, ORAL		Aldalix(Furosemide)C apsule	SS		ORAL
ORAL		Athymil(Mianserin Hydrochloride)Tablet	SS		ORAL
ORAL		Dafalgan(Paracetamol) Acticarbine(Charcoal Activated) Tablet	SS SS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hallucination, Auditory	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
800 MG (400 MG, BID), PER							
ORAL							

Heparin-Fraction, Sodium Salt)	C
(Omeprazole)	C
(Dexamethasone)	C
(Morphine Sulfate)	C
(Amitriptyline)	C
(Docusate Sodium)	C
(Nortriptyline)	C
(Metoclopramide Hydrochloride)	C
(Metronidazole)	C
(Chlorhexidine Gluconate)	C

Date:10/23/01ISR Number: 3814245-2Report Type:Expedited (15-DaCompany Report #049-0945-M0100101
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Glucose Increased	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG							
Other (DAILY), PER							

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Freedom Of Information (FOI) Report

ORAL

Tremarit (Metixene
Hydrochloride) C
Propa Retard C

Date:10/24/01ISR Number: 3813468-6Report Type:Direct
Age:58 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 300 MG PO TID Required Intervention to Prevent Permanent Impairment/Damage		Cardiac Output Decreased		Neurontin	PS		ORAL
		Ecchymosis Fluid Retention International Normalised Ratio Increased Liver Disorder Oedema					

Date:10/24/01ISR Number: 3814585-7Report Type:Expedited (15-DaCompany Report #057-0945-M0100003
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 300 MG		Respiratory Arrest	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL

(DAILY), PER

ORAL

Date:10/24/01ISR Number: 3815095-3Report Type:Expedited (15-DaCompany Report #001-0945-M0101224
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dialysis Renal Failure Chronic	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/24/01ISR Number: 3815280-0Report Type:Expedited (15-DaCompany Report #055-0945-M0100051
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Diverticulitis	Foreign	Gabapentin			
Hospitalization - 800 MG Initial or Prolonged (DAILY), PER ORAL		Nosocomial Infection	Consumer	(Gabapentin)	PS		ORAL
				(Fluoxetine)	C		
				Isosorbide-5-Mono-Ni trate)	C		

Date:10/24/01ISR Number: 3815281-2Report Type:Expedited (15-DaCompany Report #044-0945-M0100250
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (THREE TIMES DAILY), PER ORAL		Renal Impairment	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				..	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/24/01ISR Number: 3815372-6Report Type:Expedited (15-DaCompany Report #001-0945-M0101237

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged (DAILY) PER	Arterial Occlusive Disease	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL	Cardiovascular Disorder					
	Condition Aggravated		(Insulin)	C		
	Drug Effect Decreased		(Multiple Vitamins)	C		
	Feeling Abnormal					
	Skin Ulcer					
	Vein Disorder					

Date:10/24/01ISR Number: 3815655-XReport Type:Expedited (15-DaCompany Report #033-0945-M0100125

Age:87 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID)	Balance Disorder Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		
Other	Fall Gait Disturbance Hepatocellular Damage	Professional	Deroxat (Paroxetine Hydrochloride)	SS		
			Augmentin (Clavulanate Potassium, Amoxicillin Trihydrate)	SS		
			Cordarone (Amiodarone Hydrochloride)	C		
			Equanil (Meprobamate0	C		
			Tareg (Valsartan0	C		
			Aspegic (Acetylsalicylate			
			Lysine)	C		

Date:10/31/01ISR Number: 3818572-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101255

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		
600 MG (BID)		Hepatic Neoplasm					
		Malignant Pain		Oxycontin (Oxycodone Hydrochloride)	SS		
20 MG (BID)		Spinal Compression Fracture		(Prednisone)	C		
		Tremor		(Levothyroxine Sodium)	C		
				(Paroxetine Hydrochloride)	C		

Date:10/31/01ISR Number: 3818649-3Report Type:Expedited (15-DaCompany Report #001-0945-M0101251
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chronic Obstructive Pulmonary Disease	Health Professional	Neurontin (Gabapentin)	PS		

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Date:10/31/01ISR Number: 3818939-4Report Type:Expedited (15-DaCompany Report #NSADSS2001029506

Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bradycardia	Literature	Tylox (Capsule)			
		Brain Herniation	Health	(Oxycodone/Acetamino			
ORAL		Brain Hypoxia	Professional	phen)	PS		ORAL
		Brain Oedema		Doxepin (Doxepin)	SS		ORAL
ORAL		Cardiac Arrest		Amitriptyline			
		Cardio-Respiratory Arrest		(Amitriptyline)	SS		ORAL
ORAL		Completed Suicide		Roxicet (Oxycocet)	SS		ORAL
		Electrocardiogram		Morphine Sulfate			
ORAL		Abnormal		(Morphine Sulfate)	SS		ORAL
		Electrocardiogram Qrs		Neurontin			
ORAL		Complex Prolonged		(Gabapentin)	SS		ORAL
				Zoloft (Sertraline			
ORAL				Hydrochloride)	SS		ORAL
				Ketoprofen			
ORAL				(Ketoprofen)	SS		ORAL

Date:10/31/01ISR Number: 3818971-0Report Type:Expedited (15-DaCompany Report #2000COU1548

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Gastrooesophageal Reflux	Consumer	Coumadin			
Initial or Prolonged		Disease		(Crystalline			
PO		International Normalised		Warfarin Sodium)	PS		ORAL
		Ratio Decreased		Neurontin			
		Medication Error		(Gabapentin)	SS		
		Prothrombin Time		Tegretol			
		Shortened		(Carbamazepine)	SS		
		Transient Ischaemic		Ni (Baclofen)	SS		
		Attack		Vioxx (Rofecoxib)	SS		
		Trigeminal Neuralgia		Lanoxin (Digoxin)	SS		

Ni (Verapamil)	SS
Lipitor(Atorvastatin Sodium)	SS
Ni (Other (S) - Unspecified)	SS
Ni(Baclofen)	SS
Dilantin(Phenytoin Sodium)	SS

Date:10/31/01ISR Number: 3818984-9Report Type:Expedited (15-DaCompany Report #001-0945-M0101232

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Thrombotic Thrombocytopenic Purpura	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/31/01ISR Number: 3819000-5Report Type:Expedited (15-DaCompany Report #NSADSS2001029547

Age:21 YR Gender:Male I/FU:F

Outcome	PT
Death	Aspiration
Required	Completed Suicide
Intervention to	Dialysis
Prevent Permanent	Heart Rate Increased
Impairment/Damage	Pneumonia

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1200.00 MG	Gastritis	Gabapentin	SS	ORAL
TOTAL: BID: ORA	Haematemesis			
L	Nausea			
	Vomiting	Risperdal	C	
		Trazadone	C	
		Ativan	C	
		Congentin	C	
		Multivitamin	C	
		Haldol	C	
		Tylenol	C	
		Protonix	C	
		Restoril	C	

Date: 10/31/01
 ISR Number: 3819231-4
 Report Type: Expedited (15-DaCompany Report #057-0945-M0100004)
 Age: Gender: Female I/FU: I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Renal Cell Carcinoma Stage Unspecified	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG							
(DAILY), PER							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/31/01ISR Number: 3819750-0Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #2001068079US

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Arthralgia	Consumer	Celebrex (Celecoxib)			
		Drug Ineffective		Capsule	PS		
		Drug Interaction		Neurontin			
		Drug Tolerance		(Gabapentin)	SS		
1200 MG, QD		Weight Increased					

Date:10/31/01ISR Number: 3820271-XReport Type:Periodic
Age:30 YR Gender:Female I/FU:I

Company Report #2001073793US

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Complications Of Maternal	Consumer	Celebrex	PS		ORAL
200 MG, QD,		Exposure To Therapeutic					
ORAL		Drugs		Neurontin(Gabapentin)	SS		ORAL
750 MG, BID,				Elavil			
ORAL				(Amitriptyline Hydrochloride)	SS		
				Zanaflex (Tizanidine Hydrochloride)	SS		
				Oral Contraceptive Nos (Oral Contraceptive Nos)	SS		

Date:10/31/01ISR Number: 3821077-8Report Type:Periodic
Age:56 YR Gender:Female I/FU:F

Company Report #2001055409US

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Arterial Disorder	Consumer	Celebrex (Celecoxib)			
		Difficulty In Walking	Health	Capsule	PS		ORAL
200MG BID							

ORAL
 Pain In Extremity Professional
 Neurontin (Gabapentin) SS
 UNKNOWN 900MG TID
 Skelaxin C
 Unspecified Hormone C
 Cipro C

Date:10/31/01ISR Number: 3821114-0Report Type:Periodic Company Report #2001065688US
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Health Professional	Celebrex (Celecoxib) Capsule	PS		ORAL
ORAL				Neurontin (Gabapentin)	SS		ORAL

Date:11/02/01ISR Number: 3819696-8Report Type:Expedited (15-DaCompany Report #2001AP05201
 Age:36 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Literature Health Professional	Quetiapine Gabapentin Indinavir	PS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/05/01ISR Number: 3820436-7Report Type:Expedited (15-DaCompany Report #001-0945-M0101270

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Carcinoembryonic Antigen Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
PER ORAL							

Date:11/05/01ISR Number: 3820462-8Report Type:Expedited (15-DaCompany Report #001-0782-M0100200

Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angina Pectoris Burning Sensation Cerebrovascular Accident	Consumer	Nitrostat Tablets (Glyceryl Trinitrate)	PS		ORAL
(PRN), PER							
ORAL							
		Infection					
		Neuralgia		Neurontin (Gabapentin)	SS		
				(Diltiazem Hydrochloride)	C		
				(Acetylsalicylic Acid)	C		
				(Pravastatin Sodium)	C		
				(Isosorbide Mononitrate)	C		
				(Metoprolol)	C		
				(Clopidogrel)	C		

Date:11/05/01ISR Number: 3820494-XReport Type:Expedited (15-DaCompany Report #A123280

Age:71 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cerebrovascular Accident	Health	Sinequan	PS		
Hospitalization -		Chest Pain	Professional	Gabapentin	SS		
Initial or Prolonged		Coma		Roxanol	SS		
Required		Coronary Artery Disease		Ativan	SS		
Intervention to		Drug Toxicity		Humulin	C		
Prevent Permanent		Emphysema		Theodur	C		

Impairment/Damage	Hypertension	Prednisone	C
	Overdose	Imdur	C
	Pain	Aldomet	C
	Thermal Burn	Lasix	C
		Relafen	C
		Percocet	C
		Skelaxin	C

Date:11/05/01ISR Number: 3820549-XReport Type:Expedited (15-DaCompany Report #001-0945-N0101268
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrioventricular Block	Consumer	Neurontin			
1200 MG		Drug Ineffective		(Gabapentin)	PS		ORAL
(QID), PER		Tremor					
ORAL							
				Lipitor			
10 MG				(Atorvastatin)	SS		ORAL
(DAILY), PER							
ORAL							
				Glyceryl Trinitrate	C		
				Furosemide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Enalapril Maleate C
 Digoxin C
 Amlodipine Besilate C
 Valproate Semisodium C
 Theophylline C
 Salbutamol C
 Sertraline
 Hydrochloride C
 Amitriptyline C

Date:11/05/01ISR Number: 3820782-7Report Type:Expedited (15-DaCompany Report #02558
 Age:62 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature	Acetaminophen/Codeine	PS		
				Lisinopril	SS		
				Gabapentin	SS		

Date:11/05/01ISR Number: 3821687-8Report Type:Expedited (15-DaCompany Report #2001077799US
 Age:38 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Ambien (Zolpidem Tartrate) Tablet	PS		
				Gabapentin (Gabapentin)	SS		

Date:11/05/01ISR Number: 3821863-4Report Type:Expedited (15-DaCompany Report #A125321
 Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	600.00 MG	Back Pain Drug Dependence	Foreign Health Professional	Sinequan Capsules Neurontin	PS SS		ORAL
TOTAL:ORAL		Drug Toxicity Movement Disorder	Professional Other	Heroin	SS		

Paraparesis
Rhabdomyolysis
Sedation

Cocaine SS
Benzodiazepines C
Alcohol C

Date:11/05/01ISR Number: 3822118-4Report Type:Expedited (15-DaCompany Report #353-0945-M0100009
Age:28 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged UNKNOWN	Dyspnoea Heart Rate Increased	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Tricyclic Antidepressants (Unspecified) Zanaflex (Tizanidine)	PS C C		ORAL

(UNKNOWN) PER
ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/05/01ISR Number: 3822119-6Report Type:Expedited (15-DaCompany Report #044-0945-M0100250

Age:79 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG (100 Other MG, THREE TIMES DAILY), PER ORAL	Renal Impairment	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
1500 MG (500 MG, THREE TIMES DAILY, PER ORAL)			Fusidic Acid (Fusidic Acid)	SS		ORAL
			Etodolac	C		
			Allopurinol	C		
			Bumetanide	C		
			Lansoprazole	C		
			Clopidogrel	C		
			Trimethoprim	C		
			Digoxin	C		

Date:11/07/01ISR Number: 3822029-4Report Type:Expedited (15-DaCompany Report #033-0945-M0100127

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (TID)	C-Reactive Protein Increased	Foreign Health Professional	Neurontin (Gabapentin)	PS		
	Fall Pneumonia Pyrexia Radius Fracture Spinal Compression Fracture		Tegretol (Carbamazepine)	C		

Vertigo

Date:11/07/01ISR Number: 3822036-1Report Type:Expedited (15-DaCompany Report #A125321
 Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Asthenia	Foreign	Sinequan Capsules	PS		
Initial or Prolonged		Back Pain	Health	Neurontin	SS		ORAL
600.00 MG							
TOTAL :ORAL		Hepatitis C	Professional				
		Neuropathy Peripheral	Other	Heroin	SS		
		Nuclear Magnetic		Cocaine	SS		
		Resonance Imaging		Benzodiazepines	C		
		Abnormal		Alcohol	C		
		Paralysis					
		Paraparesis					
		Rhabdomyolysis					
		Sedation					
		Speech Disorder					

Date:11/07/01ISR Number: 3822385-7Report Type:Expedited (15-DaCompany Report #001-0945-M0101265
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion	Consumer	Neurontin			
Initial or Prolonged		Vomiting		(Gabapentin)	PS		ORAL
8000 MG (800							

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Freedom Of Information (FOI) Report

MG , TEN

TIMES DAILY)

PER ORAL

Date:11/07/01ISR Number: 3822387-0Report Type:Expedited (15-DaCompany Report #001-0945-M0101216
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1000 MG (TID)		Face Oedema	Health Professional	Neurontin (Gabapentin)	PS		ORAL

PER ORAL

(Trazodone)	C
(Citalopram Hydrobromide)	C

Date:11/07/01ISR Number: 3822406-1Report Type:Expedited (15-DaCompany Report #001-0945-M0101179
 Age:40 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1200 MG (300 MG QID) PER		Blood Glucose Decreased Convulsion Hypoglycaemia	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

ORAL

SUBCUTANEOUS SUBCUTANEOUS

Insulin (Insulin)	SS
(Mirtazapine)	C
(Nicotinic Acid, Meclozine Hydrochloride)	C
(Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Retinol, Riboflavin,	C

Date:11/07/01ISR Number: 3823500-1Report Type:Expedited (15-DaCompany Report #002-0945-M0100129
Age:49 YR Gender:Female I/FU:U

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 200 MG (BID), Initial or Prolonged PER ORAL	Fluid Retention Oedema	Foreign Health	Gabapentin	PS		ORAL
Other	Weight Increased	Professional	(Ipratropium)	C		
			(Salbutamol Sulfate)	C		
			(Fluticasone)	C		
			(Prednisone)	C		
			(Diltiazem)	C		
			(Furosemide)	C		
			(Spironolactone)	C		
			(Morphine Sulfate)	C		
			(Paroxetine)	C		
			(Levomepromazine)	C		
			(Calcium)	C		
			(Etidronate			
			Disodium)	C		
			(Ergocalciferol)	C		
			(Estrogens			
			Conjugated)	C		
			(Domperidone)	C		
			(Omeprazole)	C		

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Freedom Of Information (FOI) Report

(Insulin Human Injection, Isophane) C
 (Insulin Human) C
 (Levothyroxine) C
 (Paracetamol) C
 (Lorazepam) C
 (Clotrimazole) C
 (Latanoprost) C

Date:11/08/01ISR Number: 3822394-8Report Type:Direct
 Age:56 YR Gender:Female I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300MG PO TID				Neurontin 300mg	PS		ORAL
		Fatigue		Asa	C		
		Neck Pain		Trazodone	C		
		Rash Erythematous		Vanco	C		
		Rash Pruritic		Lopressor	C		
		Skin Exfoliation		Lisinopril	C		

Date:11/08/01ISR Number: 3822888-5Report Type:Expedited (15-DaCompany Report #200120976EU
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
Initial or Prolonged							
20 MG / DAY				Leflunomide (Arava)			
			Foreign	Tablets	PS		ORAL
			Other				
		Cardiac Failure					
		Chest Pain		Metronidazole			
		Depressed Level Of		(Flagyl)	SS		
		Consciousness		Metronidazole			
		Hepatic Enzyme Increased		(Flagyl)	SS		ORAL
		Renal Failure Acute					
800 MG / DAY				Flucloxacillin			
				Sodium (Heracillin)	SS		
				Flucloxacillin			
PO	15	DAY					

2.2 G/DAY	15	DAY	Sodium (Heracillin)	SS	
25 MG/ DAY			Rofecoxib (Vioxx)	SS	
20MG/DAY PO	7	DAY	Enalapril Maleate (Renitec)	SS	ORAL
50 MG/DAY PO			Spiroinolactone	SS	ORAL
40 MG/DAY PO	7	DAY	Furosemide (Furix)	SS	ORAL
QD PO			Gabapentin (Neurontin)	SS	ORAL
			Folic Acid, Calcium Phosphate (Folacin)	C	
			Vitamin E	C	
			Morfin	C	
			Prednisolon	C	
			Cisapride (Prepulsid)	C	
			Cetirizine (Zyrlex)	C	
			Morphine Sulfate (Dolcontin)	C	
			Lansoprazole (Lanzo)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/09/01ISR Number: 3822620-5Report Type:Expedited (15-DaCompany Report #301216

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2 DAY			Valium	PS	Roche	
	2 DAY			Orfidal	SS		
	2 DAY			Dolantina	SS		
	2 DAY			Neurontin	SS		

Date:11/12/01ISR Number: 3823586-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101288

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	2400 MG		Consumer	Neurontin (Gabapentin)	PS		ORAL
	(DAILY), PER						
	ORAL						
				(Sertraline Hydrochloride)	C		
				(Tetracycline)	C		
				(Clarithromycin)	C		
				(Hydroxychloroquine)	C		

Date:11/12/01ISR Number: 3824360-5Report Type:Expedited (15-DaCompany Report #049-0945-M0100064

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	600 MG, PER		Foreign	Neurontin	PS		ORAL
Initial or Prolonged	ORAL		Health				
			Professional	Doxepin (Doxepin)	SS		
				Heroin (Diamorphine)	SS		
				Cocaine (Cocaine)	SS		

Muscle Twitching	Benzodiazepine	SS
Pain	Alcohol (Ethanol)	SS
Paralysis	Methadone	
Paraparesis	(Methadone)	C
Paresis	Diazepam (Diazepam)	C
Rhabdomyolysis	Aponal (Doxepin	
Sedation	Hydrochloride)	C
Speech Disorder		

Date:11/12/01ISR Number: 3824666-XReport Type:Expedited (15-DaCompany Report #2001078865SE

Age:47 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Blood Alkaline Phosphatase Increased Depressed Level Of Consciousness	Foreign Other	Spirolakton Nm Pharma (Spironolactone) Tablet	PS		
50 MG/DAY						
	Gamma-Glutamyltransferase Increased		Gabapentin (Gabapentin)	SS		
2.4 MG/DAY						
(CAPS)	Hepatic Enzyme Increased					
	Infected Skin Ulcer Metabolic Disorder		Furosemide (Furosemide)	SS		
40 MG/DAY						
	Renal Failure Acute Sedation		Enalapril (Enalapril)	SS		
20 MG/DAY						
			Rofecoxib (Rofecoxib)	SS		
25 MG/DAY						
			Metronidazole			

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Freedom Of Information (FOI) Report

800 MG/DAY	(Metronidazole)	SS
2.2 G/DAY	Flucloxacillin (Flucloxacillin)	SS
20 MG/DAY	Leflunomide (Leflunomide)	SS
	Folic Acid	C
	Tocopherol (Tocopherol)	C
	Paracetamol	C
	Prednisolon	C
	Morphine	C
	Cisapride (Cisapride)	C
	Cetirizin (Cefatrizine	
	Propyleglycolate Sulfate)	C
	Dolcontin (Morphine Sulfate)	C
	Zolpidem (Zolpidem)	C
	Calcium	C
	Loperamide (Loperamide)	C
	Ipratropium Bromide	C
	Lansoprazole	C

Date:11/12/01ISR Number: 3824678-6Report Type:Expedited (15-DaCompany Report #01P-167-0112302-00
Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Dystonia	Foreign Health Professional Other	Epilim Tablets(Sodium Valproate) (Sodium Valproate) (Sodium Valproate)	PS		ORAL
400 MG, 1 IN							
1 D, PER ORAL							
300 MG, 1 IN				Gabapentin	SS		ORAL
1 D, PER ORAL							

Bisoprolol C
Aporex C
Acetylsalicylic Acid C

Date:11/13/01ISR Number: 3825444-8Report Type:Expedited (15-DaCompany Report #044-0945-M0100192
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Liver Function Test	Foreign	Gabapentin	PS		ORAL
900 MG (TID),		Abnormal	Health				
PER ORAL			Professional	Unspecified Oncology Treatment	C		

Date:11/13/01ISR Number: 3825447-3Report Type:Expedited (15-DaCompany Report #044-0945-M0100254
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Malaise	Foreign	Neurontin			
Initial or Prolonged		Paraesthesia	Health	(Gabapentin)	PS		ORAL
50 MG							
Other		Psychotic Disorder	Professional				
(DAILY), PER		Urinary Tract Infection					
ORAL							

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FDA - Adverse Event Reporting System (AERS)

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Date:11/13/01ISR Number: 3825576-4Report Type:Expedited (15-DaCompany Report #301216

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Respiratory Failure	Foreign Other	Valium (Diazepam) 5 Mg	PS		ORAL
5 MG ORAL				Orfidal (Lorazepam) 1 Mg	SS		ORAL
2 MG ORAL				Dolantian (Meperidine Hydrochloride) 100 Mg/2ml	SS		ORAL
100 MG ORAL				Neurontin (Gabapentin) 300 Mg	SS		ORAL
900 MG ORAL							

Date:11/13/01ISR Number: 3826125-7Report Type:Expedited (15-DaCompany Report #039-0945-M0100010

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Coordination Abnormal Dyspnoea	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
300 MG (ONCE)		Fatigue	Professional				
PER ORAL							

Date:11/14/01ISR Number: 3825078-5Report Type:Expedited (15-DaCompany Report #001-0945-M0101295

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Epistaxis	Health Professional	Neurontin (Gabapentin)	PS		
				Tylenol Arthritis Extended Relief (Paracetamol)	SS		
				Tylenol Extra Strength			

1000 MG,

(Paracetamol)

SS

ORAL

DAILY, PER

ORAL

Tocopherol	C
Selenium	C
Allopurinol	C
Carteolol	
Hydrochloride	C
Ascorbic Acid,	
Tocopheryl Acetate,	
Retinol, Zinc,	
Calcium, Vitamins	
Nos, Minerals Nos,	C

Date:11/14/01ISR Number: 3825131-6Report Type:Expedited (15-DaCompany Report #001-0945-M0100731

Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Neurontin			
		Embolism	Health	(Gabapentin)	PS		ORAL
UNKNOWN		Fatigue	Professional				
(TID), PER		Oedema Peripheral					
ORAL		Sedation		Atenolol (Atenolol)	C		
		Weight Increased		Maxzide			
				(Hydrochlorothiazide			

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Freedom Of Information (FOI) Report

, Triamterene) C

Date:11/14/01ISR Number: 3825325-XReport Type:Expedited (15-DaCompany Report #044-0945-M0100236
Age:81 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Drug Interaction	Foreign	Neurontin			
Other	International Normalised	Health	(Gabapentin)	PS		ORAL
400 MG, PER	Ratio Increased	Professional				
ORAL			Unspecified			
			Medication	SS		
			(Warfarin)	C		
			(Digoxin)	C		
			Furosemide			
			(Furosemide)	C		
			(Lisinopril)	C		

Date:11/14/01ISR Number: 3825343-1Report Type:Expedited (15-DaCompany Report #001-0945-M0101000
Age:77 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Arthralgia	Health	Neurontin			
Initial or Prolonged	Arthritis	Professional	(Gabapentin)	PS		
Other	Asthenia		Depakote (Valproate			
	Cardiac Murmur		Semisodium)	SS		
	Convulsion		(Hydrochlorothiazide			
	Drug Eruption		, Losartan			
	Limb Injury		Potassium)	C		
	Mental Impairment		(Warfarin Sodium)	C		
	Myalgia		(Metoprolol)	C		
	Oedema Peripheral		(Enalapril Maleate)	C		
	Pyrexia					
	Rales					
	Tremor					

Date:11/14/01ISR Number: 3825435-7Report Type:Expedited (15-DaCompany Report #001-0945-M0101024
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia Drooling	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 MG (BID), PER		Electrocardiogram Qt Prolonged					
ORAL 80 MG (40 MG, BID), PER		Respiratory Arrest Sudden Death		Geodon	SS		ORAL
ORAL				Risperidone	C		
				Trazodone	C		
				Temazepam	C		
				Pantoprazole	C		
				Paracetamol	C		
				Lorazepam	C		
				Haloperidol	C		
				Benzatropine Mesilate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/14/01ISR Number: 3826403-1Report Type:Expedited (15-DaCompany Report #045-0945-M0100014

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), PER ORAL		Syncope	Foreign Health Professional Company Representative	Gabapentin (Gabapentin)	PS		ORAL

Date:11/14/01ISR Number: 3826509-7Report Type:Expedited (15-DaCompany Report #044-0945-M0100088

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Hypoglycaemia	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:11/14/01ISR Number: 3826510-3Report Type:Expedited (15-DaCompany Report #044-0945-M0100090

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cardiac Failure Lower Respiratory Tract Infection Transient Ischaemic Attack	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:11/14/01ISR Number: 3826869-7Report Type:Expedited (15-DaCompany Report #045-0945-M0100010

Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 300 MG		Emphysema Sudden Death	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL

Initial or Prolonged
(DAILY), PER
Other
ORAL

Professional

(Theophylline) C
(Prednisolone) C
(Budesonide) C
(Salmeterol) C
(Acetylcysteine) C
(Ipratropium) C
(Montelukast) C
(Salbutamol) C
(Alprazolam) C
(Sertraline) C
(Mianserin) C

Date:11/14/01ISR Number: 3826872-7Report Type:Expedited (15-DaCompany Report #057-0945-M0100005
Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Infection	Foreign	Neurontin			
Other		Lung Disorder	Consumer	(Gabapentin)	PS		ORAL
300 MG							

(DAILY), PER

ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/16/01ISR Number: 3825376-5Report Type:Direct
Age:43 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neutrophil Count		Neurontin 300mg	PS		
1000MG/DAY IN		Decreased					
7 DIVIDED		White Blood Cell Count					
DOSES		Decreased		Neurontin 100mg	SS		
				Amoxicillin	C		
				Depakene	C		

Date:11/16/01ISR Number: 3826923-XReport Type:Expedited (15-DaCompany Report #033-0945-M0100141
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Agitation	Foreign	Neurontin			
Initial or Prolonged		Condition Aggravated	Health	(Gabapentin)	PS		
		Delusion	Professional	Neuroleptics	C		
		Schizophrenia					

Date:11/16/01ISR Number: 3826942-3Report Type:Expedited (15-DaCompany Report #033-0945-M0100135
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Level Above Therapeutic	Foreign Literature	Gabapentin (Gabapentin)	PS		
400 MG		Neurotoxicity	Health				
(DAILY)		Petit Mal Epilepsy	Professional	(Diazepam)	C		
				(Clonazepam)	C		
				(Lamotrigine)	C		

Date:11/16/01ISR Number: 3826953-8Report Type:Expedited (15-DaCompany Report #061-0945-M0100064
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2700 (900 MG , TID), PER		Cognitive Disorder Confusional State	Foreign Health Professional	Neurontin(Gabapentin)	PS		ORAL
ORAL				(Sertraline Hydrochloride)	C		
				(Methadone)	C		
				(Gliclazide)	C		
				(Metronidazole)	C		

Date:11/19/01ISR Number: 3825858-6Report Type:Expedited (15-DaCompany Report #WAES 01112380
Age:35 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Unevaluable Event		Crixivan Gabapentin Quetiapine Fumarate [Composition Unspecified]	PS SS SS SS	Merck & Co., Inc	ORAL ORAL ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/19/01ISR Number: 3825859-8Report Type:Expedited (15-DaCompany Report #WAES 01111661

Age:21 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Cogentin	PS	Merck & Co., Inc	ORAL
				Haloperidol	SS		
				Gabapentin	SS		

Date:11/19/01ISR Number: 3827583-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101301

Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abdominal Pain	Literature	Gabapentin			
Hospitalization -		Aggression	Health	(Gabapentin)	PS		
Initial or Prolonged		Agitation	Professional	Haloperidol			
Other		Aspartate		(Haloperidol)	SS		
15 MG (5 MG,		Aminotransferase					
TID)		Increased		Olanzapine			
		Blood Albumin Increased		(Olanzapine)	SS		
		Blood Calcium Decreased		Nortriptyline			
		Blood Creatine		(Nortriptyline)	SS		
		Phosphokinase Increased		Zolpedem (Zolpidem)	SS		
		Blood Pressure Increased		Fluoxetine	SS		
		Blood Sodium Increased		Multivitamins			
		Cardiac Arrest		(Ergocalciferol,			
		Chest X-Ray Abnormal		Ascorbic Acid, Folic			
		Constipation		Acid, Thiamine			
		Delirium		Hydrochloride,	SS		
		Dry Skin		Unspecified			
		Dysuria		Laxatives	SS		
		Excoriation		Clomipramine			
		Haematuria		(Clomipramine)	SS		
25 MG (DAILY)		Haemodialysis		Benztropine			
		Headache		(Benzatropine			
		Heart Rate Increased		Mesilate)	SS		
4 MG (BID)		Insomnia		Buspirone Sr			
		Lethargy		(Buspirone)	SS		
100 MG							

(DAILY)

Metabolic Acidosis

Multi-Organ Failure
Muscle Rigidity

Clonazepam
(Clonazepam)

SS

1.5 MG (TID)

Neuroleptic Malignant
Syndrome
Proteinuria
Prothrombin Time
Prolonged
Renal Failure
Respiratory Rate
Increased
Serotonin Syndrome
Tachypnoea
Urinary Casts
White Blood Cell Count
Increased

Date:11/19/01ISR Number: 3827623-2Report Type:Expedited (15-DaCompany Report #001-0945-M0100592
Age:75 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

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG	(DAILY), PER	Antibody Test Positive Arthralgia	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Asthenia Bronchopneumonia Burning Sensation C-Reactive Protein Increased Cardiac Disorder Chest Pain Circulatory Collapse Diabetes Mellitus Disorientation Dizziness Dysphagia Ecchymosis Electrocardiogram St Segment Depression Fall Fear Gastrooesophageal Reflux Disease Headache Laceration Livedo Reticularis Myalgia Nausea Pain Pallor Paraesthesia Pneumonia Pyrexia Red Blood Cell Sedimentation Rate Increased Sedation Tremor Vasculitis Vision Blurred		Deltasone (Prednisone) Glucotrol (Glipizide) Avapro (Irbesartan) Hydrodiuril (Hydrochlorothiazide) Oxycodone (Oxycodone)	C C C C C		

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Interaction	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
Life-Threatening (TID), PER Hospitalization - ORAL		Respiratory Failure	Professional				
Initial or Prolonged 5 MG (DAILY), Other PER ORAL				Diazepam (Diazepam)	SS		ORAL
1 MG (BID), PER ORAL				Orfidal (Lorazepam)	SS		ORAL
50 MG (BID), PER ORAL				Pethidine Hydrochloride (Pethidine Hydrochloride)	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/01ISR Number: 3827923-6Report Type:Expedited (15-DaCompany Report #200120976EU

Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Alanine Aminotransferase Abnormal	Foreign Other	Leflunomide (Arava) Tablets	PS		ORAL
20 MG/DAY PO		Aspartate Aminotransferase Abnormal Blood Alkaline Phosphatase Increased		Metronidazole (Flagyl) Metronidazole (Flagyl)	SS SS		ORAL
800 MG/DAY PO	15 DAY	Blood Potassium Decreased Blood Potassium Increased Cardiac Failure Chest Pain Depressed Level Of		Flucloxacillin Sodium (Heracillin) Flucloxacillin Sodium (Heracillin)	SS SS		
2.2. G/DAY 25 MG/DAY	15 DAY	Consciousness		Rofecoxib (Vioxx)	SS		
20 MG/DAY PO	1 WK	Fluid Overload Hepatic Enzyme Increased Liver Function Test		Enalapril Maleate (Renitec) Spironolactone	SS SS		ORAL ORAL
50 MG/DAY PO 40 MG/DAY PO	1 WK	Abnormal Localised Infection Metabolic Disorder		Furosemide (Furix) Gabapentin (Neurontin)	SS SS		ORAL ORAL
QD PO		Pain Renal Failure Acute Weight Increased		Folic Acid, Calcium Phosphate (Folacin) Vitamin E Morfin Prednisolon Cisapride (Prepulsid) Cetirizine (Zyrlex) Morphine Sulfate (Dolcontin) Lansoprazole (Lanzo)	C C C C C C C C		

Date:11/20/01ISR Number: 3828046-2Report Type:Expedited (15-DaCompany Report #044-0945-M0100241
Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged PER ORAL	Leukaemia Neutropenia Sepsis	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:11/21/01ISR Number: 3829069-XReport Type:Expedited (15-DaCompany Report #049-0945-M0100129
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG (THREE TIMES A DAY), PER ORAL	Drug Ineffective Myalgia	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Fentanyl	PS C		ORAL

PER ORAL

Mania

Medication Error
Nightmare
Panic Attack
Sedation
Thinking Abnormal
Tremor
Weight Increased

Date:11/21/01ISR Number: 3829785-XReport Type:Expedited (15-DaCompany Report #001-0945-M0101306

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 900 MG (TID), Initial or Prolonged PER ORAL		Abdominal Distension Dysgeusia	Consumer	Neurontin	PS		ORAL
		Headache Hyperhidrosis Movement Disorder Salivary Hypersecretion Thirst Weight Increased		Tegretol (Carbamazepine)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/21/01ISR Number: 3829791-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100387
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetes Mellitus Flatulence	Consumer Health	Neurontin (Gabapentin)	PS		
1200 MG (400 MG, TID)		Neuropathy Peripheral Post Procedural Complication	Professional				

Date:11/23/01ISR Number: 3828765-8Report Type:Expedited (15-DaCompany Report #001-0945-M0101316
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ependymoma Grand Mal Convulsion	Health Professional	Neurontin (Gabapentin)	PS		
4800 MG (DAILY)		Loss Of Consciousness		Celebrex (Celecoxib) Prozac (Fluoxetine Hydrochloride) Ibuprofen (Ibuprofen) Prilosec (Omeprazole)	SS C C C		

Date:11/23/01ISR Number: 3829284-5Report Type:Expedited (15-DaCompany Report #001-0945-990839
Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged PER ORAL		Abnormal Dreams Asthenia	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Confusional State Depressed Level Of Consciousness Hallucination		(Unknown Drugs) Tegretol (Carbamazepine) Unspecified Blood	SS C		

Lethargy	Pressure Medication	C
Listless	Vicodin	
Paranoia	(Paracetamol,	
Pneumonia	Hydrocodone	
Sedation	Bitartrate)	C
Urinary Incontinence	Gammaglobulin	
Weight Increased	Infusions	
	(Immunoglobulin	
	Human Normal)	C

Date:11/23/01ISR Number: 3829287-0Report Type:Expedited (15-DaCompany Report #001-0945-M0101354
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Asthenia	Health	Neurontin			
Other		Epistaxis	Professional	(Gabapentin)	PS		
		Hip Fracture		(Amitriptyline			
		Myeloid Metaplasia		Hydrochloride)	C		
		Pancytopenia		(Paracetamol)	C		
				(Paracetamol,			
				Dextropropoxyphene)	C		
				(Temazepam)	C		
				(Magnesium			
				Hydroxide)	C		

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Freedom Of Information (FOI) Report

(Pregabalin) C

Date:11/23/01ISR Number: 3829587-4Report Type:Expedited (15-DaCompany Report #033-0945-M0100147
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Blood Creatine Phosphokinase Increased	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
PER ORAL				Mediator (Benfluorex Hydrochloride)	SS		ORAL
PER ORAL				Befizal (Bezafibrate)	SS		ORAL
				Glucophage (Metformin Hydrochloride)	C		
				Cibadrex (Benazepril)	C		

Date:11/26/01ISR Number: 3829366-8Report Type:Direct Company Report #
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent 100MG BID Impairment/Damage INITIATE ON		Pain Skin Discolouration Swelling		Neurontin (Gabapentin) 300mg Po Twice Daily	PS		ORAL
8/12/01-300MG							
PO TWICE							
DAILY	1	WK		Ibuprofen	C		

Date:11/26/01ISR Number: 3829678-8Report Type:Expedited (15-DaCompany Report #WAES 01112380
Age:35 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Unevaluable Event	Literature	Cap Crixivan	PS		ORAL
PO			Health	Gabapentin	SS		ORAL
PO			Professional	Quetiapine Fumarate	SS		ORAL
PO				[Composition Unspecified]	SS		ORAL
PO							

Date:11/26/01ISR Number: 3830043-8Report Type:Expedited (15-DaCompany Report #WAES 01111661
Age:21 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Tab Cogentin	PS		ORAL
PO			Health	Haloperidol	SS		
			Professional	Gabapentin	SS		

Date:11/27/01ISR Number: 3831123-3Report Type:Expedited (15-DaCompany Report #061-0945-M0100064
Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cognitive Disorder Confusional State	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
2700 MG (900 MG, TID), PER			Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Methadone
(Methadone) SS ORAL

PER ORAL

(Sertraline
Hydrochloride) C
(Gliclazide) C
(Metronidazole) C

Date:11/27/01ISR Number: 3831142-7Report Type:Expedited (15-DaCompany Report #055-0945-M0100013
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 800 MG		Cough Diabetes Mellitus	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
(DAILY), PER		Dyspnoea					
ORAL		Hypertension					
		Renal Disorder		Insulin (Insulin)	C		
				Captopril (Captopril)	C		
				Dipyridamole (Dipyridamole)	C		
				Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C		
				Furosemide (Furosemide)	C		
				Cinnarizine (Cinnarizine)	C		

Date:11/28/01ISR Number: 3831312-8Report Type:Expedited (15-DaCompany Report #001-0945-M0101201
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG		Cataract Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL

(UNKNOWN),
Macular Oedema
Nausea
PER ORAL
Photosensitivity Reaction Thyroid C
Visual Disturbance Vioxx (Rofecoxib) C

Date:11/28/01ISR Number: 3831318-9Report Type:Expedited (15-DaCompany Report #001-0945-M0101323
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Cough	Consumer	Neurontin			
600 MG		Difficulty In Walking		(Gabapentin)	PS		
(DAILY),				Zestril (Lisinopril)	SS		
5 MG (DAILY),				Levodopa (Levodopa)	C		
				Carbidopa			
				(Carbidopa)	C		
				Comtran	C		
				Benzotinate Perles	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/28/01ISR Number: 3831869-7Report Type:Expedited (15-DaCompany Report #033-0945-M0100061

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MNG (400 Other MG, TID), PER ORAL	Alopecia Haemorrhagic Stroke	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Zocor (Simvastatin) Depakine (Valproate Sodium) Stilnox (Zolpidem) Fraxiparine (Heparin-Fraction, Calcium Salt)	C C C C C		

Date:11/28/01ISR Number: 3832218-0Report Type:Expedited (15-DaCompany Report #PHHO2001IT09280

Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG, QD, ORAL	Drug Interaction Myalgia Rhabdomyolysis	Foreign Study Health Professional	Gleevec Ms Contin (Morphine Sulfate)	PS SS		ORAL ORAL
20 MG, ORAL TID, ORAL		Other	Neurontin (Gabapentin)	SS		ORAL

Date:11/28/01ISR Number: 3832253-2Report Type:Expedited (15-DaCompany Report #033-0945-M0100149

Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Condition Aggravated Intraocular Pressure	Foreign Health	Neurontin (Gabapentin)	PS		

Increased
Open Angle Glaucoma

Professional

Date:11/28/01ISR Number: 3832664-5Report Type:Direct
Age:75 YR Gender:Male I/FU:I

Company Report #

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia Balance Disorder		Taxol (80mg/M2) Q Wk-Iv	PS		
INTRAVENOUS	80MG/M2	Q WK Dyspnoea					
IV		Fatigue		Gabapentin (300 Mg)	SS		ORAL
300 MG TID PO		Pulmonary Embolism					

Date:11/29/01ISR Number: 3832724-9Report Type:Expedited (15-DaCompany Report #001-0945-M0101183
Age:51 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability Other	Amnesia Blood Pressure Increased Depressed Level Of Consciousness Dry Mouth Dysarthria Dysuria 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
		Gait Disturbance Lethargy Miosis					
600 MG (BID), PER ORAL		Nausea Sedation	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Vioxx (Rofecoxib) Zanaflex (Tizanidine)	SS SS		

Date:11/29/01ISR Number: 3833136-4Report Type:Expedited (15-DaCompany Report #PHHO2001IT09280
Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG, QD, ORAL		Blood Creatinine Increased	Foreign Study	Gleevec	PS		ORAL
20 MG, ORAL		Drug Interaction Renal Impairment	Health Professional	Ms Contin (Morphine Sulfate)	SS		ORAL
TID, ORAL		Rhabdomyolysis	Other	Neurontin(Gabapentin)	SS		ORAL

Date:11/30/01ISR Number: 3832922-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101353
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2400 MG (DAILY)		Intraocular Pressure Increased	Consumer	Neurontin (Gabapentin)	PS		
		Post Procedural Complication					

Date:11/30/01ISR Number: 3832954-6Report Type:Expedited (15-DaCompany Report #001-0945-M0101265
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 8000 MG		Vomiting	Consumer Health	Neurontin(Gabapentin)	PS		ORAL
(800MG, TEN TIMES DAILY), PER ORAL			Professional				

Date:11/30/01ISR Number: 3832955-8Report Type:Expedited (15-DaCompany Report #001-0945-M0101288
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2400 MG		Convulsion Dizziness	Consumer Health	Neurontin(Gabapentin)	PS		ORAL
(DAILY), PER ORAL		Infection Loss Of Consciousness	Professional				
		Somnolence Tongue Biting		Zoloft (Sertraline Hydrochloride) Tetracycline (Tetracycline) Hydroxychloroquine(H ydroxychloroquine)	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/30/01ISR Number: 3833592-1Report Type:Expedited (15-DaCompany Report #064-0945-M0100012

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG, PER	Medication Residue	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Professional				

Date:11/30/01ISR Number: 3833595-7Report Type:Expedited (15-DaCompany Report #045-0945-M0100014

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1500 MG (DAILY), PER	Arrhythmia Fall Hypotonia Syncope	Foreign Health Professional Company Representative	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Digoxin (Digoxin) Captol (Captopril) Veraloc (Verapamil Hydrochloride) Metadon (Methadone Hydrochloride) Saroten (Amitriptyline Hydrochloride)	C C C C C		

Date:12/03/01ISR Number: 3833660-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101354

Age:73 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Hospitalization - Initial or Prolonged Other	Anaemia Asthenia Epistaxis Fall Gallbladder Disorder	Consumer Health Professional	Neurontin (Gabapentin) (Amitriptyline Hydrochloride) (Paracetamol)	PS C C		

Haemorrhage	(Paracetamol,	C
Hip Fracture	Dextropropoxyphene)	C
Myeloid Metaplasia	(Temazepam)	C
Pancytopenia	(Magnesium	C
Sepsis	Hydroxide)	C
Somnolence	(Pregabalin)	C
	(Lorazepam)	C
	(Oxycodone	C
	Hydrochloride)	C
	(Morphine Sulfate)	C
	(Bethanechol	C
	Chloride)	C
	(Glimepiride)	C

Date:12/03/01ISR Number: 3833661-6Report Type:Expedited (15-DaCompany Report #001-0945-M0101197
Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Health	Neurontin			
		Finger Deformity	Professional	(Gabapentin)	PS		ORAL
PER ORAL		Tendon Rupture		(Paracetamol,			
				Hydrocodone			
				Bitartrate)	C		
				(Codeine Phosphate,			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Paracetamol) C

Date:12/03/01ISR Number: 3833663-XReport Type:Expedited (15-DaCompany Report #001-0945-M0101368
Age:86 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 200 MG (DAILY), PER ORAL	Condition Aggravated Depressed Level Of Consciousness Difficulty In Walking Disorientation Gastrooesophageal Reflux Disease Incoherent Nausea Oral Pain Pneumonia Somnolence	Consumer	Neurontin (Gabapentin) Morphine (Morphine) Plavix (Clopidogrel) Pancrelipase (Pancrelipase) Aciphex (Rabeprazole Sodium) Metoclopramide (Metoclopramide) Remeron (Mirtazapine) Trazodone (Trazodone)	 PS C C C C C C C C C		ORAL

Date:12/03/01ISR Number: 3833737-3Report Type:Expedited (15-DaCompany Report #001-0945-M0101362
Age:57 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 300 MG (THREE TIMES DAILY), PER ORAL	Convulsion Drug Effect Decreased Insomnia Muscle Twitching	Consumer	Neurontin (Gabapentin) Pepcid (Famotidine) Enalapril (Enalapril) Synthroid (Levothyroxine)	 PS C C C		ORAL

Sodium)

C

Date:12/03/01ISR Number: 3833740-3Report Type:Expedited (15-DaCompany Report #001-0945-M0101315
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia	Consumer	Neurontin (Gabapentin)	PS		
		Epistaxis	Health	Elavil (Amitriptyline Hydrochloride)	SS		
		Myelodysplastic Syndrome	Professional	Paracetamol	C		
		Myeloid Metaplasia		Paracetamol	C		
		Pancytopenia		Dextropropoxyphene	C		
		Urinary Retention		Temazepam	C		
				Magnesium Hydroxide	C		
				Pregabalin	C		
				Lorazepam	C		
				Oxycodone Hydrochloride	C		
				Morphine Sulfate	C		
				Bethanecol Chloride	C		
				Glimepiride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/04/01ISR Number: 3833102-9Report Type:Direct
Age:74 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Anaemia		Neurontin	PS		
SEE						
Hospitalization -	Anxiety					
PRESCRIPTION						
Initial or Prolonged	Asthenia					
	Drug Ineffective					

Date:12/06/01ISR Number: 3836202-2Report Type:Direct
Age:56 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Acute Respiratory		Neurontin (600 Mg			
Initial or Prolonged	Distress Syndrome		Per Tube Q 1 D)	PS		
600 MG PER						
	Arrhythmia					
TUBE QID						
	Cardiac Failure		Dyazide	C		
	Congestive		Kcl	C		
			Ferrous Sulfate	C		
			Valproic Acid	C		
			Liorisel	C		
			Celebrex	C		
			Vit E	C		
			Congentin	C		
			Seroquel	C		
			Provent	C		
			Glucose	C		
			Levaquin	C		

Date:12/06/01ISR Number: 3836527-0Report Type:Expedited (15-DaCompany Report #001-0945-M0101376
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Anaphylactic Reaction	Consumer	Neurontin			
	Asthma		(Gabapentin)	PS		ORAL
900 MG						

(DAILY), PER

Blood Pressure Increased

Irritable Bowel Syndrome

ORAL

Loss Of Consciousness

Multiple Allergies

Urticaria

Flovent (Fluticasone Propionate)	C
Serevent (Salmeterol Xinafoate)	C
Ventolin (Salbutamol)	C
Unspecified Nebulizer	C

Date:12/06/01ISR Number: 3836532-4Report Type:Expedited (15-DaCompany Report #2014953

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Drug Toxicity	Health	Oxycodone			
		Toxicologic Test Abnormal	Professional	Hydrochloride			
			Other	(Similar To Nda			
				20-553)	PS		
				Gabapentin	SS		
				Temazepam	SS		
				Caffeine Anhydride	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/06/01ISR Number: 3836538-5Report Type:Expedited (15-DaCompany Report #001-0945-M0101372

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Endometrial Cancer	Consumer	Neurontin (Gabapentin)	PS		ORAL
1500 MG (500							
MG, TID), PER							
ORAL							

Date:12/06/01ISR Number: 3837433-8Report Type:Expedited (15-DaCompany Report #2014944

Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Tamponade	Health	Oxycodone			
		Cardiomegaly	Professional	Hydrochloride	PS		
		Coronary Artery	Other	Acetaminophen	SS		
		Atherosclerosis		Gabapentin	SS		
		Drug Toxicity		Propoxyphene Hcl	SS		
		Hepatic Steatosis		Mirtazipine Hcl	SS		
		Pericardial Haemorrhage		Alprazolam	SS		
		Pericarditis					

Date:12/07/01ISR Number: 3835609-7Report Type:Expedited (15-DaCompany Report #B0128407A

Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blister		Wellvone	PS	Glaxo Wellcome	ORAL
2UNIT per day							
Initial or Prolonged		Cerebral Toxoplasmosis		Forlax	SS		ORAL
1UNIT per day 4	DAY						
		Dermatitis Exfoliative		Videx	SS		ORAL
1UNIT per day							
		Erythema		Zerit	SS		ORAL
2UNIT per day							
				Neurontin	SS		ORAL
5UNIT per day							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Ageusia Dysgeusia	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), PER ORAL		Glossodynia					
12 MG (TID), PER ORAL		Influenza Like Illness Tongue Ulceration		Gabitril (Tiagabine)	SS		ORAL
		Vomiting Weight Decreased		(Metaxalone) (Lidocaine Hydrochloride)	C C		

Date:12/07/01ISR Number: 3837255-8Report Type:Expedited (15-DaCompany Report #033-0945-M0100152
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 MG, BID), PER ORAL		Rash Erythematous	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Lovenox (Heparin-Fraction, Sodium Salt)	SS		

(DAILY),

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Freedom Of Information (FOI) Report

SUBCUTANEOUS

Valproate Sodium) C

Date:12/07/01ISR Number: 3837544-7Report Type:Expedited (15-DaCompany Report #200121709FR
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Erythema Pneumonia	Foreign Other	Heparin-Fraction, Sodium Salt (Lovenox) Solution For Injection	PS		
SUBCUTANEOUS	0.4 ML QD SC			Albendazole (Zentel) Tablet	SS		ORAL
1500 MG/DAY				Omeprazole(Mopral) Capsules	SS		ORAL
PO				Roxithromycin (Rulid) Tablets	SS		ORAL
20 MG QD PO				Metronidazole (Flagyl) Tblets	SS		ORAL
100 MG QD PO	11 DAY			Gabapentin (Neurontin) Capsules	SS		ORAL
1500 MG /DAY							
PO							
300 MG BID PO	2 DAY						

Date:12/07/01ISR Number: 3837727-6Report Type:Expedited (15-DaCompany Report #001-0945-M0101373
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blepharospasm Fall	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG Disability (DAILY), PER		Visual Disturbance					

ORAL

Date:12/07/01ISR Number: 3837731-8Report Type:Expedited (15-DaCompany Report #001-0945-M0101157

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Unevaluable Event	Consumer	Neurontin			
Initial or Prolonged		Health	(Gabapentin)	PS		
800 MG (BID)		Professional				

Date:12/10/01ISR Number: 3837989-5Report Type:Expedited (15-DaCompany Report #001-0945-M0101387

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Blood Glucose Increased	Consumer	Neurontin			
Initial or Prolonged	Disturbance In Attention		(Gabapentin)	PS		ORAL
600 MG (300						
Other	Dry Mouth					
MG, BID), PER	Headache					
ORAL	Nausea		Lipitor			
	Pain		(Atorvastatin)	C		
	Vomiting		Zyrtec (Cetirizine			
			Hydrochloride)	C		
			Lasix (Furosemide)	C		
			Potassium			
			(Potassium)	C		
			Accupril (Quinapril			
			Hydrochloride)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Minitran(Glyceryl
Trinitrate) C
Glucophage
(Metformin
Hydrochloride) C
Plavix (Clopidogrel) C
Tiazac (Diltiazem
Hydrochloride) C
Benadryl
(Diphenhydramine
Hydrochloride) C
Metoprolol
(Metoprolol) C
Flovent (Fluticasone
Propionate) C
Aspirin
(Acetylsalicylic
Acid) C
Albuterol Inhaler
(Salbutamol) C
Albuterol Solution
(Salbutamol) C
Multivitamin
(Vitamins Nos) C

Date:12/11/01ISR Number: 3837411-9Report Type:Direct
Age: Gender: I/FU:I

Company Report #USP 54642

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin	PS	Pfizer	
Other		Medication Error		Noroxin (Norfloxacin)	SS	Merck	

Date:12/11/01ISR Number: 3838358-4Report Type:Expedited (15-DaCompany Report #GBR003253
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Meridia	PS		ORAL
Other		Blood Pressure Increased	Health	Hydrocodone/Apap	SS		
10 MG QD PO		Cystitis	Professional				
OTHER OTHER							

1 TAB QOD	Diabetes Mellitus	Other	Soma	SS	
600 MG TID PO	Nausea		Neurontin	SS	ORAL
260 MG NOCTE	Sinusitis		Reglan	SS	
PO	Vomiting		Quinine Sulfate	SS	ORAL
1 TAB QD PO			Mobic	SS	ORAL
			Advil	SS	

Date:12/11/01ISR Number: 3838405-XReport Type:Expedited (15-DaCompany Report #2014962
Age:43 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Blood Disorder
Initial or Prolonged	Blood Testosterone
Other	Decreased
	Electroencephalogram
	Abnormal
	Grand Mal Convulsion
	Hypothyroidism

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Oedema Peripheral Pain In Extremity Polycythaemia Vera	Report Source	Product	Role	Manufacturer	Route
560 MG Q12H			Health Professional	Oxycontin Cr (Oxycodone Hydrochloride)	PS		ORAL
PO	1 YR			Neurontin (Gabapentin)	SS		ORAL
2100 MG HS PO				Vioxx (Rofecoxib)	SS		ORAL
PO				Celebrex (Celecoxib)	SS		ORAL
PO				Dalmane (Flurazepam)	C		
				Folic Acid	C		
				Prozac (Fluoxetine)	C		
				Vistaril (Hydroxizine Pamoate)	C		
				Synthroid (Levothyroxine)	C		
				Lasix (Furosemide)	C		
				Aspirin (Acetylsalicylic Acid)	C		
				Multivitamin	C		
				Colace (Docusate)	C		

Date:12/11/01ISR Number: 3838488-7Report Type:Expedited (15-DaCompany Report #049-0945-M0100130
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Confusional State Coordination Abnormal	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		Uraemic Neuropathy	Professional Company Representative				

Outcome	PT
Hospitalization -	Abdominal Adhesions
Initial or Prolonged	Abdominal Pain
Disability	Atelectasis
Other	Back Pain
	Bronchitis
	Burning Sensation
	Chest X-Ray Abnormal
	Cholelithiasis
	Colonic Polyp
	Cough
	Crepitations
	Depression
	Diabetes Mellitus
	Inadequate Control
	Difficulty In Walking
	Dyspnoea Exertional
	Ear Discomfort
	Fatigue
	Foot Deformity

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
200 MG (DAILY)		Fungal Infection				
		Fungal Rash				
		Fungal Skin Infection				
		Glucose Urine Present				
		Haemorrhoidal Haemorrhage	Rezulin (Troglitazone)	PS		
		Headache				
		Hepatic Cirrhosis	Vasotec (Enalapril			
		Hepatic Steatosis	Maleate)	SS		
		Hepatitis	Hyoscyamine			
		Hepatomegaly	(Hyoscyamine)	SS		
		Hiatus Hernia	Amitriptyline			
		Hypergammaglobulinaemia	(Amitriptyline)	SS		
		Hypertension	Neurontin			
		Insomnia	(Gabapentin)	SS		ORAL
SEE IMAGE		Ketonuria	Dilaudid			
		Neck Pain	(Hydromorphone			
		Oedema Peripheral	Hydrochloride)	SS		
		Ovarian Cyst Ruptured	Trazodone			
		Pharyngolaryngeal Pain	(Trazodone)	SS		
		Plantar Fasciitis	Diabeta			
		Proteinuria	(Glibenclamide)	C		
		Rales	Glucophage			
		Rash	(Metformin			
		Rash Maculo-Papular	Hydrochloride)	C		
		Rhinitis	Spironolactone			
		Seasonal Allergy	(Spironolactone)	C		
		Sinusitis	Zyrtec (Cetirizine			
		Tinea Pedis	Hydrochloride)	C		
		Umbilical Hernia	Vancenase			
		Urinary Tract Infection	(Beclometasone			
		Uterine Leiomyoma	Dipropionate)	C		
		Viral Pharyngitis	Axid (Nizatidine)	C		
		Vomiting	Glyburide			
		Weight Increased	(Glibenclamide)	C		
			Insulin 70/30			
			(Insulin)	C		
			St. Johns Wart			
			(Hypericum			
			Perforatum)	C		
			Zithromax			
			(Azithromycin)	C		
			Zovia 1/35	C		
			Guiatex			

(Phenylephrine
Hydrochloride,
Guaifenesin,
Phenylpropanolamine C
Propoxyphene
(Dextropropoxyphene) C
Darvon-N
(Dextropropoxyphene
Napsilate) C
Cephalexin
(Cefalexin) C
Tylenol
(Paracetamol) C
Benadryl
(Diphenhydramine
Hydrochloride) C
Demerol (Pethidine
Hydrochloride) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vicodin (Paracetamol, Hydrocodone Bitartrate)	C
Heparin (Heparin)	C
Phenergen (Promethazine Hydrochloride)	C
Cefazolin (Cefazolin)	C
Compazine (Prochlorperazine Edisylate)	C
Vitamins	C
Dovonex (Calcipotriol)	C
Ultravate (Ulobetasol Propionate)	C

Date:12/12/01ISR Number: 3839093-9Report Type:Expedited (15-DaCompany Report #001-0945-M0101395
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Health	Neurontin			
		Hallucination	Professional	(Gabapentin)	PS		
		Paranoia		Xanax (Alprazolam)	SS		
		Screaming					

Date:12/12/01ISR Number: 3839467-6Report Type:Expedited (15-DaCompany Report #001-0945-M0101157
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1200 MG (300	No Adverse Drug Effect	Consumer	Neurontin			
			Health	(Gabapentin)	PS		
			Professional				

MG, QID)

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour Anhedonia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1200 MG (TID), PER ORAL		Cognitive Disorder Diet Refusal	Professional				
				Aspegic (Acetylsalicylate Lysine)	C		
				Calcidose Vitamin D (Colecalciferol, Calcium Carbonate)	C		
				Cetornan (Ornithineoxoglurate)	C		
				Hydrosol Polyvitamin (Vitamins Nos)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/12/01ISR Number: 3839959-XReport Type:Expedited (15-DaCompany Report #046-0945-M0100062

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2.4 GM Other (DAILY), PER ORAL		Blood Alkaline Phosphatase Increased Cardiac Failure Chest Pain	Foreign Consumer	Neurontin Tablets(Gabapentin)	PS		ORAL
40 MG (DAILY), PER ORAL		Depressed Level Of Consciousness Fluid Retention		Furix(Furosemide)	SS		ORAL
50 MG (DAILY), PER ORAL		Infected Skin Ulcer Liver Function Test Abnormal Metabolic Disorder		Spironolakton(Spiron olactone)	SS		ORAL
20 MG (DAILY), PER ORAL		Renal Failure Acute Skin Ulcer Weight Increased		Renitec (Enalapril Maleate)	SS		ORAL
25 MG (DAILY), PER ORAL				Vioxx(Rofecoxib)	SS		ORAL
800 MG (DAILY), PER ORAL				Flagyl(Metronidazole)	SS		ORAL
				Heracillin(Flucloxac			

PER ORAL
 20 MG
 (DAILY), PER
 ORAL

illin Sodium) SS ORAL
 Arava(Leflunomide) SS ORAL
 Folic Acid, Calcium Phosphate C
 Tocopheryl Acetate C
 Paracetamol C
 Prednisolone C
 Morphine C
 Cisapride C
 Cetirizine C
 Morphine Sulfate C
 Lansoprazole C
 Zolpidem Tartrate C
 Loperamide C
 Calcium Carbonate C
 Ipratropium Bromide,
 Salbutamol Sulfate C

Date:12/12/01ISR Number: 3840282-8Report Type:Expedited (15-DaCompany Report #S01-USA-02257-01
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Health Professional Company Representative	Celexa (Citalopram Hydrobromide) Quetiapine (Quetiapine Fumarate) Neurontin (Gabapentin) Desyrel (Trazodone Hydrochloride)	PS SS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/12/01ISR Number: 3840290-7Report Type:Expedited (15-DaCompany Report #001-0945-M0101407
 Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2400 MG (DAILY), PER ORAL		Condition Aggravated Dizziness Feeling Cold Visual Acuity Reduced	Consumer	Neurontin (Gabapentin)	PS		ORAL
				(Verapamil Hydrochloride) (Clopidogrel)	C C		

Date:12/13/01ISR Number: 3839676-6Report Type:Expedited (15-DaCompany Report #001-0073-M0100506
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (100 MG, TID)		Chest Pain Drug Level Above Therapeutic	Consumer	Dilantin (Phenytoin Sodium)	PS		
300 MG (100 MG, TID)		Dyspepsia Epilepsy Grand Mal Convulsion		Neurontin (Gabapentin)	SS		
2 DROPS				Hall'S (Menthol, Eucaliptus Oil)	SS		
				Phenobarbital	C		

Date:12/13/01ISR Number: 3840213-0Report Type:Direct Company Report #
 Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 80MG QHS		Alanine Aminotransferase		Simvastatin	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Initial or Prolonged 50MG QHS		Increased		Trazodone	SS		
300MG TID				Gabapentin	SS		
Date:12/14/01ISR Number: 3839432-9Report Type:Expedited (15-DaCompany Report #046-0945-M0100062 Age: Gender:Female I/FU:F							
Hospitalization - Initial or Prolonged 2.4 GM Other (DAILY), PER ORAL		Blood Alkaline Phosphatase Increased Blood Creatinine Increased	Foreign Consumer	Neurontin Tablets (Gabapentin)	PS		ORAL
40 MG (DAILY), PER ORAL		Blood Potassium Increased Cardiac Failure Chest Pain		Furix (Furosemide)	SS		ORAL
50 MG (DAILY), PER ORAL		Depressed Level Of Consciousness Infected Skin Ulcer Liver Function Test		Spironolakton (Spironolactone)	SS		ORAL
20 MG (DAILY), PER ORAL		Abnormal Oedema Pain Weight Increased		Renitec (Enalapril Maleate)	SS		ORAL
25 MG (DAILY), PER ORAL				Vioxx (Rofecoxib)	SS		ORAL
800 MG (DAILY), PER				Flagyl (Metronidazole)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Heracillin (Flucloxacillin Sodium)	SS	ORAL
--	----	------

PER ORAL

Arava (Leflunomide)	SS	ORAL
---------------------	----	------

20 MG

(DAILY), PER

ORAL

Folic Acid, Calcium	
Phosphate	C
Tocopheryl Acetate	C
Paracetamol	C
Prednisolone	C
Morphine	C
Cisapride	C
Cetirizine	C
Morphine Sulfate	C
Lansoprazole	C
Zolpidem Tartrate	C
Loperamide	C
Calcium Carbonate	C
Ipratropium Bromide, Salbutamol Sulfate	C

Date:12/14/01ISR Number: 3839594-3Report Type:Expedited (15-DaCompany Report #001-0945-M0101415

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged DAILY, PER Other ORAL	Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL
			(Morphine Sulfate)	C		
			(Methocarbamol)	C		

Date:12/14/01ISR Number: 3839596-7Report Type:Expedited (15-DaCompany Report #001-0945-M0101398

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

Outcome
Hospitalization -
Initial or Prolonged
Other

PT
Arthralgia
Arthropathy
Blood Albumin Decreased
Blood Calcium Decreased
Blood Creatinine
Decreased
Blood Potassium Decreased
Dilatation Atrial
Electrocardiogram
Abnormal
Haematocrit Increased
Intentional Misuse
Mental Status Changes
Multiple Sclerosis
Muscular Weakness
Neck Pain
Pco2 Decreased
Pco2 Increased
Platelet Count Decreased
Po2 Decreased
Poisoning
Rhinitis Allergic
Road Traffic Accident

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Toxicologic Test Abnormal

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1200 MG (DAILY), PER ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
	20 MG		Copaxone (Glatiramer Acetate)	SS		
0.25 MG (DAILY), PER ORAL			Tylenol W/Codeine No. 3 (Codeine Phosphate, Paracetamol) Xanax (Alprazolam)	SS SS		ORAL
			Roxicet (Paracetamol, Oxycodone Hydrochloride) (Fluticasone Propionate) (Tolterodine Tartrate)	SS C C		

Date:12/14/01ISR Number: 3840652-8Report Type:Expedited (15-DaCompany Report #033-0945-M0100154

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1200 MG (TID)		Diabetes Mellitus	Foreign Health	Neurontin (Gabapentin)	PS		

Professional (Propranolol) C
(Acebutolol) C
(Atenolol) C

Date:12/17/01ISR Number: 3839972-2Report Type:Direct
Age:77 YR Gender:Female I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 5MG AM/10MG Initial or Prolonged HS PO 300MG BID	Asthenia Blood Glucose Increased Blood Pressure Increased Lethargy		Zyprexa Neurontin	PS SS		ORAL

Date:12/17/01ISR Number: 3840101-XReport Type:Direct
Age:45 YR Gender:Male I/FU:I

Company Report #

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability Other 4 X 300 MG PER DAY 4 MON	Eye Irritation Eyelid Function Disorder Visual Disturbance		Neurontin Parke-Davis Pfizer. Inc	PS	Parke-Davis Pfizer. Inc	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/17/01ISR Number: 3840251-8Report Type:Expedited (15-DaCompany Report #001-0945-M0101421

Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Uveitis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
100 MG							
(DAILY), PER							
ORAL				(Amfebutamone Hydrochloride)	C		

Date:12/17/01ISR Number: 3840252-XReport Type:Expedited (15-DaCompany Report #001-0945-M0101319

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cholelithiasis	Health Professional	Neurontin (Gabapentin)	PS		
Initial or Prolonged		Hepatitis					
900 MG (300							
Other		Hepatotoxicity	Company Representative	(Tazobactam Sodium)	SS		
MG, TID)							
INTRAVENOUS	INTRAVENOUS						

Date:12/17/01ISR Number: 3840375-5Report Type:Expedited (15-DaCompany Report #049-0945-M0100098

Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG		Myasthenia Gravis					
(DAILY), PER							
ORAL				Immunosuppressive Agents	C		

Date:12/17/01ISR Number: 3840834-5Report Type:Expedited (15-DaCompany Report #001-0073-M0100514
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Dyskinesia Fall	Consumer	Dilantin (Phenytoin Sodium)	PS		ORAL
PER ORAL						
Other 900 MG (300 MG, TID), PER ORAL	Muscle Rigidity Psychomotor Hyperactivity Somnolence		(Gabapentin)	SS		ORAL
			(Levothyroxine Sodium)	C		
			(Amantadine)	C		

Date:12/18/01ISR Number: 3841299-XReport Type:Expedited (15-DaCompany Report #200121819FR
Age:39 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Skin Exfoliation	Foreign Other	Lovenox 40 Mg/0.4 Ml Solution For Injection	PS		
SUBCUTANEOUS	40 MG QD SC					
1 U QD PO			Didanosine (Videx)	SS		ORAL
			Gabapentin (Neurontin) Capsules	SS		ORAL
5 U PO						
2 U PO			Stavudine (Zerit)	SS		ORAL
			Macrogol 4000 (Forlax 10 G) Powder (Lyophilisate)	SS		ORAL
10 G QD PO	4 DAY					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

500 MG PO Atovaquone (Wellvone) Coated Tablets SS ORAL

Date:12/19/01ISR Number: 3842256-XReport Type:Expedited (15-DaCompany Report #039-0945-M0100012
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG	Rash Scarlatiniform	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER							
ORAL							

(Acenocoumarol) C
(Allopurinol) C
(Digoxin) C
(Phenobarbital Sodium) C
(Furosemide) C
(Omeprazole) C
(Tiagabine) C
(Verapamil Hydrochloride) C

Date:12/19/01ISR Number: 3842746-XReport Type:Expedited (15-DaCompany Report #001-0945-M0101189
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	600 MG (300 MG, BID), PER	Balance Disorder Condition Aggravated	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Epistaxis					
		Hyperhidrosis					
		Impetigo					
		Skin Infection					
		Staphylococcal Infection					

Stevens-Johnson Syndrome

Date:12/19/01ISR Number: 3846642-3Report Type:Expedited (15-DaCompany Report #033-0945-M0100159
 Age:86 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG (DAILY), PER ORAL	Condition Aggravated Neuromyopathy Paraesthesia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			(Acetylsalicylate Calcium)	C		
			(Zolpidem)	C		
			(Enalapril)	C		
			(Amlodipine)	C		

Date:12/20/01ISR Number: 3843576-5Report Type:Expedited (15-DaCompany Report #033-0945-M0100157
 Age:87 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Confusional State Depressed Level Of Consciousness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
800 MG (400 MG, DAILY) PER OERAL		Encephalopathy Hypernatraemia Inflammation Renal Failure Somnolence Status Epilepticus	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Clopidogrel	C
Furosemide	C
Colchicine	C
Pantoprazole	C

Date:12/21/01
Age:53 YR
Gender:Female
I/FU:I

Company Report #

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other TAKEN 1 AT N PILL			Dyspepsia Faecal Incontinence		Neurontin	PS		
					Effexor Xr 75 Mg Wyeth Ayerst Celexa 20	SS SS	Wyeth Ayerst	

Date:12/24/01
Age:39 YR
Gender:Female
I/FU:I

Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-11638772

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Delirium Tremens Dermatitis Exfoliative Hallucination Nervous System Disorder Toxoplasmosis		Zerit Caps Videx Neurontin Lovenox Forlax Wellvone	PS SS SS C C C	Bristol-Myers Squibb Company Bristol-Myers Squibb Company	ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Cognitive Disorder	Foreign	Docetaxel	PS		
INTRAVENOUS 169 MG	Q3W IV 85 DAY					
Initial or Prolonged	Confusional State	Other	Gabapentin	SS		ORAL
500 MG TID PO 2 WK						
	Constipation		Benzydamine	C		
	Diarrhoea		Fentanyl	C		
	Fall		Dalteparin Sodium	C		
	Pulmonary Embolism		Ranitidine	C		
	Ulna Fracture		Salbutamol			
			W/Ipratropium	C		
			Dexamethasone	C		
			Chloramphenicol	C		
			Metoclopramide	C		
			Lactulose	C		
			Ciprofloxacin	C		
			Ampicillin	C		
			Frusemide	C		
			Nystatin	C		
			Cyclizine	C		
			Loperamide	C		
			Paracetamol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Co-Danthramer C

Date:12/26/01ISR Number: 3844613-4Report Type:Expedited (15-DaCompany Report #034-0945-M0100011

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death (TID), PER Life-Threatening ORAL		Drug Interaction	Foreign Health	Gabapentin	PS		ORAL
Hospitalization - 5 MG (DAILY), Initial or Prolonged PER ORAL		Respiratory Failure	Professional	Diazepam (Diazepam)	SS		ORAL
Other 1 MG (BID), PER ORAL				Orfidal (Lorazepam)	SS		ORAL
				Pethidine Hydrochloride (Pethidine Hydrochloride)	SS		
INTRAMUSCULAR (DAILY), INTRAMUSCULAR	50 MG						

Date:12/26/01ISR Number: 3844614-6Report Type:Expedited (15-DaCompany Report #064-0945-M0100012

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG, PER ORAL		Medication Residue	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 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		

Age:44 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Other	Confusional State Dysarthria Dyskinesia Electrocardiogram Abnormal Hallucination, Visual Mental Impairment Persecutory Delusion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Suspiciousness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1200 MG	(TID), PER	Foreign Health Professional Company	Neurontin (Gabapentin)	PS		ORAL
1500 MG	(BID), PER	Representative	(Valproic Acid)	SS		ORAL
			(Pyrimethamine)	C		
			(Clindamycin)	C		
			(Efavirenz)	C		
			(Stavudine)	C		
			(Lamivudine)	C		

Date:12/26/01
 Age:72 YR
 Gender:Female
 I/FU:I
 ISR Number: 3844814-5
 Report Type:Expedited (15-DaCompany Report #044-0945-M0100281

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1500 MG (500	Cognitive Disorder Confusional State	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other (MG, TID), PER		Fall					
		Ulna Fracture					
ORAL				(Docetaxel)	SS		
INTRAVENOUS	169 MG (EVERY						
THREE WEEKS),							
INTRAVENOUS							
				Benzydamine	C		
				Ranitidine	C		
				Dexamethasone	C		
				Furosemide	C		

Paracetamol C
Dantron, Poloxamer C

Date:12/26/01ISR Number: 3845066-2Report Type:Expedited (15-DaCompany Report #033-0945-M0100158
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (DAILY), PER		Cerebellar Syndrome Confusional State	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Disorientation	Professional				
(DAILY), PER		Erythema		(Didanosine)	SS		ORAL
ORAL		Hallucination					
SUBCUTANEOUS	40 MG	Skin Exfoliation Skin Nodule		(Heparin-Fraction, Sodium Salt)	SS		
(DAILY),		Toxoplasmosis					
SUBCUTANEOUS				(Stavudine)	SS		ORAL
(DAILY), PER							
ORAL				(Macrogol)	SS		ORAL
10 GM							
(DAILY), PER							
ORAL				(Atovaquone)	SS		ORAL
500 MG							
(DAILY), PER							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/26/01ISR Number: 3845070-4Report Type:Expedited (15-DaCompany Report #049-0945-M0100143
 Age:69 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 800 MG (DAILY), PER ORAL	Collagen Disorder Juvenile Arthritis Pericardial Effusion	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:12/26/01ISR Number: 3845216-8Report Type:Expedited (15-DaCompany Report #001-0945-M0101183
 Age:51 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG Disability (ONCE), PER Other ORAL 50 MG (ONCE), PER ORAL	Amnesia Blood Pressure Increased Confusional State Difficulty In Walking Dysarthria Dysuria	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
2 MG (ONCE), PER ORAL	Fatigue Gait Disturbance Hemiparesis Lethargy Loss Of Consciousness Memory Impairment Mental Status Changes Miosis Nausea Photophobia Somnolence Speech Disorder		Vioxx (Rofecoxib) Zanaflex (Tizanidine) (Clopidogrel)	SS SS C		ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Agitation	Consumer	Neurontin			
Initial or Prolonged	Anhedonia		(Gabapentin)	PS		
Disability	Anxiety		(Atropine)	SS		
1 MG						
	Back Pain		(Oxygen)	C		
	Confusional State					
	Emotional Distress					
	Eye Pain					
	Faecal Incontinence					
	Fall					
	Haematoma					
	Hip Fracture					
	Joint Dislocation					
	Lethargy					
	Overdose					
	Pain					
	Respiratory Distress					
	Sleep Walking					
	Staphylococcal Infection					
	Urinary Tract Infection					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/26/01ISR Number: 3845218-1Report Type:Expedited (15-DaCompany Report #001-0945-M0101268

Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG (QID), PER ORAL		Coronary Artery Occlusion Tremor	Consumer	Neurontin (Gabapentin)	PS		ORAL
10 MG (DAILY), PER ORAL				Lipitor (Atorvastatin)	SS		ORAL
				(Glyceryl Trinitrate)	C		
				(Furosemide)	C		
				(Enalapril Maleate)	C		
				(Digoxin)	C		
				(Amlodipine Besilate)	C		
				(Valproate Semisodium)	C		
				(Theophylline)	C		
				(Salbutamol)	C		
				(Sertraline Hydrochloride)	C		
				(Amitriptyline)	C		

Date:12/26/01ISR Number: 3845228-4Report Type:Expedited (15-DaCompany Report #001-0945-M0101368

Age:86 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (TWICE DAILY AS		Condition Aggravated Depressed Level Of Consciousness	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

NEEDED), PER Difficulty In Walking

Disorientation

ORAL

Fatigue	Morphine (Morphine)	C
Gastroesophageal Reflux Disease	Plavix (Clopidogrel)	C
Incoherent	Pancrelipase (Pancrelipase)	C
Nausea	Aciphex (Rabeprazole Sodium)	C
Pneumonia	Metoclopramide (Metoclopramide)	C
Somnolence	Remeron (Mirtazapine)	C
	Trazodone (Trazodone)	C

Date:12/26/01ISR Number: 3845259-4Report Type:Expedited (15-DaCompany Report #049-0945-M0100092
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bone Disorder Fibrosis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER		Pathological Fracture	Professional				

ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/26/01ISR Number: 3845360-5Report Type:Expedited (15-DaCompany Report #039-0945-M0100013

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	600 MG (TWICE DAILY) PER ORAL	Depression Sedation	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:12/27/01ISR Number: 3864582-0Report Type:Periodic Company Report #WAES 01112908

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
25 MG/ PO		Asthenia	Health	Tab Vioxx	PS		ORAL
		Drug Interaction Fatigue	Professional Company Representative	Ultram Zanaflex Neurontin	SS SS SS		

Date:12/28/01ISR Number: 3845543-4Report Type:Direct Company Report #

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - INTRAVENOUS Initial or Prolonged WK	80MG /M2 IV Q	Chest Discomfort Chest Pain		Paclitaxel	PS		
600MG TID QD		Computerised Tomogram		Gabapentin	SS		
		Abnormal Dyspnoea Lung Infection Po2 Po2 Abnormal Sputum Abnormal		Augmentin Albuterol Inh Tamoxifen Dyazide Ativan Mvi	C C C C C C		

Date:12/28/01ISR Number: 3846712-XReport Type:Expedited (15-DaCompany Report #061-0945-M0100089
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder Confusional State	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
1200 MG, PER		Dizziness	Professional				
ORAL		Fall Nausea Sedation	Company Representative	Chemotherapy	C		

Date:12/28/01ISR Number: 3846733-7Report Type:Expedited (15-DaCompany Report #002-0945-M0100152
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Oedema Pneumonia	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
PER ORAL			Professional				

Date:12/31/01ISR Number: 3846639-3Report Type:Expedited (15-DaCompany Report #044-0945-M0100281
Age:72 YR Gender:Female I/FU:F

Outcome
Hospitalization -
Initial or Prolonged

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Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1500 MG (500 MG, TID), PER ORAL		Cardiac Failure Congestive	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Cognitive Disorder	Professional				
		Confusional State					
INTRAVENOUS THREE WEEKS)	169 MG	Constipation (EVERY Decreased Appetite		(Docetaxel)	SS		
INTRAVENOUS		Dyspnoea					
		Eye Infection		(Benzydamine)	C		
		Fall		(Ranitidine)	C		
		Mouth Ulceration		(Dexamethasone)	C		
		Nausea		(Furosemide)	C		
		Neutropenia		(Paracetamol)	C		
		Oral Candidiasis		(Dantron, Poloxamer)	C		
		Ulna Fracture					

Date:12/31/01ISR Number: 3846838-0Report Type:Expedited (15-DaCompany Report #002-0945-M0100153
Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), PER ORAL		Anger Condition Aggravated	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
		Confusional State	Professional				
		Coordination Abnormal		(Etidronate Disodium)	C		
		Disorientation		(Risedronic Acid)	C		
		Fall		(Calcium Carbonate)	C		
		Hallucination		(Ergocalciferol)	C		
		Lumbar Vertebral Fracture		(Fluoxetine)	C		
		Memory Impairment		(Morphine Sulfate)	C		
		Mental Impairment		(Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine			
		Muscle Spasms					
		Paralysis					
		Pneumonia					

Pulmonary Fibrosis
Urinary Incontinence

Hydrochloride,
Retinol, Riboflavin, C
(Hydroxychloroquine
Sulfate C
(Prednisone) C
(Baclofen) C

Date:12/31/01ISR Number: 3847037-9Report Type:Expedited (15-DaCompany Report #001-0945-M0101278
Age:53 YR Gender:Male I/FU:I

Outcome PT
Disability Anorexia
Other Bursitis
Cervical Spine Flattening
Cyst
Depressed Level Of
Consciousness
Depression
Disturbance In Attention
Dizziness
Dyspepsia
Exostosis
Feeling Abnormal
Insomnia

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Sensation In Eye Condition Aggravated	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
2400 MG		Dizziness	Professional				
(DAILY), PER		Drug Ineffective					
ORAL		Visual Acuity Reduced		(Verapamil Hydrochloride) (Clopidogrel)	C C		

Date:12/31/01ISR Number: 3847121-XReport Type:Expedited (15-DaCompany Report #001-0945-M0101445
Age:90 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia Drug Ineffective	Health Professional	Neurontin (Gabapentin)	PS		ORAL
100 MG		Drug Toxicity					
(DAILY), PER		Eyelid Ptosis					
ORAL		Lethargy		(Tramadol Hydrochloride) (Digoxin) (Lisinopril) (Trazone)	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/31/01ISR Number: 3847122-1Report Type:Expedited (15-DaCompany Report #001-0945-M0101443

Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Arthralgia Erythema	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Hypoaesthesia Intervertebral Disc Protrusion Oedema Peripheral		(Clonidine) (Benazepril Hydrochloride) (Hydrochlorothiazide)	C C C		

Date:01/02/02ISR Number: 3847890-9Report Type:Direct Company Report #CTU 158173

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1000MG TID		Fall		Valproic Acid	PS		
500MG QD		Somnolence		Clonazepam	SS		
300MG TID				Gabapentin	SS		

Date:01/03/02ISR Number: 3847908-3Report Type:Expedited (15-DaCompany Report #001-0945-M0101387

Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 MG, BID), PER ORAL		Blood Glucose Increased Disturbance In Attention	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other		Dry Mouth	Professional				
		Headache					
		Infection Pain Vomiting		Lipitor (Atorvastatin) Zyrtec (Cetirizine Hydrochloride) Lasix (Furosemide)	C C C		

Potassium	C
(Potassium)	
Accupril (Quinapril	
Hydrochloride)	C
Minitran (Glyceryl	
Trinitrate)	C
Glucophage	
(Metformin	
Hydrochloride)	C
Plavix (Clopidogrel)	C
Prilosec	
(Omeprazole)	C
Tiazac (Diltiazem	
Hydrochloride)	C
Benadryl	
(Diphenhydramine	
Hydrochloride)	C
Metoprolol	
(Metoprolol)	C
Flovent (Fluticasone	
Propionate)	C
Aspirin	
(Acetylsalicylic	
Acid)	C
Albuterol Inhaler	
(Salbutamol)	C
Albuterol Solution	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Salbutamol) C
 Multivitamin
 (Vitamins Nos) C

Date:01/03/02ISR Number: 3847963-0Report Type:Expedited (15-DaCompany Report #044-0945-M0100287
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Abdominal Pain	Foreign	Neurontin			
PER ORAL		Drug Withdrawal Syndrome	Health	(Gabapentin)	PS		ORAL
		Malaise	Professional				
		Nausea					

Date:01/04/02ISR Number: 3850017-0Report Type:Expedited (15-DaCompany Report #200124000GDDC
 Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Cognitive Disorder	Foreign	Docetaxel	PS		
INTRAVENOUS	169 MG	Q3W IV 85 DAY					
Initial or Prolonged		Confusional State	Other	Gabapentin	SS		ORAL
500 MG TID PO 2	WK						
		Constipation		Benzydamine	C		
		Diarrhoea		Fentanyl	C		
		Fall		Dalteparin Sodium	C		
		Pulmonary Embolism		Ranitidine	C		
		Ulna Fracture		Salbutamol			
				W/Ipratropium	C		
				Dexamethasone	C		
				Chloramphenicol	C		
				Lactulose	C		
				Ciprofloxacin	C		
				Ciprofloxacin	C		
				Ampicillin	C		
				Frusemide	C		
				Nystatin	C		
				Loperamide	C		
				Paracetamol	C		
				Co-Danthramer	C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Convulsion	Foreign	Gabapentin			
1200 MG (400		Educational Problem	Consumer	(Gabapentin)	PS		ORAL
MG, TID), PER		Fatigue	Other				
ORAL		Headache					
		Migraine		(Immunoglobulin			
		Oedema Mouth		Human Normal)	C		
		Pharyngeal Oedema					

Age:79 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Angina Unstable
Initial or Prolonged	Arterial Occlusive
Other	Disease
	Blood Pressure Abnormal
	Chest Discomfort

Freedom Of Information (FOI) Report

Joint Swelling

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
600 MG (TWICE DAILY), PER ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
			(Etodolac)	C		
			(Paracetamol, Hydrocodone Bitartrate)	C		
			(Atenolol)	C		
			(Psyllium Hydrophilic Mucilloid)	C		

Date:01/08/02 ISR Number: 3850130-8 Report Type:Expedited (15-DaCompany Report #034-0945-M0100012
 Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1800 MG, PER ORAL	Alanine Aminotransferase Increased	Foreign Literature	Gabapentin (Gabapentin)	PS		ORAL
		Aspartate	Health				
		Aminotransferase Abnormal	Professional	(Metamizole)	C		
		Blood Alkaline		(Tramadol)	C		
		Phosphatase Increased		(Morphine)	C		
		Eosinophilia		(Clonazepam)	C		
		Gamma-Glutamyltransferase Increased		(Omeprazole)	C		
		Hepatomegaly		(Fentanyl)	C		
		Hepatotoxicity		(Ciprofloxacin)	C		
		Jaundice		(Methylprednisolone)	C		
		Pruritus		Dexchlorpheniramine	C		
		Rash Erythematous					
		Toxic Skin Eruption					
		White Blood Cell Count Increased					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Rash Erythematous	Foreign	Neurontin			
Initial or Prolonged	Rash Pruritic	Health	(Gabapentin)	PS		ORAL
PER ORAL	Rash Vesicular	Professional	(Trimebutine)	C		
			(Buflomedil)	C		
			(Docusate Sodium)	C		
			(Acetylsalicylate			
			Lysine)	C		
			(Alprazolam)	C		
			(Domperidone)	C		
			(Sodium Bicarbonate)	C		
			(Mineral Oil			
			Emulsion)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/10/02ISR Number: 3851095-5Report Type:Expedited (15-DaCompany Report #033-0945-M0200002

Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged PER ORAL	Deep Vein Thrombosis Gouty Arthritis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other PER ORAL	Oedema Peripheral Pyrexia Synovitis	Professional	(Clopidogrel)	SS		ORAL

Date:01/10/02ISR Number: 3851103-1Report Type:Expedited (15-DaCompany Report #046-0945-M0100018

Age:47 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2400 MG Other (DAILY), PER ORAL	Blood Alkaline Phosphatase Increased Blood Potassium Increased Cardiac Failure	Foreign Consumer	Neurontin Tablets (Gabapentin)	PS		ORAL
40 MG (DAILY), PER ORAL	Chest Pain Depressed Level Of Consciousness		Furosemide	SS		ORAL
50 MG (DAILY), PER ORAL	Electroencephalogram Abnormal		Spirolactone	SS		ORAL
20 MG (DAILY), PER ORAL	Infected Skin Ulcer Liver Disorder Liver Function Test Abnormal		Enalapril Maleate	SS		ORAL
25 MG	Metabolic Disorder		Rofecoxib	SS		ORAL

(DAILY), PER	Renal Failure Acute			
ORAL	Skin Ulcer			
800 MG	Weight Increased	Metronidazole	SS	ORAL
(DAILY), PER				
ORAL		Flucloxacillin		
PER ORAL		Sodium	SS	ORAL
20 MG		Leflunomide	SS	ORAL
(DAILY), PER				
ORAL		Folic Acid, Calcium		
		Phosphate	C	
		Tocopheryl Acetate	C	
		Paracetamol	C	
		Prednisolone	C	
		Morphine	C	
		Cisapride	C	
		Cetirizine	C	
		Morphine Sulfate	C	
		Lansoprazole	C	
		Zolpidem Tartrate	C	
		Loperamide	C	
		Calcium Carbonate	C	
		Combivent		
		(Ipratropium		
		Bromide, Salbutamol		
		Sulfate)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/10/02ISR Number: 3851108-0Report Type:Expedited (15-DaCompany Report #044-0945-M0200001
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (100 Other MG, TID), PER ORAL		Cerebrovascular Accident Constipation Embolism Gastrointestinal Haemorrhage Vomiting	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Lansoprazole	C		

Date:01/10/02ISR Number: 3851408-4Report Type:Expedited (15-DaCompany Report #A200280
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose Sleep Attacks Suicidal Ideation	Health Professional	Atarax Tablets Neurontin Hydrocodone Fentanyl	PS SS C C		

Date:01/10/02ISR Number: 3851420-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200033
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 700 MG (DAILY), UNKNOWN		Constipation Decreased Activity Insomnia Social Avoidant Behaviour Vision Blurred	Consumer	Neurontin (Gabapentin)	PS		

Date:01/10/02ISR Number: 3851421-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200023
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chromaturia	Health Professional	Neurontin (Gabapentin)	PS		

Date:01/10/02ISR Number: 3851422-9Report Type:Expedited (15-DaCompany Report #001-0945-M0101475
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional	Neurontin (Gabapentin)	PS		
Life-Threatening		Suicidal Ideation		Hydroxyzine	SS		
		Toxicologic Test Abnormal		Hydrocodone	C		
				Fentanyl	C		

Date:01/11/02ISR Number: 3851832-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200026
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG, PER		Staphylococcal Infection					
ORAL		Vision Blurred					
		Visual Disturbance		(Paracetamol, Oxycodone Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Rofecoxib) C
 (Zyvox) C
 (Clonazepam) C

Date:01/11/02ISR Number: 3851857-4Report Type:Expedited (15-DaCompany Report #USA021029
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abdominal Pain		Dilaudid	PS		
Initial or Prolonged		Back Pain		Dilaudid	SS		
		Blood Glucose Increased		Dilaudid	SS		
		Bronchitis		Dilaudid	SS		
		Cholecystitis		Neurontin	SS		ORAL
300 MG OD PO							
		Cholelithiasis		Neurontin	SS		ORAL
300 MG BID PO							
		Colonic Polyp		Neurontin	SS		ORAL
300 MG TID PO							
		Cough		Neurontin	SS		ORAL
OTHER TID PO							
		Dyspnoea Exertional		Neurontin	SS		ORAL
300 MG TID PO							
		Ear Discomfort		Trazodone	SS		
		Fatigue		Diabeta	C		
		Foot Deformity		Glucophage	C		
		Fungal Infection		Spirocholactone	C		
		Fungal Rash		Zyrtec	C		
		Fungal Skin Infection		Vancenase	C		
		Glucose Urine Present		Axid	C		
		Glycosylated Haemoglobin		Glyburide	C		
		Increased		Insulin Humulin			
		Haematochezia		70/30	C		
		Haemorrhage		St. John'S Wort	C		
		Headache		Zithromax	C		
		Hepatic Steatosis		Zovia 1/35	C		
		Hepatomegaly		Guiatex	C		
		Nausea		Propoxyphene	C		
		Neck Pain		Darvon-N	C		
		Oedema Peripheral		Cephalexin	C		
		Ovarian Cyst Ruptured		Tylenol	C		
		Paranasal Sinus		Benadryl	C		
		Hypersecretion		Demerol	C		
		Proteinuria		Vicodin	C		
		Rales		Heparin	C		

Rhinitis
Sinusitis
Tinea Pedis
Umbilical Hernia
Urine Ketone Body Present
Vomiting
Weight Increased

Phenergan C
Cefazolin C
Compazine C
Vitamins C
Dovonex C
Ultravate C

Date:01/14/02ISR Number: 3852904-6Report Type:Expedited (15-DaCompany Report #055-0945-M0200002
Age:72 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 800 MG	Leg Amputation	Foreign	Gabapentin	PS		ORAL
Initial or Prolonged (DAILY), PER	Spinal Disorder	Consumer				

ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/15/02ISR Number: 3852867-3Report Type:Direct
 Age:49 YR Gender:Female I/FU:I

Company Report #CTU 159134

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Balance Disorder		Gabapentin			
		Dizziness		(Neurontin) 300mg	PS		
300MG 1 BID		Fatigue					

Date:01/15/02ISR Number: 3853220-9Report Type:Expedited (15-DaCompany Report #001-0945-M0101395
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Abnormal	Health	Neurontin			
Other		Drug Interaction	Professional	(Gabapentin)	PS		
		Hallucination		Alprazolam	SS		
		Oxygen Saturation		Oxygen	C		
		Increased					
		Paranoia					
		Thinking Abnormal					

Date:01/15/02ISR Number: 3853382-3Report Type:Expedited (15-DaCompany Report #002-0945-M0100131
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Foreign	Gabapentin			
Hospitalization -		Alanine Aminotransferase	Health	(Gabapentin)	PS		ORAL
Initial or Prolonged		Increased	Professional				
300 MG		Aspartate					
Other		Aminotransferase		(Hydromorphone)	SS		ORAL
(DAILY), PER		Increased					
ORAL		Blood Albumin Decreased		(Hydromorphone Sr)	SS		ORAL
		Blood Alkaline					
8 MG, PER							
ORAL							
18 MG (EVERY							
12 HOURS),							

Phosphatase Increased		
Blood Calcium Decreased	(Paracetamol)	C
Blood Phosphorus	(Magnesium	
Increased	Hydroxide)	C
Blood Pressure Systolic	(Glycerol)	C
Increased	(Bisacodyl)	C
Blood Urea Increased	(Sodium Phosphate	
Cervical Cord Compression	Dibasic, Sodium	
Clonic Convulsion	Phosphate)	C
Coma	(Sennoside A)	C
Depressed Level Of	(Docusate Calcium)	C
Consciousness	(Metoclopramide)	C
Eye Rolling		
Haematuria		
Heart Rate Increased		
Hypoventilation		
International Normalised		
Ratio Increased		
Neoplasm		
Respiratory Rate		
Decreased		
White Blood Cell Count		
Increased		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/16/02ISR Number: 3853618-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200044

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Accidental Overdose	Consumer	Neurontin (Gabapentin)	PS		
	Alopecia		(Diphenhydramine			
	Bipolar Disorder		Hydrochloride)	SS		ORAL
PER ORAL	Bladder Disorder		(Morphine)	SS		
	Condition Aggravated		(Unspecified Pain			
	Divorced		Medication)	SS		
	Dyspnoea		(Unspecified			
	Post Procedural		Insomnia Medication)	C		
	Complication					
	Pruritus					
	Stress					

Date:01/16/02ISR Number: 3853623-2Report Type:Expedited (15-DaCompany Report #A200681

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 10.00 MG	Diarrhoea	Consumer	Norvasc Tablets	PS		ORAL
Intervention to TOTAL:DAILY:0	Hypertension					
Prevent Permanent RAL	Hypoaesthesia					
Impairment/Damage 1200.00 MG	Neuropathy Peripheral		Neurontin	SS		ORAL
TOTAL:DAILY:0	Oedema Peripheral					
RAL	Paraesthesia					
40.00 MG	Therapeutic Response		Accupril	SS		ORAL
TOTAL:DAILY:0	Unexpected					
RAL			Coumadin	C		
			Lipitor	C		

Date:01/16/02ISR Number: 3855064-0Report Type:Expedited (15-DaCompany Report #DSA_21487_2002
Age: Gender:Not Specified/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG DAILY		Gastrointestinal Motility Disorder	Foreign Health	Diltiazem Neurontin	PS SS		ORAL
PO			Professional Other				

Date:01/17/02ISR Number: 3853848-6Report Type:Direct Company Report #CTU 159515
Age:83 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 300MG TWICE DAILY		Aspiration Gastric Haemorrhage Haemorrhagic Stroke Pneumonia		Neurontin 300mg Twice Daily	PS		

Date:01/17/02ISR Number: 3854159-5Report Type:Expedited (15-DaCompany Report #353-0945-M0100011
Age:73 YR Gender:Female I/FU:I

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Coordination Abnormal Fall Fibula Fracture	Foreign 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
PER ORAL		Gabapentin (Gabapentin)	PS		ORAL

Date:01/17/02ISR Number: 3854200-XReport Type:Expedited (15-DaCompany Report #A200833
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900.00 MG Required TOTAL:TID		Alanine Aminotransferase Increased	Foreign Health	Atarax Tablets Gabapentin	PS SS		
Intervention to Prevent Permanent 1000.00 MG Impairment/Damage TOTAL:PID		Ascites	Professional				
		Aspartate Aminotransferase		Ranitidine Valproic Acid	SS SS		
		Increased					
		Blood Alkaline Phosphatase Increased Gamma-Glutamyltransferase Increased Liver Disorder Lung Disorder Urinary Tract Infection		Glyceryl Trinitrate Noctran 10 Lactulose	C C C		

Date:01/17/02ISR Number: 3854817-2Report Type:Expedited (15-DaCompany Report #2002CG00039
Age:82 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 20 MG QD PO Initial or Prolonged 500 MG Q6H PO		Anti Factor Viii Antibody Positive	Foreign Health	Mopral 20 Mg Dafalgan 500	PS SS		ORAL ORAL
		Coagulation Factor Viii Level Decreased Coagulation Time	Professional Other	Prothrombinkomplex-K onzentrat Fragmine	SS SS		

TRANSDERMAL	5 MG QD TD	Prolonged		Discotrine	SS	
300 MG Q6H PO				Neurontin	SS	ORAL

Date:01/17/02ISR Number: 3855128-1Report Type:Expedited (15-DaCompany Report #N133866
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 75 MG PO	2 YR	Arthralgia Deep Vein Thrombosis	Health Professional	Plavix (Clopidogrel Sulfate)	PS		ORAL
4 U PO		Gout Middle Insomnia		Neurontin (Gabapentin)	SS		ORAL
		Oedema Peripheral Osteoarthritis Pyrexia Synovitis		...	C		

Date:01/17/02ISR Number: 3855160-8Report Type:Expedited (15-DaCompany Report #033-0945-M0200007
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG		Hypokalaemia Megacolon	Foreign Health Professional	Neurontin (Gabapentin)	PS		
(TID),							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/18/02ISR Number: 3855775-7Report Type:Expedited (15-DaCompany Report #A119156

Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	80.00 MG	Arrhythmia	Health	Ziprasidone Po	PS		ORAL
TOTAL:	BID:ORA	Cardiac Arrest	Professional				
L		Drooling					
	1200.00 MG	Drug Withdrawal		Gabapentin	SS		ORAL
TOTAL:	BID:ORA	Convulsions					
L		Electrocardiogram Qt					
		Prolonged		Risperdal	C		
		Haematemesis		Trazadone	C		
		Nausea		Ativan	C		
		Respiratory Arrest		Cogentin	C		
		Vomiting		Multivitamin	C		
				Haldol	C		
				Tylenol	C		
				Protonix	C		
				Restoril	C		

Date:01/18/02ISR Number: 3855988-4Report Type:Expedited (15-DaCompany Report #044-0945-M0200004

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG (THREE TIMES DAILY), PER ORAL	Cardiac Failure Congestive	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:01/18/02ISR Number: 3855999-9Report Type:Expedited (15-DaCompany Report #033-0945-M0200006

Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Alkaline Phosphatase Increased	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 Other MG, TID), PER		Condition Aggravated					
ORAL		Gamma-Glutamyltransferase					
		Increased Liver Function Test		(Hydroxyzine Hydrochloride)	SS		ORAL
PER ORAL		Abnormal Lung Disorder		(Ranitidine Hydrochloride)	SS		ORAL
PER ORAL		Urinary Tract Infection		(Valproic Acid, Valproate Sodium)	SS		ORAL
1000 MG (500 MG, BID) PER							
ORAL							
				(Glyceryl Trinitrate) 10 Mg, Transdermal	SS		
TRANSDERMAL	10 MG,						
TRANSDERMAL							
				(Acepromazine, Aceprometazine, Clorazepate Dipotassium)	SS		ORAL
PER ORAL				(Lactulose)	C		ORAL
(PRN), PER							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/22/02ISR Number: 3855483-2Report Type:Expedited (15-DaCompany Report #10842557
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthralgia		Videx Tabs	PS	Bristol-Myers Squibb Company	ORAL
-also from		Hypertriglyceridaemia					
August 1995		Osteonecrosis					
to May 1996.				Zerit	SS	Bristol-Myers Squibb Company	
				Sustiva	SS	Bristol-Myers Squibb Company	
				Neurontin	SS		
				Ziagen	SS		
				Bactrim Forte	SS		
				Lipur	C		

Date:01/22/02ISR Number: 3857541-5Report Type:Expedited (15-DaCompany Report #033-0945-M0100149
Age:71 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		
500 MG		Hyperprolactinaemia					
Other (DAILY)		Open Angle Glaucoma	Professional				
				Brimonidine	SS		
				(Veralipride)	SS		
				(Timolol)	C		
				(Latanoprost)	C		
				(Ursodeoxycholic Acid)	C		
				(Flavonoides)	C		
				(Alpha-Tocopherol)	C		
				(Silymarin)	C		

Date:01/22/02ISR Number: 3858038-9Report Type:Expedited (15-DaCompany Report #033-0945-M0100156
Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2400 MG	Clonic Convulsion Drug Tolerance Decreased	Foreign Health	Neurontin (Gabapentin)	PS		
(DAILY)		Fall	Professional				
	200 MG	Hyperaesthesia Hyponatraemia	Company Representative	Tegretol (Carbamazepine)	SS		
(DAILY)		Paresis					
		Sciatica		(Potassium Chloride)	C		
		Sialoadenitis		(Aciclovir)	C		
		Somnolence		(Ramipril)	C		
				(Digoxin)	C		
				(Amiodarone Hydrochloride)	C		
				(Allopurinol)	C		
				(Furosemide)	C		

Date:01/22/02ISR Number: 3858049-3Report Type:Expedited (15-DaCompany Report #033-0945-M0200008
Age:82 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Anti Factor Viii Antibody Positive

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Coagulation Factor Viii Level Decreased Coagulation Time	Report Source	Product	Role	Manufacturer	Route
1200 MG (300 MG, QID), PER ORAL		Prolonged	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
2000 MG (500 MG, QID), PER ORAL				(Paracetamol)	SS		ORAL
INTRAVENOUS MILLILITERS (ONCE), INTRAVENOUS	40			Kaskadil	SS		
SUBCUTANEOUS 0.2ML (DAILY), SUBCUTANEOUS	2500 IU/			Heparin-Fraction, Sodium Salt	SS		
20 MG (DAILY), PER ORAL				Omeprazole	SS		ORAL
TRANSDERMAL TRANSDERMAL	5 MG (DAILY),			(Glyceryl Trinitrate)	SS		

Date:01/22/02ISR Number: 3858271-6Report Type:Expedited (15-DaCompany Report #0018910
Age:82 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - TRANSDERMAL 5 MG Initial or Prolonged TRANSDERMAL	Anti Factor Viii Antibody	Health	Discotrine (5 Mg)	PS		
2000 MG ORAL 34 DAY SUBCUTANEOUS 0.2 ML SUBCUTANEOUS 42 DAY 20 MG ORAL 1200 MG ORAL	Coagulation Factor Viii Level Decreased Coagulation Time Prolonged	Professional	Paracetamol (500 Mg) Fragmine Mopral Neurontin Kaskadil	SS SS SS SS SS		ORAL ORAL ORAL
INTRAVENOUS	INTRAVENOUS					

Date:01/23/02ISR Number: 3856177-XReport Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-11665429
Age:44 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Disability Other	Cholelithiasis Colonic Polyp Condition Aggravated Hepatic Cirrhosis Hepatic Steatosis Ovarian Cyst Ruptured		Trazodone Hcl Tabs Rezulin Vasotec Hyoscyamine Amitriptyline Neurontin Dilaudid	PS SS SS SS SS SS SS	Apothecon	 ORAL

Date:01/23/02ISR Number: 3859490-5Report Type:Expedited (15-DaCompany Report #001-0719-M0100407
Age: Gender:Male I/FU:F

Outcome
Hospitalization -
Initial or Prolonged
Disability

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(Gabapentin) SS
Lithium (Lithium) C

Date:01/23/02ISR Number: 3860143-8Report Type:Expedited (15-DaCompany Report #033-0945-M0200006
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG, TID), PER ORAL		Alanine Aminotransferase Increased	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
		Ascites	Health				
		Aspartate	Professional				
		Aminotransferase Increased		(Hydroxyzine Hydrochloride)	SS		ORAL
PER ORAL		Blood Alkaline Phosphatase Increased		(Ranitidine Hydrochloride)	SS		ORAL
PER ORAL		Liver Disorder Liver Function Test		(Valproic Acid, Valproate Sodium)	SS		ORAL
1000 MG (500 MG, BID), PER ORAL		Abnormal					
		Lung Disorder					
		Prothrombin Level Abnormal		(Glyceryl Trinitrate)	SS		
TRANSDERMAL TRANSDERMAL	10 MG,	Urinary Tract Infection					
				(Acepromazine, Aceprometazine, Clorazepate Dipotassium)	SS		ORAL
10 MG (DAILY), PER							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL
 (PRN), PER Lactulose SS ORAL

ORAL
 (Heparin-Fraction, Sodium Salt) C

Date:01/23/02ISR Number: 3860170-0Report Type:Expedited (15-DaCompany Report #055-0945-M0200008
 Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	800 MG (BID)	Bone Neoplasm Malignant Prostate Cancer	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL

PER ORAL

Date:01/23/02ISR Number: 3860233-XReport Type:Direct Company Report #CTU 159883
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage	200MG 1 X DAY	Choreoathetosis Drug Hypersensitivity		Celebrex 200mg Searle/Pfizer	PS	Searle/Pfizer	ORAL
	300 MG 3 X DAY	Drug Interaction Dysstasia Medication Error Muscle Spasms Muscle Twitching Muscular Weakness		Neurontin 300mg Parke-Davis	SS	Parke-Davis	ORAL

Date:01/24/02ISR Number: 3858562-9Report Type:Direct Company Report #CTU 160033
 Age:61 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Condition Aggravated Erectile Dysfunction		Gabapentin (Neurontin)	PS		
400MG QD					Hctz	C		
					Novolin N	C		
					Alprostadil	C		
					Fluticasone Nasal And Oral Inhaler	C		
					Lisinopril	C		
					Serevent	C		
					Novolin R	C		

Date:01/24/02ISR Number: 3858917-2Report Type:Direct Company Report #CTU 160053
Age:78 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300MG PO TID	3 DAY	Confusional State Mental Status Changes		Gabapentin 300mg Po Tid	PS		ORAL
20MG Q H			Sedation		Elavil 20mg Q H	SS		
INTRATHECAL	20MG/DAY		Urinary Tract Infection		Morphine Sulfate 20mg/Day	SS		
INTRATHECAL					Asa	C		
					Claritin	C		
					Neurontin	C		
					Norvasc	C		
					Pravachol	C		
					Roxicodone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Senokot C
 Vioxx C
 Xanax C

Date:01/24/02ISR Number: 3860931-8Report Type:Expedited (15-DaCompany Report #055-0945-M0000020
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	1200 MG (600 MG, BID), PER	Blindness Unilateral	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:01/24/02ISR Number: 3861103-3Report Type:Expedited (15-DaCompany Report #044-0945-M0200007
 Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Confusional State Dysphasia	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
				(Oxycodone)	C		
				(Dexamethasone)	C		
				(Lansoprazole)	C		
				(Paroxetine Hydrochloride)	C		
				(Paraffin, Liquid, Milk Of Magnesia)	C		
				(Haloperidol)	C		
				(Fluconazole)	C		
				(Zopiclone)	C		
				(Paracetamol, Dextropropoxyphene Hydrochloride)	C		
				(Diltiazem)	C		

Date:01/25/02ISR Number: 3857851-1Report Type:Expedited (15-DaCompany Report #A0170813A
Age:17 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - YR	Convulsion		Wellbutrin	PS	Glaxo Wellcome	ORAL
Initial or Prolonged UNKNOWN	Euphoric Mood		Soma	SS		
	Overdose		Neurontin	SS		
UNKNOWN	Suicide Attempt		Effexor	C		

Date:01/25/02ISR Number: 3860053-6Report Type:Expedited (15-DaCompany Report #033-0945-M0100125
Age:87 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (300 Other MG, TID) PER	Balance Disorder Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
	Fall	Professional				
ORAL	Gait Disturbance					
	Hepatitis Hepatocellular Damage		(Paroxetine Hydrochloride) (Amiodarone Hydrochloride) (Meprobamate) (Valsartan)	SS C C C		

Freedom Of Information (FOI) Report

(Acetylsalicylate
Lysine) C
(Paroxetine) C
(Glyceryl
Trinitrate) C
(Furosemide) C
(Loprazolam) C
(Risperidone) C
(Macrogol, Sodium
Bicarbonate, Sodium
Chloride, Potassium
Chloride) C

Date:01/25/02ISR Number: 3860325-5Report Type:Expedited (15-DaCompany Report #PHRM2002FR00527
Age:63 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Alanine Aminotransferase Increased Ascites Aspartate Aminotransferase	Foreign Health Professional Other	Nitriderm Tts (Glyceryl Trinitrate) Trans-Therapeutic System			PS
TRANSDERMAL TRANSDERMAL	10 MG, QD, Increased					
PRN, ORAL	Blood Alkaline Phosphatase Increased Echography Abnormal		Atarax(Hydroxyzine Hydrochloride) Tablet	SS		ORAL
ORAL	Gamma-Glutamyltransferase Increased Lung Disorder		Azantac (Ranitidine Hydrochloride) Unknown	SS		ORAL
300 MG, TID, ORAL	Urinary Tract Infection		Neurontin (Gabapentin) Unknown	SS		ORAL
500 MG, BID,			Depakine Chrono (Valproate Sodium, Valproic Acid) Slow Release Tablet	SS		ORAL

ORAL

Noctran (Clorazepate
Dipotassium,
Aceprometazine,
Acepromazine) Tablet SS

ORAL

10 MG, QD,

ORAL

Lactulose
(Lactulose) Solution SS

ORAL

ORAL

Date:01/25/02ISR Number: 3860604-1Report Type:Expedited (15-DaCompany Report #001-0945-M0101278
Age:53 YR Gender:Male I/FU:F

Outcome	PT
Disability	Anorexia
Other	Asthenia
	Cervical Spine Flattening
	Cyst
	Depressed Level Of
	Consciousness
	Depression
	Disturbance In Attention
	Dizziness
	Dyspepsia

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ORAL

Dizziness

Drug Ineffective
Drug Toxicity
Eyelid Ptosis
Lethargy

(Tramadol
Hydrochloride) C
(Digoxin) C
(Lisinopril) C
(Trazodone) C
(Calcitonin) C
(Ergocalciferol,
Calcium Phosphate,
Calcium Sodium) C
(Paracetamol) C
(Vitamins Nos,
Minerals Nos) C

Date:01/28/02ISR Number: 3859768-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200033
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Constipation Drug Withdrawal Syndrome	Consumer	Neurontin (Gabapentin)	PS		
UNKNOWN	700 MG	Eye Pain					
(DAILY),		Impaired Driving Ability					
UNKNOWN		Vision Blurred					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/02ISR Number: 3859769-7Report Type:Expedited (15-DaCompany Report #001-0945-M0101421

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG (DAILY), PER ORAL		Eyelid Infection	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				(Amfebutamone Hydrochloride)	C		

Date:01/28/02ISR Number: 3859779-XReport Type:Expedited (15-DaCompany Report #001-0945-M0101185

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (300 MG, BID), PER ORAL		Back Pain Blood Pressure Systolic Increased Convulsion	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
UNKNOWN UNKNOWN	60 MG (TID),	Dizziness Headache Hypoaesthesia		(Propranolol Hydrochloride)	SS		
		Intracranial Haemangioma Migraine Neck Pain Pharyngolaryngeal Pain Somnolence Stupor Vision Blurred Visual Disturbance Visual Field Defect		(Estradiol)	C		

Date:01/28/02ISR Number: 3859787-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200102
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erectile Dysfunction	Health Professional	Neurontin (Gabapentin)	PS		ORAL
2400 MG							
(TID), PER							
ORAL				(Morniflumate)	C		

Date:01/28/02ISR Number: 3860675-2Report Type:Expedited (15-DaCompany Report #039-0945-M0100013
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Depression Sedation	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (TWICE							
DAILY), PER		Self Injurious Behaviour	Professional				
ORAL				(Atenolol)	C		

Date:01/28/02ISR Number: 3860696-XReport Type:Expedited (15-DaCompany Report #044-0945-M0200009
Age:73 YR Gender:Female I/FU:I

Outcome	PT	Report Source
Disability	Disorientation Psychomotor Hyperactivity	Foreign Health

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
900 MG (THREE TIMES DAILY) PER ORAL		Neurontin (Gabapentin)	PS		ORAL
		(Nifedipine)	C		
		(Insulin Human, Insulin Human Injection, Isophane)	C		
		(Furosemide, Amiloride Hydrochloride)	C		
		(Ranitidine)	C		
		(Theophylline)	C		

Date:01/28/02ISR Number: 3861234-8Report Type:Expedited (15-DaCompany Report #A201067
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Alanine Aminotransferase	Foreign	Atarax Tablets	PS		ORAL
Initial or Prolonged 900.00 MG		Increased	Health	Gabapentin	SS		ORAL
TOTAL:TID:ORA		Ascites	Professional				
L		Aspartate	Other				
ORAL		Aminotransferase		Ranitidine	SS		ORAL
1000.00 MG		Increased		Valproic Acid	SS		ORAL
TOTAL:BID:ORA		Blood Alkaline					
L		Phosphatase Increased					
		Echography Abnormal		Trinitrine	C		
		Gamma-Glutamyltransferase		Noctran	C		
		Increased		Lactulose	C		

Liver Disorder
 Lung Disorder
 Prothrombin Level
 Increased
 Urinary Tract Infection

Omeprazole C
 Enoxaparine C
 Lamotrigine C

Date:01/28/02ISR Number: 3861315-9Report Type:Expedited (15-DaCompany Report #055-0945-M0200009
 Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebral Infarction	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
Hospitalization - 400 MG Initial or Prolonged (DAILY), PER							ORAL

Date:01/28/02ISR Number: 3861316-0Report Type:Expedited (15-DaCompany Report #039-0945-M0200002
 Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (DAILY), PER		Dermatitis Atopic Rash Maculo-Papular	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Celecoxib	SS		ORAL
200 MG (DAILY), PER							ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/29/02ISR Number: 3860878-7Report Type:Expedited (15-DaCompany Report #B0133358A

Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	78 DAY	Alanine Aminotransferase		Azantac	PS	Glaxo Wellcome	ORAL
Initial or Prolonged	300MG Three	Increased		Gabapentin	SS		ORAL
times per day		Ascites					
500MG Twice		Aspartate		Valproic Acid	SS		ORAL
per day		Aminotransferase					
		Increased		Hydroxyzine			
		Blood Alkaline		Hydrochloride	SS		ORAL
		Phosphatase Increased		Nitroglycerin	SS	Glaxo Wellcome	
TRANSDERMAL	10MG Per day	Condition Aggravated		Noctran	SS		ORAL
UNKNOWN		Gamma-Glutamyltransferase		Lactulose	C		
		Increased					
		Liver Disorder					
		Lung Disorder					
		Prothrombin Time					
		Prolonged					
		Ultrasound Scan Abnormal					
		Urinary Tract Infection					

Date:01/29/02ISR Number: 3904257-2Report Type:Periodic Company Report #A123238

Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1000.00 MCG	Abdominal Pain	Consumer	Tikosyn Capsules	PS		ORAL
TOTAL, BID,		Constipation					
ORAL		Diarrhoea					
900.00 MG		Drug Ineffective		Neurontin	SS		ORAL
TOTAL, TID,							

ORAL

Methadone

SS

ORAL

ORAL

Coumadin
Tylenol
Altace
Coreg

C
C
C
C

Date:01/30/02ISR Number: 3862079-5Report Type:Expedited (15-DaCompany Report #033-0945-M0100125
Age:87 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG, TID), PER		Balance Disorder Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Cytolytic Hepatitis	Professional				
ORAL		Fall Gait Disturbance		(Clavulanate Potassium Amoxicillin Trihydrate) (Amiodarone Hydrochloride) (Meprobamate) (Valsartan) (Acetylsalicylate Lysine) (Paroxetine) (Glyceryl Trinitrate) (Furosemide)	C C C C C C		

Freedom Of Information (FOI) Report

(Loprazolam) C
 (Risperidone) C
 (Macrogol, Sodium
 Bicarbonate, Sodium
 Chloride, Potassium
 Chloride) C

Date:01/30/02ISR Number: 3862222-8Report Type:Expedited (15-DaCompany Report #A200378
 Age:50 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - INTRAVENOUS 40.00 MG Initial or Prolonged TOTAL; DAILY;	Bradycardia Confusional State	Foreign Health	Feldene Capsules	PS		
INTRAVENOUS 2200.00 MG TOTAL; ORAL	Drug Abuser Drug Interaction Drug Level Increased	Professional	Gabapentin	SS		ORAL
12.00 MG TOTAL; ORAL	Dysarthria Dysphagia		Tizanidine	SS		ORAL
	Hallucination Hypertension Hypotension Hypovolaemia		Seropram Voltaren Vioxx	C C C		

Date:01/30/02ISR Number: 3862989-9Report Type:Expedited (15-DaCompany Report #001-0719-M0100407
 Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability Other	Abdominal Mass Angina Pectoris Anhedonia Anxiety Atherosclerosis Back Pain Benign Prostatic Hyperplasia

Blood Cholesterol
Increased
Chest Pain
Condyloma Acuminatum
Coronary Artery Disease
Coronary Artery Occlusion
Diabetes Mellitus
Drug Interaction
Dyspepsia
Ear Pain
Emphysema
Food Poisoning
Gastritis
Headache
Heart Rate Abnormal
High Density Lipoprotein
Increased
Hypertension
Hypothyroidism
Insomnia
Intermittent Claudication
Low Density Lipoprotein
Increased
Lumbar Spinal Stenosis

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Initial or Prolonged 100 MG	Toxic Skin Eruption	Consumer	(Gabapentin)	PS	ORAL
(DAILY), PER					
ORAL			(Celecoxib)	SS	ORAL
200 MG					
(DAILY), PER					
ORAL					

Date:01/31/02ISR Number: 3862896-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200121
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Consumer	Neurontin			
		Balance Disorder		(Gabapentin)	PS		
		Convulsion		Sertraline			
100 MG DAILY		Decreased Appetite		Hydrochloride	SS		
		Depression		Clonazepam	SS		
		Emotional Disorder		Levetiracetam	C		
		Energy Increased		Gaba	C		
		Fatigue					
		Feeling Abnormal					
		Headache					
		Irritability					
		Libido Decreased					
		Palpitations					
		Sedation					
		Sleep Disorder					
		Weight Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/31/02ISR Number: 3862917-6Report Type:Expedited (15-DaCompany Report #001-0945-M0200098

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health	Neurontin			
		Medication Error	Professional	(Gabapentin)	PS		
				Fluoxetine			
				Hydrochloride	C		
				Zolpidem Tartrate	C		

Date:01/31/02ISR Number: 3862920-6Report Type:Expedited (15-DaCompany Report #A201293

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Anxiety	Consumer	Zoloft Tablets	PS		
100.00 MG							
Intervention to		Balance Disorder					
TOTAL:DAILY							
Prevent Permanent		Convulsion		Neurontin	SS		
Impairment/Damage		Decreased Appetite		Klonopin	SS		
		Depression		Keppra	C		
		Energy Increased		Gaba	C		
		Fatigue					
		Headache					
		Irritability					
		Libido Decreased					
		Malaise					
		Palpitations					
		Sedation					
		Sleep Disorder					
		Weight Decreased					

Date:01/31/02ISR Number: 3862931-0Report Type:Expedited (15-DaCompany Report #001-0945-M0101395

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Glucose Abnormal	Health	Neurontin			
		Drug Interaction	Professional	(Gabapentin)	PS		
400 MG							

(DAILY),
UNKNOWN
Hallucination
Hallucination, Visual
Paranoia

Alprazolam SS
Oxygen C
Insulin C

Date:01/31/02ISR Number: 3862935-8Report Type:Expedited (15-DaCompany Report #001-0945-M0101376
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaphylactic Reaction	Consumer	Neurontin			
900 MG		Irritable Bowel Syndrome	Health	(Gabapentin)	PS		ORAL

(DAILY), PER

ORAL

Flovent (Fluticasone
Propinonate) C
Serevent (Salmeterol
Xinafoate) C
Ventolin
(Salbutamol) C
Unspecified
Nebulizer C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/31/02ISR Number: 3862941-3Report Type:Expedited (15-DaCompany Report #001-0945-M0101181

Age:79 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG (BID, PER ORAL 25 MG (DAILY) (ONE DOSE), PER ORAL	Bone Pain Contusion Depressed Level Of Consciousness Fall	Health Professional	Neurontin (Gabapentin) Vioxx (Rofecoxib) (Glibenclamide)	PS SS SS		ORAL ORAL
	Hypoglycaemia Inflammation Loss Of Consciousness Medication Error Respiratory Rate Increased		(Methadone) Fosamax (Alendronate Sodium) Nasal Calcitonin (Calcitonin) Paracetamol Enalapril	C C C C C C		

Date:02/01/02ISR Number: 3861873-4Report Type:Expedited (15-DaCompany Report #WAES 0201USA02611

Age:44 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Disability Other	Abdominal Adhesions Ammonia Increased Atelectasis Blood Urea Increased Burns Second Degree Cholelithiasis Colonic Polyp Condition Aggravated Cough Creatinine Renal Clearance Increased Diabetes Mellitus Diabetic Neuropathy Duodenal Varices Fatigue Fungal Rash Gallbladder Disorder

Gastric Varices
Hepatic Cirrhosis
Hepatic Steatosis
Hepatitis
Hepatomegaly
Hypergammaglobulinaemia
Intertrigo
Irritable Bowel Syndrome
Neck Pain
Obesity
Ovarian Cyst Ruptured
Pharyngolaryngeal Pain
Platelet Count Decreased
Polycystic Ovaries
Portal Hypertension
Psoriasis
Purpura
Rales
Red Blood Cell Count
Decreased
Serum Ferritin Increased
Somnolence

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Varices Oesophageal Weight Decreased	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
204	DAY			Vasotec	PS	Merck & Co., Inc	ORAL
				Rezulin	SS		
113	DAY			Rezulin	SS		ORAL
1	DAY			Neurontin	SS		ORAL
				Elavil	SS		ORAL
				Dilaudid	SS		
				Hyoscyamine	SS		
1	DAY			Neurontin	SS		ORAL
28	DAY			Neurontin	SS		ORAL
12	DAY			Neurontin	SS		ORAL
22	DAY			Neurontin	SS		ORAL
				Trazodone	SS		
				Diabeta	C		
				Glucophage	C		
				Spironolactone	C		
				Zyrtec	C		
				Vancenase	C		
				Axid	C		
				Glyburide	C		
				Insulin, Biphasic			
				Isophane [Injection]	C		
				Hypericin	C		
				Zithromax	C		
				Zovia	C		
				Guiatex-La	C		
				Propoxyphene			
				Hydrochloride	C		
				Ultravate	C		
				Dovonex	C		
				Vitamins			
				(Unspecified)	C		
				Compazine	C		
				Cefazolin	C		
				Phenergan			
				(Promethazine			
				Hydrochloride)	C		

Heparin	C
Vicodin Tablets	C
Demerol	C
Benadryl	C
Tylenol	C
Darvon-N	C
Cephalexin	C

Date:02/01/02ISR Number: 3863935-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200126
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG,							
DAILY, PER							
ORAL				Citalopram Hydrobromide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/02ISR Number: 3863946-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200128

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder Cataract Diplopia Ivth Nerve Paralysis Paralysis Visual Field Defect	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG (400 MG, TID), PER ORAL				(Metformin Hydrochloride) (Glibenclamide) (Verapamil) (Lisinopril)	C C C C		

Date:02/01/02ISR Number: 3864017-8Report Type:Expedited (15-DaCompany Report #061-0945-M0200010

Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness Memory Impairment Muscle Twitching	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
DAILY , PER ORAL							

Date:02/01/02ISR Number: 3864444-9Report Type:Expedited (15-DaCompany Report #033-0945-M0200003

Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged PER ORAL		Rash Erythematous Rash Pruritic Toxic Skin Eruption		Neurontin (Gabapentin)	PS		ORAL
Other				Trimebutine Buflomedil Docusate Sodium Acetylsalicylate	C C C		

Lysine	C
Alprazolam	C
Domperidone	C
Sodium Bicarbonate	C
Mineral Oil Emulsion	C
Betamethasone	
Dipropionate	C
Calcipotriol	C

Date:02/01/02ISR Number: 3864447-4Report Type:Expedited (15-DaCompany Report #055-0945-M0200011
 Age:57 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG (TWICE DAILY), PER ORAL	Haemorrhoid Operation	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
			Osseine-Hydroxyapati te Complex Atorvastatin	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/01/02ISR Number: 3864882-4Report Type:Expedited (15-DaCompany Report #2002CG00039
Age:82 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 20 MG QD PO	Anti Factor Viii Antibody	Foreign	Mopral 20 Mg	PS		ORAL
Initial or Prolonged 500 MG Q6H PO	Positive	Health	Dafalgan 500	SS		ORAL
	Coagulation Factor Vii Level Decreased	Professional Other	Prothrombinkomplex-K onzentrat	SS		
	Coagulation Time Prolonged		Fragmine	SS		
TRANSDERMAL	5 MG QD TD		Discotrine	SS		
300 MG Q6H PO	Coagulopathy		Neurontin	SS		ORAL
	Depressed Level Of Consciousness Disorientation Haematoma Somnolence					

Date:02/01/02ISR Number: 3865057-5Report Type:Expedited (15-DaCompany Report #033-0945-M0200007
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2400 MG (TID)	Gastrointestinal Disorder Hypokalaemia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
PER ORAL	Megacolon	Professional	(Acetylsalicylate Lysine)	C		

Date:02/04/02ISR Number: 3863819-1Report Type:Direct Company Report #CTU 160624
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose 300MG TID PO	Diplopia		Gabapentin 300mg Tid Parke-Davis	PS	Parke Davis	ORAL

Ativan C
Trazodone C
Celexa C

Date:02/04/02ISR Number: 3865016-2Report Type:Expedited (15-DaCompany Report #033-0945-M0200012
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia	Foreign	Neurontin			
1200 MG		Cardiac Disorder	Consumer	(Gabapentin)	PS		
(TID),		Confusional State					
120 MG (60		Face Oedema		(Morphine Sulfate)	SS		
MG, BID)		Insomnia					
10 MG (DAILY)		Pallor		(Diazepam)	SS		

Date:02/05/02ISR Number: 3865862-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200146
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Anaphylactic Reaction	Health	Neurontin			
PER ORAL			Professional	(Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/05/02ISR Number: 3865981-3Report Type:Expedited (15-DaCompany Report #033-0945-M0200008
 Age:82 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (300 MG, QID), PER ORAL		Anti Factor Viii Antibody Positive	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Coagulation Factor Viii Level Decreased	Professional				
2000 MG (500 MG, QID), PER ORAL		Coagulation Time Prolonged		(Paracetamol)	SS		ORAL
		Depressed Level Of Consciousness		(Kaskadil)	SS		
INTRAVENOUS MILLILITERS (ONCE), INTRAVENOUS	40	Disorientation Haematoma Somnolence					
SUBCUTANEOUS (DAILY), SUBCUTANEOUS	2500 IU/0.2ML	Subdural Haematoma		(Heparin-Fraction, Sodium Salt)	SS		
20 MG(DAILY), PER ORAL				(Omeprazole)	SS		ORAL
TRANSDERMAL TRANSDERMAL	5 MG (DAILY),			(Glyceryl Trinitrate)	SS		

Date:02/05/02ISR Number: 3865983-7Report Type:Expedited (15-DaCompany Report #046-0945-M0200005
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction Loss Of Consciousness	Foreign Health Professional Company Representative	Neurontin (Gabapentin) (Ketobemidone Hydrochloride, Dimethyl-3, 3-Diphenyl-1-Methyla llylamine Hcl)	PS SS		

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
Hospitalization - Initial or Prolonged 1500 MG (QID)		Asthma Blood Glucose Fluctuation	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other PER ORAL		Hypoglycaemia Incontinence		 (Bupropion)	 SS		 ORAL
450 MG (TID) PER ORAL				(Advair) (Prednisone) (Diazepam)	C C C		

Date:02/07/02ISR Number: 3867257-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200142
Age: Gender:Female I/FU:I

Outcome	PT
Other	Ankle Fracture Anxiety Bronchitis Cognitive Disorder Confusional State

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
PER ORAL		Health Professional	Neurontin (Gabapentin)	PS		ORAL
1.5 MG, PER ORAL			Clonazepam	SS		ORAL
			Paroxetine Hydrochloride	C		

Date:02/07/02ISR Number: 3867270-XReport Type:Expedited (15-DaCompany Report #WAES 0201USA02611
 Age:44 YR Gender:Female I/FU:I

Outcome
 Hospitalization -
 Initial or Prolonged
 Disability
 Other

PT
 Abdominal Adhesions
 Abdominal Pain
 Alanine Aminotransferase
 Increased
 Angiopathy
 Aspartate
 Aminotransferase
 Increased
 Atelectasis
 Back Pain
 Blood Ketone Body Absent
 Bronchitis
 Burning Sensation
 Chest X-Ray Abnormal
 Cholelithiasis
 Colonic Polyp
 Cough
 Crepitations
 Depression
 Diabetes Mellitus
 Inadequate Control
 Difficulty In Walking
 Dyspnoea Exertional
 Fatigue
 Foot Deformity
 Fungal Infection
 Fungal Rash

Fungal Skin Infection
Gastrointestinal Disorder
Haemorrhoidal Haemorrhage
Headache
Hepatic Cirrhosis
Hepatic Steatosis
Hepatitis
Hepatomegaly
Hiatus Hernia
Hypergammaglobulinaemia
Hypertension
Injury
Irritable Bowel Syndrome
Jaundice
Nausea
Neck Pain
Neuralgia
Oedema Peripheral
Ovarian Cyst Ruptured

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG (TWICE DAILY), PER ORAL	Haemorrhoid Operation	Foreign Health Professional	Gabapentin (Gabapentin) (Osseine - Hydroxyapatite Complex) (Atorvastatin) (Levothyroxine Sodium)	PS C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/08/02ISR Number: 3867504-1Report Type:Expedited (15-DaCompany Report #002-0945-M0200013

Age:87 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Nephritis Interstitial	Foreign Literature	Gabapentin (Gabapentin)	PS		ORAL
Hospitalization - Initial or Prolonged (300 MG (100 MG, TID)), PER			Health Professional				
ORAL				Omeprazole	SS		ORAL
20 MG (DAILY), PER							
ORAL							

Date:02/08/02ISR Number: 3867508-9Report Type:Expedited (15-DaCompany Report #039-0945-M0200002

Age:81 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (DAILY), PER		Dermatitis Atopic Rash Macular	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Toxic Skin Eruption	Health Professional	Celecoxib	SS		ORAL
200 MG DAILY, PER							
ORAL							

Date:02/08/02ISR Number: 3868035-5Report Type:Expedited (15-DaCompany Report #001-0945-M0100970

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other (PER ORAL)		Atonic Urinary Bladder Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Convulsion		(Chinese Herbs)	C		

Depression
 Gastrointestinal Pain
 Hyperacusis
 Lyme Disease
 Muscular Weakness
 Suicidal Ideation
 Vertigo

(Levothyroxine
 Sodium) C

Date:02/11/02ISR Number: 3867410-2Report Type:Direct
 Age:4 YR Gender:Female I/FU:I

Company Report #CTU 161390

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300MG BID		Hallucination		Gabapentin 300mg Parke-Davis	PS	Parke-Davis	ORAL
ORAL				Acetaminophen	C		
				Morphine	C		
				Ivig	C		

Date:02/11/02ISR Number: 3868577-2Report Type:Expedited (15-DaCompany Report #020-0945-M0200001
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (TID), PER		Coma Hepatic	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
ORAL				"Levatec" (Insulin)	C C		

Freedom Of Information (FOI) Report

(Heparin-Fraction,
Sodium Salt) C
(Tocopherol) C
(Phytomenadione) C

Date:02/11/02ISR Number: 3868903-4Report Type:Expedited (15-DaCompany Report #A200833
Age:63 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL	Alanine Aminotransferase	Foreign	Atarax Tablets	PS		ORAL
Initial or Prolonged 900.00 MG	Increased	Health	Gabapentin	SS		ORAL
Required TOTAL TID;	Ascites	Professional				
Intervention to ORAL	Aspartate	Other				
Prevent Permanent ORAL	Aminotransferase		Ranitidine	SS		ORAL
Impairment/Damage 1000.00 MG	Increased		Valproic Acid	SS		ORAL
TOTAL: BID:	Blood Alkaline					
ORAL	Phosphatase Increased					
	Gamma-Glutamyltransferase Increased		Glyceryl Trinitrate	C		
	Hepatic Function Abnormal		Noctran 10	C		
	Liver Disorder		Lactulose	C		
	Lung Disorder		Omeprazole	C		
	Prothrombin Level Decreased		Enoxaparine Sodique	C		
	Urinary Tract Infection		Lamotrigine	C		
			Ceftriazone	C		
			Spiramycine	C		

Date:02/11/02ISR Number: 3868911-3Report Type:Expedited (15-DaCompany Report #044-0945-M0200019
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1500 MG	Liver Function Test Abnormal	Foreign Health	Neurontin(Gabapentin)	PS		ORAL

(DAILY), PER
 ORAL
 Date:02/11/02ISR Number: 3868927-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200160
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Discomfort	Health	Neurontin			
Other		Abdominal Distension	Professional	(Gabapentin)	PS		ORAL
2800 MG		Abnormal Faeces					
(THREE TIMES		Angioneurotic Oedema					
DAILY) PER		Benign Gastric Neoplasm					
ORAL		Change Of Bowel Habit		Hydrochlorothiazide	C		
		Constipation		Diltiazem			
		Defaecation Urgency		Hydrochloride	C		
		Dyskinesia		Doxazosin Mesilate	C		
		Facial Palsy		Atorvastatin	C		
		Gait Disturbance		Rabeprazole Sodium	C		
		Urticaria		Domperidone	C		
		Weight Decreased		Clonazepam	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/11/02ISR Number: 3868999-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200169

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue	Consumer	Neurontin			
		Road Traffic Accident		(Gabapentin)	PS		
		Vision Blurred					
3600 MG							
(DAILY),							
UNKNOWN							

Date:02/11/02ISR Number: 3869377-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200142

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ankle Fracture	Health	Neurontin			
		Anxiety	Professional	(Gabapentin)	PS		ORAL
PER ORAL		Bronchitis		(Clonazepam)	SS		ORAL
1.5 MG, PER		Cognitive Disorder					
ORAL		Confusional State		Paroxetine			
		Costochondritis		Hydrochloride	C		
		Depression					
		Ear Infection					
		Gastritis					
		Laryngitis					
		Sinusitis					

Date:02/11/02ISR Number: 3869623-2Report Type:Expedited (15-DaCompany Report #046-0945-M0200005

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Foreign	Neurontin			
		Loss Of Consciousness	Health	(Gabapentin)	PS		
			Professional	(Ketobemidone			
			Company	Hydrochloride,			
			Representative	Dimethyl-3,			

3-Diphenyl-1-Methylam
llylamine Hcl SS

Date:02/11/02ISR Number: 3869625-6Report Type:Expedited (15-DaCompany Report #045-0945-M0100010
Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Foreign	Gabapentin			
Hospitalization -		Dizziness	Health	(Gabapentin)	PS		ORAL
300 MG							
Initial or Prolonged		Emphysema	Professional				
DAILY, PER							
Other		Headache					
ORAL							
		Malaise		Theophylline	C		
				Prednisolone	C		
				Budesonide	C		
				Salmeterol	C		
				Acetylcysteine	C		
				Ipratropium	C		
				Montelukast	C		
				Salbutamol	C		
				Alprazolam	C		
				Sertraline	C		
				Mianserin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/13/02ISR Number: 3869484-1Report Type:Periodic
Age:61 YR Gender:Male I/FU:I

Company Report #2013152

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional Company Representative	Oxycontin Cr Tablets, 40 Mg (Oxycodone Hydrochloride)	PS		ORAL
PO				Neurontin (Gabapentin)	SS		
				Phenergan (Promethazine)	SS		
				Flexeril (Cyclobenzaprine)	SS		
				Vancomycin	C		

Date:02/13/02ISR Number: 3869546-9Report Type:Periodic
Age:37 YR Gender:Female I/FU:I

Company Report #2013265

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		
				Alprazolam	SS		
				Promethazine	SS		
				Gabapentin	SS		
				Meperidine	SS		

Date:02/13/02ISR Number: 3870436-6Report Type:Expedited (15-DaCompany Report #NSADSS2002004183
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agranulocytosis Mental Disorder Neutropenia	Consumer Health Professional	Risperdal (2 Mg Tablet) (Risperidone)	PS		ORAL

2 MG, 1 IN 1

NIGHT (S),

ORAL

Neurontin
(Gabapentin)

SS

ORAL

ORAL

Depakote (Valproate
Semisodium)

C

Date:02/13/02ISR Number: 3870586-4Report Type:Periodic
Age:46 YR Gender:Male I/FU:I

Company Report #2012646

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		
				Alprazolam	SS		
				Diphenhydramine	SS		
				Nicotine	SS		
				Olanzapine	SS		
				Barbiturate	SS		
				Benzodiazepine	SS		
				Salicylate	SS		
				Valproic Acid	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Gabapentin	SS
Butalbital	SS
Carisoprodol	C
Neurontin	
(Gabapentin)	C
Depakote (Divalproex Sodium)	C
Zyprexa	C
Viagra	C
Risperdal	C
Nystatin Cream Usp	C
Diphenhydramine	C

Date:02/13/02ISR Number: 3870588-8Report Type:Periodic
Age:37 YR Gender:Female I/FU:I

Company Report #2012633

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Health Professional Other	Oxycontin Cr Tablets, 40 Mg (Oxycodone Hydrochloride)	PS		ORAL
PO				Carisoprodol	SS		
				Ranitidine	SS		
				Neurontin (Gabapentin)	SS		
				Propoxyphene	SS		
				Meprobamate	SS		
				Valproic Acid	SS		
				Roxicodone (Oxycodone)	SS		
				Caffeine Anhydride	SS		
				Hydrocodone/Apap	C		
				Prozac (Fluoxetine)	C		
				Depakote (Divalproex Sodium)	C		
				Skelaxin	C		
				Zyprexa	C		
				Remeron (Mirtazapine)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional Other	Oxycodone Hydrochloride Ethanol Diazepam Mirtazapine Fluoxetine Gabapentin Oxazepam Temazepam	PS SS SS SS SS SS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/13/02ISR Number: 3870615-8Report Type:Periodic
 Age:43 YR Gender:Female I/FU:I

Company Report #2013549

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Health Professional Other	Oxycodone Hydrochloride Sertraline Hydrochloride Gabapentin Clonazepam Acetaminophen Propoxyphene Hcl	PS SS SS SS SS SS		

Date:02/13/02ISR Number: 3870620-1Report Type:Periodic
 Age:42 YR Gender:Male I/FU:I

Company Report #2012272

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		ORAL
PO				Diazepam Oxazepam Temazepam Lorazepam Cannabinoids Diphenhydramine Hcl Gabapentin	SS SS SS SS SS SS SS		

Date:02/13/02ISR Number: 3870622-5Report Type:Periodic
 Age:44 YR Gender:Female I/FU:I

Company Report #2012275

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride Alprazolam Diphenhydramine Hcl Gabapentin Propoxyphene Hcl	PS SS SS SS SS		

Temazepam	SS
Acetaminophen	SS
A-Hydroxyalprazolam	SS
Metoprolol	SS
Oxazepam	SS

Date:02/13/02ISR Number: 3870699-7Report Type:Expedited (15-DaCompany Report #0018910
Age:82 YR Gender:Male I/FU:I

Outcome PT
Hospitalization - Anti Factor Viii Antibody
Initial or Prolonged Positive
Coagulation Factor Viii
Level Decreased
Coagulation Time
Prolonged
Convulsion
Depressed Level Of
Consciousness
Disorientation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Haematoma Somnolence Subdural Haematoma	Report Source	Product	Role	Manufacturer	Route
TRANSDERMAL	5 MG		Foreign	Discotrine (5 Mg)	PS		
TRANSDERMAL			Health				
2000 MG ORAL	34 DAY		Professional	Paracetamol (500 Mg)	SS		ORAL
SUBCUTANEOUS	0.2 ML			Fragmine	SS		
SUBCUTANEOUS	42 DAY			Mopral	SS		ORAL
20 MG ORAL				Neurontin	SS		ORAL
1200 MG ORAL				Kaskadil	SS		
INTRAVENOUS	INTRAVENOUS						

Date:02/13/02ISR Number: 3871032-7Report Type:Periodic Company Report #2012365
 Age:52 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 20-553)	PS		
				Cocaine	SS		
				Acetaminophen	SS		
				Promethazine	SS		
				Dextromethorphan	SS		
				Lidocaine	SS		
				Carbinoxamine	SS		
				Ephedrine	SS		
				Pseudoephedrine	SS		
				Fluoxetine	SS		
				Diazepam	SS		
				Clonazepam	SS		
				Gabapentin	SS		
				Nicotine	SS		

Date:02/13/02ISR Number: 3871065-0Report Type:Periodic
Age:41 YR Gender:Male I/FU:I

Company Report #2014946

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Interaction Intentional Misuse	Health Professional Other	Oxycontin Cr Tablets, 40 Mg (Oxycodone Hydrochloride)	PS		ORAL
MG PO				Hydrocodone Bitartrate	SS		
				Caffeine Anhydride	SS		
				Diphenhydramine Hcl	SS		
				Codeine	SS		
				Gabapentin	SS		
				Acetaminophen	SS		
				Cocaine	SS		
				Alprazolam	SS		

Date:02/13/02ISR Number: 3871124-2Report Type:Periodic
Age:31 YR Gender:Male I/FU:I

Company Report #2014152

Outcome	PT
Death	Abdominal Pain 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
MG PO		Health Professional Other	Oxycontin Cr Tablets, 10 Mg (Oxycodone Hydrochloride)	PS		ORAL
			Clonazepam Neurontin (Gabapentin)	SS		
			Clonidine Hcl Combivent (Ipratropium & Albuterol)	C		
			Lithium	C		

Date:02/13/02ISR Number: 3872412-6Report Type:Periodic Company Report #2014930
 Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		
				Gabapentin	SS		
				Mirtazapine	SS		
				Cocaine	SS		
				Nicotine	SS		
				Caffeine Anhydride	SS		

Date:02/13/02ISR Number: 3873193-2Report Type:Periodic Company Report #2014950
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		
				Diazepam	SS		

Gabapentin SS
Amitriptyline SS
Paroxetine Hcl SS
Caffeine Anhydride SS

Date:02/13/02ISR Number: 3876133-5Report Type:Periodic Company Report #2013156
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		
				Cocaine	SS		
				Cyclobenzaprine	SS		
				Gabapentin	SS		
				Ticlopidine	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/13/02ISR Number: 3876136-0Report Type:Periodic
Age:45 YR Gender:Male I/FU:I

Company Report #2013163

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) Ethanol (Alcohol, Anhydrous) Cocaine Gabapentin Cyclobenzaprine Hcl	PS SS SS SS SS		

Date:02/13/02ISR Number: 3876139-6Report Type:Periodic
Age:33 YR Gender:Male I/FU:I

Company Report #2013138

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	40 MG BID; PO	Accidental Overdose	Health Professional Other	Oxycontin Cr Tablets Neurontin (Gabapentin) Diphenhydramine Hcl Prozac (Fluoxetine)	PS SS SS SS		ORAL

Date:02/13/02ISR Number: 3876251-1Report Type:Periodic
Age:42 YR Gender:Female I/FU:I

Company Report #2012528

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) Paxil (Paroxetine) Meprobamate Carisoprodol Neurontin (Gabapentin)	PS SS SS SS SS		

Date:02/13/02ISR Number: 3876313-9Report Type:Periodic
Age:51 YR Gender:Female I/FU:I

Company Report #2012128

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion	Health Professional	Oxycontin Cr Tablets, 10 Mg (Oxycodone Hydrochloride)	PS		ORAL
MG; PO				Methadone Gabapentin	SS SS		

Date:02/13/02ISR Number: 3877725-XReport Type:Periodic
Age:46 YR Gender:Male I/FU:I

Company Report #2013077

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) Neurontin	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

500 MG QD; PO	(Gadapentin)	SS	
100 MG 1 QAM;	Levaquin		
PO	(Levofloxacin)	SS	ORAL
240 MG QD; PO	Zoloft (Sertraline)	SS	ORAL
20 MG QD; PO	Verapamil	SS	ORAL
QD; PO	Prilosec		
TRANSDERMAL 100 MG; TD	(Omeprazole)	SS	ORAL
10 MG QD; PO	Claritin-D		
	(Loratadine/Pseudoephedrine)	SS	ORAL
	Duragesic (Fentanyl)		
	Patch	SS	
	Potassium Chloride	SS	ORAL
	Alprazolam	SS	
	Propoxyphene	SS	
	Clidinium		
	W/Chlordiazepoxide	SS	

Date:02/13/02ISR Number: 3877752-2Report Type:Periodic
 Age:45 YR Gender:Male I/FU:I

Company Report #2013035

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)	PS		ORAL
PO				Neurontin (Gabapentin)	SS		ORAL
PO				Temazepam	SS		ORAL
PO				Secrogod	C		
				Buspar (Buspirone)	C		
				Promethazine	C		

Date:02/13/02ISR Number: 3879049-3Report Type:Periodic Company Report #2014479
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Oxycontin Cr Tablets, 40 Mg (Oxycodone Hydrochloride)	PS		ORAL
MG Q12H	PO			Diphenhydramine Hcl Promethazine Hcl Effexor (Venlafaxine) Gabapentin Risperidone Clonazepam	SS SS SS SS SS		ORAL
MG	PO						

Date:02/14/02ISR Number: 3870230-6Report Type:Expedited (15-DaCompany Report #049-0945-M0100097
Age:79 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other 600 MG		Atrioventricular Block Bradyarrhythmia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER							
ORAL				Atehexal (Atenolol)	C		

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Freedom Of Information (FOI) Report

Lorzaar (Losartan Potassium) (Acetylsalicylic Acid)	C
Bezafibrat (Bezafibrate)	C
Humalog Mix (Insulin Lispro)	C
Ossofortin (Colecalciferol, Calcium Gluconate, Calcium Phosphate)	C
Presomen (Estrogens Conjugated) (Magnesium)	C

Date:02/14/02ISR Number: 3870231-8Report Type:Expedited (15-DaCompany Report #046-0945-M0200005
Age: Gender:Unknown I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dialysis Drug Interaction Loss Of Consciousness Medication Error Somnolence	Foreign Health Professional Company Representative	Neurontin (Gabapentin) (Ketobemidone Hydrochloride, Dimethyl-3, 3-Diphenyl -1- Methylallylamine	PS		SS
INTRAVENOUS	7.5 MG,						
INTRAVENOUS ;							
2.5 MG,							
SUBCUTANEOUS							

Date:02/14/02ISR Number: 3870232-XReport Type:Expedited (15-DaCompany Report #044-0945-M0200022
Age:39 YR Gender:Male I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Circulatory Collapse	Foreign	Neurontin			

600 MG (200 Ecchymosis Health (Gabapentin) PS ORAL
 MG, TID), PER Professional
 ORAL (Tramadol) C

Date:02/14/02ISR Number: 3870446-9Report Type:Expedited (15-DaCompany Report #055-0945-M0100016
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign	Gabapentin			
Other		Drug Toxicity	Health	(Gabapentin)	PS		ORAL
800 MG (BID),		Photophobia	Professional				
PER ORAL							

Date:02/14/02ISR Number: 3870447-0Report Type:Expedited (15-DaCompany Report #033-0945-M0200012
 Age:56 YR Gender:Male I/FU:F

Outcome	PT
Death	Asthenia
	Back Pain
	Cardiac Disorder
	Confusional State

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Face Oedema Insomnia Osteoarthritis					
1200 MG (400 MG, TID)			Foreign Consumer	Neurontin (Gabapentin)	PS		
120 MG (60 MG, BID)		Skin Discolouration	Health Professional	(Morphine Sulfate)	SS		
10 MG (DAILY)				(Diazepam)	SS		
				(Tetracosactide)	C		
				(Tetrazepam)	C		
				(Floctafenine)	C		

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
Required Intervention to Prevent Permanent Impairment/Damage		Blood Testosterone Decreased Hyponatraemia	Health Professional	Zoloft Tablets	PS		
				Neurontin	SS		
				Paxil	SS		
				Wellbutrin	SS		
				Ambien	C		
				Albuterol Inhaler	C		
				Atenolol	C		
				Folic Acid	C		
				Zocor	C		
				Hydrochlorothiazide	C		

Date:02/14/02ISR Number: 3870481-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200170
Age:85 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG		Blood Glucose Increased Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Other (DAILY), PER	Drug Toxicity				
ORAL	Lethargy				
PER ORAL	Mental Status Changes		Phenytoin	SS	ORAL
	Mydriasis		Sulfamethoxazole,		
	Orthostatic Hypotension		Trimethoprim	SS	

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		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/15/02ISR Number: 3870895-9Report Type:Expedited (15-DaCompany Report #A0356185A

Age:5 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 1 TABLET, 3 Initial or Prolonged TIMES PER Other DAY, GASTR	Enterobacter Infection	Health	Calcium Carbonate	PS		ORAL
1 TABLET THREE TIMES PER DAY, GASTR	Haematemesis	Professional				
2 YR	Implant Site Reaction					
GASTROSTOMY TUBE	Inappropriate Antidiuretic Hormone Secretion		Calcium Carbonate Tablet 1 Tablet (Calcium Carbonate)	SS		ORAL
GASTROSTOMY TUBE	Medical Device Complication					
GASTROSTOMY TUBE	Pancreatitis					
GASTROSTOMY TUBE	Urinary Tract Infection		Gabapentin (Formulation Unknown)	SS		ORAL
GASTROSTOMY TUBE			Oxcarbazepine (Formulation Unknown)	SS		ORAL
GASTROSTOMY TUBE			Cefuroxime Sodium (Formulation Unknown) (Cefuroxime Sodium)	SS		ORAL
200 MG, THREE TIMES PER			Topiramate (Formualtion Unknown) (Topiramate)	SS		ORAL

DAY, GASTR

Modafinil C
Ranitidine
Hydrochloride C
Metoclopramide Hcl C
Polyvitamin With
Iron C
Loratadine C
Ibuprofen C
Docusate Sodium C

Date:02/19/02ISR Number: 3871599-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200187
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Chest Discomfort Cough	Consumer	Neurontin Tablets (Gabapentin)	PS		ORAL
2400 MG (QID), PER		Depression					
ORAL		Dyspnoea					
25 MG (DAILY), PER		Fluid Retention Throat Tightness		(Sertraline Hydrochloride)	SS		ORAL
ORAL		Weight Increased					
				(Paracetamol, Hydrocodone Bitartrate) (Levothyroxine Sodium)	C C		

Calcium Carbonate)	C
(Pancreatin)	C
Thyronajod	
(Levothyroxine	
Sodium)	C
Migraine Medication	C
Vioxx (Rofecoxib)	C
Karil "Novartis"	C
Zantac (Ranitidine	
Hydrochloride)	C
Ergenyl "Labaz	
(Valproate Sodium)	C
(Diane / Androcur	
10)	C
Laxoberal (Sodium	
Picosulfate)	C
Valproic Acid	
(Valproic Acid)	C
(Ass 100)	C

Date:02/19/02ISR Number: 3872721-0Report Type:Expedited (15-DaCompany Report #PHRM2001FR02891
Age:37 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

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG,		Blood Fibrinogen Decreased C-Reactive Protein Increased	Foreign Health Professional Other	Tegretol Lp (Carbamazepine) Extended Release Tablet	PS		ORAL
ONCE/SINGLE,		Drug Hypersensitivity					
ORAL		Ecchymosis					
1 DF,		Haemoglobin Decreased Medication Error		Alepsal (Belladonna Extract, Caffeine)	SS		ORAL
ONCE/SINGLE,		Purpura					
ORAL		Pyrexia					
1 DF,		Rash Skin Lesion		Neurontin (Gabapentin) Tablet	SS		ORAL
ONCE/SINGLE,							
ORAL							
1 DF,				Lepticur (Tropatepine Hydrochloride) Tablet	SS		ORAL
ONCE/SINGLE,							
ORAL							
1				Urbanyl (Clobazam)	SS		ORAL
DF/ONCE/SINGL							
E, ORAL							
1 DF,				Tercian (Cyamemazine) Tablet	SS		ORAL
ONCE/SINGLE,							

ORAL

Date:02/19/02ISR Number: 3872925-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200198
Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 5100 MG, PER Initial or Prolonged ORAL	Medication Error	Health Professional	Neurontin (Sertraline Hydrochloride)	PS C		ORAL

Date:02/20/02ISR Number: 3873001-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200176
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 1600 MG (DAILY), UNKNOWN	Arachnoiditis Back Pain Hypoaesthesia Joint Dislocation	Consumer	Neurontin (Gabapentin)	PS		
			Unspecified Medications	C		

Date:02/20/02ISR Number: 3873603-0Report Type:Expedited (15-DaCompany Report #370-0945-M0200001
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Other	Anuria Cardiac Disorder Muscular Weakness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pulmonary Embolism Renal Impairment	Report Source	Product	Role	Manufacturer	Route
(DAILY), PER			Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
ORAL				(Digoxin)	C		
				(Pentostatin)	C		
				(Warfarin)	C		
				(Diclofenac)	C		
				(Unspecified Diuretics)	C		
				(Ramipril)	C		

Date:02/20/02ISR Number: 3873847-8Report Type:Expedited (15-DaCompany Report #041-0945-M0200003
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2200 MG		Bradycardia Confusional State	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER		Drug Abuser					
ORAL		Drug Interaction					
INTRAMUSCULAR	40 MG ,	Drug Level Above Therapeutic		(Piroxicam)	SS		
INTRAMUSCULAR							
12 MG, PER		Dysarthria Dysphagia		(Tizanidine Hydrochloride)	SS		ORAL
ORAL		Hallucination					
PER ORAL		Hypertension Hypotension		(Citalopram Hydrobromide)	SS		ORAL
		Hypovolaemia		(Diclofenac Sodium) (Rofecoxib)	C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600MG QID		Back Pain Crystal Urine Present Muscle Strain		Neurontin 300mg Parkedavis	PS	Parkedavis	ORAL
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG/BID + 400 MG/QHS		Haematemesis Hyponatraemia Inappropriate Antidiuretic Hormone Secretion Pancreatitis Procedural Complication Vomiting	Consumer	Trileptal (Oxcarbazepine)	PS		
INTRAGASTRIC	730 DAY			Neurontin (Gabapentin) Topamax (Topiramate) Tums (Calcium Carbonate) Provigil (Modafinil) Zantac Solutions For Parenteral Nutrition Reglan (Metoclopramide)	SS SS SS C C C C		

Date:02/22/02ISR Number: 3875318-1Report Type:Expedited (15-DaCompany Report #001-0073-M0200076
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Drug Level Decreased	Consumer	Dilantin (Phenytoin Sodium)	PS		ORAL
300 MG (TID), PER ORAL		Drug Level Increased					
900 (TID), PER ORAL		Fluid Retention Grand Mal Convulsion		Gabapentin	SS		ORAL
		Hypersomnia Nasopharyngitis Oedema Peripheral Paraesthesia Somnolence					

Date:02/22/02ISR Number: 3875659-8Report Type:Expedited (15-DaCompany Report #002-0945-M0200013
Age:87 YR Gender:Male I/FU:F

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
300 MG (100 MG, TID), PER ORAL		Alanine Aminotransferase Increased	Foreign Literature	Neurontin (Gabapentin)	PS		ORAL
20 MG (DAILY), PER ORAL		Aspartate Aminotransferase Increased	Health Professional	(Omeprazole)	SS		ORAL
0.125 MG (DAILY), PER ORAL		Asthenia Blood Lactate Dehydrogenase Increased		(Digoxin)	SS		ORAL
4 MG (DAILY), PER ORAL		Blood Phosphorus Increased Confusional State Dehydration		(Warfarin Sodium)	SS		ORAL
		Drug Interaction		(Diltiazem)	C		
		Dyspnoea Exertional		(Furosemide)	C		
		Influenza Like Illness		(Venlafaxine Hydrochloride)	C		
		International Normalised Ratio Increased		(Ergocalciferol, Ascorbid Acid, Folic Acid, Thiamine Hydrochloride, Retinol, Riboflavin, (Tamsulosin)	C C C C		
		Nephritis Interstitial		(Oxazepam)	C		
		Oral Intake Reduced		(Latanoprost)	C		
		Renal Failure		(Dorzolamide Hcl/Timolol Maleate)	C C		
				(Insulin Human Injection, Isophane)	C C		
				(Calcium Carbonate)	C		
				(Ferrous Sulfate)	C		

Date:02/25/02ISR Number: 3874255-6Report Type:Expedited (15-DaCompany Report #061-0945-M0200032
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence Panic Disorder	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
3000 (DAILY),			Professional				
PER ORAL			Company Representative				

Date:02/25/02ISR Number: 3874297-0Report Type:Expedited (15-DaCompany Report #032-0945-M0100012
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness Fatigue	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300		Nausea	Professional				
MG, TID), PER		Post Procedural	Company				
ORAL		Complication	Representative	(Calcium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/25/02ISR Number: 3874301-XReport Type:Expedited (15-DaCompany Report #033-0945-M0200012

Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	800 MG (400 MG, BID), PER ORAL	Asthenia Coma	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
		Confusional State	Health				
		Face Oedema	Professional				
	60 MG (30 MG, BID), PER ORAL	Insomnia		(Morphine Sulfate)	SS		ORAL
		Skin Discolouration					
	5 MG (DAILY), PER ORAL			(Diazepam)	SS		ORAL
				(Tetracosactide)	C		
				(Tetrazepam)	C		
				(Floctafenine)	C		

Date:02/25/02ISR Number: 3874338-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200218

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1200 MG (BID)	Anaemia Anorgasmia	Health Professional	Neurontin (Gabapentin)	PS		
		Hypothyroidism Parkinsonism Tardive Dyskinesia Urinary Incontinence		(Lithium)	C		

Date:02/25/02ISR Number: 3874526-3Report Type:Expedited (15-DaCompany Report #049-0945-M0200019

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1000 MG		Coma	Foreign	Neurontin	PS		ORAL
Initial or Prolonged (DAILY), PER		Neutropenia	Health				
Other ORAL			Professional				
				Lorazepam	C		
				Acetylcysteine	C		

Date:02/25/02ISR Number: 3874550-0Report Type:Expedited (15-DaCompany Report #001-0945-M0101440
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2700 MG		Autoimmune Disorder Decreased Appetite	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER		Pain In Extremity	Professional				
ORAL		Vomiting					

Date:02/25/02ISR Number: 3889480-8Report Type:Periodic Company Report #001-0945-M0100221
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Pneumonia	Health Professional	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/27/02ISR Number: 3874878-4Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 162400

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1500MG PO QHS	Confusional State		Gabapentin	PS		ORAL
Initial or Prolonged PTA	Fall					
Required Intervention to Prevent Permanent Impairment/Damage	Lethargy Memory Impairment Oral Intake Reduced Visual Disturbance		Atorvastatin Glipizide Atenolol Lisinopril Fluoxetine Morphine Ir Morphine Sr Clonazepam Tramadol Actos Loratadine Dipyridamole/Aspirin	C C C C C C C C C C C C		

Date:02/27/02ISR Number: 3876619-3Report Type:Expedited (15-DaCompany Report #061-0945-M0200036
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 300 MG, PER	Asthenia Coordination Abnormal	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL	Difficulty In Walking	Professional Company Representative				

Date:02/27/02ISR Number: 3876999-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200233
 Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged PER ORAL	Condition Aggravated Drug Withdrawal Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL

Trigeminal Neuralgia
Weight Decreased

(Medroxyprogesterone
Acetate, Estrogens
Conjugated) C

Date:02/28/02ISR Number: 3877482-7Report Type:Expedited (15-DaCompany Report #061-0945-M0200034
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (TID), PER ORAL	Drug Toxicity Overdose Respiratory Depression	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:02/28/02ISR Number: 3878169-7Report Type:Expedited (15-DaCompany Report #2002-02-1107
Age: Gender: I/FU:I

Outcome	PT
Other	Abortion Induced Amniocentesis Abnormal Foetal Disorder Pregnancy Transmission Of Drug Via

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Semen
Ultrasound Scan Abnormal

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1200 MG QD		Health Professional	Rebetol (Ribavirin) Capsules	PS		ORAL
ORAL		Company Representative	Intron A (Interferon Alfa-2b Recombinant) Injectable Solution	SS		
SUBCUTANEOUS	3 MU TIW					
SUBCUTANEOUS			Gabapentin	SS		
			Olanzapine	SS		

Date:02/28/02ISR Number: 3878219-8Report Type:Expedited (15-DaCompany Report #HQ0996826FEB2002
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional Company Representative	Phenergan With Codeine (Promethazine Hydrochloride/Codeine Phosphate, Unspec)	PS		ORAL
2 OZ				Darvocet-N (Dextropropoxyphene/Paracetamol,)	SS		
(OVERDOSE AMOUNT)							
"3" (OVERDOSE AMOUNT)				Gabapentin (Gabapentin,)	SS		
"3" (OVERDOSE AMOUNT)				Glucovance			

OVERDOSE

(Glibenclamide/Metformin Hydrochloride,) SS

AMOUNT

UNKNOWN

Trazodone (Trazodone) SS

"3" (OVERDOSE AMOUNT)

Xanax (Alprazolam,) SS

"2" (OVERDOSE AMOUNT)

Date:02/28/02ISR Number: 3878420-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200232
Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aneurysm Hypoaesthesia	Consumer	Neurontin (Gabapentin)	PS		
900 MG (300 MG, TID)		Pain In Extremity					
UNKNOWN							

Date:02/28/02ISR Number: 3878421-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200245
Age: Gender:Male I/FU:I

Outcome	PT
Other	Cognitive Disorder Convulsion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Electroencephalogram Abnormal				
Dose	Duration		Report Source	Product	Role	Manufacturer
			Consumer	Neurontin (Gabapentin)	PS	
PER ORAL				Unspecified Decongestant	C	ORAL

Date:02/28/02ISR Number: 3889398-0Report Type:Periodic Company Report #001-0945-M0100608
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		
Other		Cerebrovascular Accident Convulsion Headache					

Date:02/28/02ISR Number: 3889400-6Report Type:Periodic Company Report #001-0945-M0100687
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional	Neurontin (Gabapentin)	PS		
Other		White Blood Cell Count Decreased					

Date:02/28/02ISR Number: 3889402-XReport Type:Periodic Company Report #001-0945-M0100694
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional	Neurontin (Gabapentin)	PS		
Other		Retinopathy					

Date:02/28/02ISR Number: 3889403-1Report Type:Periodic Company Report #001-0945-M0100701
Age: Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Hospitalization -		Contusion	Health	Neurontin			
Initial or Prolonged		Haemorrhagic Stroke	Professional	(Gabapentin)	PS		ORAL
PER ORAL			Company Representative				

Date:02/28/02ISR Number: 3889404-3Report Type:Periodic Company Report #001-0945-M0100735
 Age: Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Convulsion	Health	Neurontin			
			Professional	(Gabapentin)	PS		

Date:02/28/02ISR Number: 3889406-7Report Type:Periodic Company Report #001-0945-M0100756
 Age:59 YR Gender:Male I/FU:I

Outcome	PT
Other	Anxiety
	Asthenia
	Burning Sensation
	Drug Ineffective
	Heart Rate Decreased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
900 MG TID		Hypoaesthesia	Consumer Health	Neurontin (Gabapentin)	PS		
		Hypotension					
		Insomnia					
		Paraesthesia	Professional	Tizanidine Hydrochloride Voixx (Rofecoxib)	SS C		
		Pruritus					
		Tremor					

Date:02/28/02ISR Number: 3889407-9Report Type:Periodic Company Report #001-0945-M0100785
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		
Hospitalization -		Anaemia					
Initial or Prolonged		Eye Pain					
SEE IMAGE							

Date:02/28/02ISR Number: 3889408-0Report Type:Periodic Company Report #001-0945-M0100827
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Convulsion					
300 MG (100		Coordination Abnormal					
MG, TID), PER		Depressed Level Of					
ORAL		Consciousness					
		Eye Disorder					
		Hypokinesia					
		Muscle Twitching					

Date:02/28/02ISR Number: 3889409-2Report Type:Periodic Company Report #001-0945-M0100829
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Blood Glucose Fluctuation	Consumer Health	Neurontin (Gabapentin)	PS
1200 MG TID		Professional	Celebrex (Celecoxib)	C
			Prevacid (Lansoprazole)	C
			Remeron (Mirtazapine)	C
			Lorazepam (Lorazepam)	C
			Phenergan (Promethazine Hydrochloride)	C

Date:02/28/02ISR Number: 3889410-9Report Type:Periodic Company Report #001-0945-M0100856
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Glaucoma	Health Professional	Neurontin (Gabapentin)	PS		
1200 MG DAILY				Effexor (Venlafaxine Hydrochloride)	SS		
150 MG DAILY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3889411-0Report Type:Periodic
 Age: Gender: I/FU:I

Company Report #001-0945-M0100470

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889412-2Report Type:Periodic
 Age: Gender: I/FU:I

Company Report #001-0945-M0100471

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889413-4Report Type:Periodic
 Age:44 YR Gender:Female I/FU:I

Company Report #001-0945-M0100480

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Sudden Hearing Loss	Health Professional	Neurontin (Gabapentin)	PS		

300 MG (TID)

/ SEE IMAGE

SUBCUTANEOUS 0.25 MG

(EVERY OTHER

DAY),

SUBCUTANEOUS

Betaseron (Glucose,
 Albumin Human,
 Interferon Beta) SS

Famvir (Famciclovir) C

Date:02/28/02ISR Number: 3889414-6Report Type:Periodic
 Age:54 YR Gender:Female I/FU:I

Company Report #001-0945-M0100503

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Alopecia Oedema	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), PER ORAL			Professional				
				Zoloft (Sertraline Hydrochloride)	SS		
				Zantac (Ranitidine Hydrochloride)	C		
				Adderall (Amfetamine Sulfate)	C		
				Vitamin D (Ergocalciferol)	C		
				Vitamin E (Tocopherol)	C		
				Vitamin B Complex (Pyridoxine Hydrochloride, Thiamine Hydrochloride,	C		
				Ginkgo Biloba (Ginkgo Biloba)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Beta One, 3-D Lukan Trazodone (Trazodone)	C		

Freedom Of Information (FOI) Report

Valium (Diazepam)	C
Tenuate Dospan (Amfepramone Hydrochloride)	C
Vioxx (Rofecoxib)	C
Chlorzoxazone (Chlorzoxazone)	C
Folic Acid (Folic Acid)	C
Darvocet (Paracetamol, Dextropropoxyphene)	C
Fosamax (Alendronate Sodium)	C
Estrogen Nos (Estrogen Nos)	C
Benadryl (Diphenhydramine Hydrochloride)	C
Lucosamine W/Chondroitin Sulfates (Glucosamine, Chondroitin)	C
Betacarotene (Betacarotene)	C
Sonata	C
Prilosec (Omeprazole)	C
Parafon Forte (Chlorzoxazone, Paracetamol)	C
Serzone (Nefazodone Hydrochloride)	C

Date:02/28/02ISR Number: 3889415-8Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-M0100551

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Health	Neurontin			
Other		Sleep Disorder	Professional	(Gabapentin)	PS		ORAL
900 MG (TID),			Company				
PER ORAL							

Representative		
	Toprol (Metoprolol	
	Succinate)	C
	Diovan (Valsartan)	C
	Lipitor	
	(Atorvastatin)	C
	Tricor (Fenofibrate)	C
	Avandia	
	(Rosiglitazone)	C
	Amaryl (Glimepiride)	C
	Glucovance	
	(Metformin	
	Hydrochloride,	
	Glibenclamide)	C
	Glucosamine	
	(Glucosamine)	C
	Dilantin (Phenytoin	
	Sodium)	C
	Alprazolam	C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Welchol 3 C

Date:02/28/02ISR Number: 3889416-XReport Type:Periodic Company Report #001-0945-M0100577
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Confusional State Dyspnoea	Health Professional	Neurontin (Gabapentin)	PS		
Other (DAILY)	Overdose					

Date:02/28/02ISR Number: 3889417-1Report Type:Periodic Company Report #001-0945-M0100586
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Hyponatraemia	Health Professional	Neurontin (Gabapentin)	PS		
			Phenobarbital (Phenobarbital)	C		

Date:02/28/02ISR Number: 3889422-5Report Type:Periodic Company Report #001-0945-M0100589
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Hallucination	Consumer	Neurontin (Gabapentin)	PS		
300 MG (100 MG, TID)			Levofloxacin (Levofloxacin)	C		
			Furosemide (Furosemide)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3600 MG (TID) 900 MG (DAILY) / SEE IMAGE		Drug Ineffective Mania Overdose Schizophrenia	Health Professional	Neurontin (Gabapentin) Seroquel Geodon	PS SS SS		 ORAL
40 MG (BID), PER ORAL / SEE IMAGE				Beer	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 300 M (DAILY), PER ORAL		Coordination Abnormal Depression Headache Sleep Disorder Somnolence Suicidal Ideation Thinking Abnormal	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3889426-2Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #001-0945-M0100460

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eructation	Consumer	Neurontin			
		Loss Of Consciousness	Health	(Gabapentin)	PS		ORAL
2700 MG (900							
MG TID) PER		Overdose	Professional				
ORAL							

Evista (Raloxifene Hydrochloride)	C
Oxycontin (Oxycodone Hydrochloride)	C
Singulair (Montelukast)	C
Paxil (Paroxetine Hydrochloride)	C
Imitrex (Sumatriptan Succinate)	C
Dihydroergotamine (Dhe)	C
Phenergan (Promethazine Hydrochloride)	C

Date:02/28/02ISR Number: 3889429-8Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-M0100461

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaemia	Health	Neurontin			
		Overdose	Professional	(Gabapentin)	PS		ORAL
2700 MG TID							
PER ORAL							
				Seroquel (Quetiapine)	SS		
650 MG DAILY				Aspirin (Acetylsalicylic Acid)	C		
				Multivitamin			

(Ergocalciferol,
 Ascorbic Acid, Folic
 Acid, Thiamine
 Hydrochloride, C
 Vitamin E
 (Tocopherol) C
 Buspar (Buspirone
 Hydrochloride) C
 Vistaril
 (Hydroxyzine
 Embonate) C
 Insulin (Insulin) C

Date:02/28/02ISR Number: 3889430-4Report Type:Periodic Company Report #001-0945-M0100462
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3889432-8Report Type:Periodic Company Report #001-0945-M0100463
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889434-1Report Type:Periodic Company Report #001-0945-M0100464
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889436-5Report Type:Periodic Company Report #001-0945-M0100465
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889437-7Report Type:Periodic Company Report #001-0945-M0100466
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889456-0Report Type:Periodic Company Report #001-0945-M0100467
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889458-4Report Type:Periodic Company Report #001-0945-M0100468
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889459-6Report Type:Periodic Company Report #001-0945-M0100469
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889461-4Report Type:Periodic Company Report #001-0945-M0100215
Age: 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

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion Overdose Suicide Attempt	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889478-XReport Type:Periodic Company Report #001-0945-M0100216
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Haemorrhage Intracranial	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889482-1Report Type:Periodic Company Report #001-0945-M0100307
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other SEE IMAGE		Dizziness Medication Error	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Memory Impairment Nystagmus Overdose Sedation Vith Nerve Paralysis	Professional	Hydrocodone W/Acetaminophen	C		

Date:02/28/02ISR Number: 3889483-3Report Type:Periodic Company Report #001-0945-M0100312
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Visual Field Defect	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889485-7Report Type:Periodic Company Report #001-0945-M0100313

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Visual Field Defect	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3889486-9Report Type:Periodic Company Report #001-0945-M0100335

Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Hypertension	Consumer	Neurontin (Gabapentin)	PS		ORAL

900 MG (TID)

PER ORAL

Date:02/28/02ISR Number: 3889487-0Report Type:Periodic Company Report #001-0945-M0100409

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness	Health Professional	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3889488-2Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-M0100436

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG		Pyrexia	Health Professional	Neurontin (Gabapentin)	PS		ORAL

(DAILY) PER

ORAL

Cardizem Cd (Diltiazem Hydrochloride)	C
Visken (Pindolol)	C
Xanax (Alprazolam)	C
Bentyl (Dicycloverine Hydrochloride)	C
Aldactazide (Spironolactone, Hydrochlorothiazide)	C

Date:02/28/02ISR Number: 3889489-4Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #001-0945-M0100441

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 900 MG		Asthenia Overdose	Consumer	Neurontin (Gabapentin)	PS		ORAL

(DAILY) PER

ORAL

Date:02/28/02ISR Number: 3890064-6Report Type:Periodic
 Age:85 YR Gender:Female I/FU:I

Company Report #001-0945-M0100865

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Deafness	Consumer	Neurontin			

100 MG, (DAILY), PER ORAL
 Headache (Gabapentin) PS ORAL
 Hearing Impaired
 Keratoconjunctivitis

Sicca Naldol C
 Neck Pain Insulin C
 Lotrel Amlodipine
 Besylate, Benazepril
 Hydrochloride) C
 Prilosec
 (Omeprazole) C
 Plavix (Clopidogrel) C
 Buspar (Buspirone
 Hydrochloride) C
 Zocor (Simvastatin) C

Date:02/28/02ISR Number: 3890067-1Report Type:Periodic Company Report #001-0945-M0100914
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Areflexia	Consumer	Neurontin			
		Neuropathy Peripheral		(Gabapentin)	PS		ORAL
1800 MG (600							
MG, TID), PER							
ORAL				Captopril	C		
				Levothyroxine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Piroxicam C
 Oxybutynin C
 Cetirizine
 Hydrochloride C
 Acetylsalicylic Acid C
 Ethinylestradiol,
 Norethisterone C

Date:02/28/02ISR Number: 3890069-5Report Type:Periodic
 Age:56 YR Gender:Female I/FU:I

Company Report #001-0945-M0100980

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Mouth	Consumer	Neurontin			
		Libido Decreased		(Gabapentin)	PS		
		Oedema Peripheral		Baclofen	SS		
		Overdose		Acetylsalicylic			
		Pain		Acid, Caffeine			
		Weight Increased		Anhydrous,			
		White Blood Cell Count		Butalbital	C		
				Unknown			
				Chemotherapeutic			
				Agent	C		

Date:02/28/02ISR Number: 3890072-5Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-M0101010

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
		Drug Withdrawal Syndrome	Health Professional	(Gabapentin)	PS		
				Amitriptyline	C		
				Methadone	C		
				Unspecified Muscle			
				Relaxant	C		
				Montelukast	C		
				Lansoprazole	C		
				Ipratropium Bromide	C		
				Pirbuterol Acetate	C		
				Triamcinolone			
				Acetonide	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Renal Failure Acute	Health Professional	Neurontin (Gabapentin)	PS		
1800 MG							
(TID),							
UNKNOWN							
				Amitriptyline	C		
				Hydroxychloroquine	C		
				Synthroid (Levothyroxine)	C		
				Zoloft (Sertraline Hydrochloride)	C		
				Oxycontin (Oxycodone Hydrochloride)	C		
				Percocet (Paracetamol,			

Freedom Of Information (FOI) Report

Oxycodone
 Hydrochloride) C
 Prednisone C
 Lactulose C
 Evista (Raloxifene
 Hydrochloride) C
 Calcium C
 Miacalcin Nasal
 Spray (Calcitonin,
 Salmon) C
 Multivitamin
 (Ergocalciferol,
 Ascorbic Acid, Folic
 Acid, Thiamine
 Hydrochloride, C
 Metamucil (Psyllium
 Hydrophilic
 Mucilloid) C
 Compazine
 (Prochlorperazine
 Edisylate) C

Date:02/28/02ISR Number: 3890078-6Report Type:Periodic
 Age:74 YR Gender:Female I/FU:I

Company Report #001-0945-M0101025

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG, DAILY, UNKNOWN		Dizziness Heart Rate Increased Hyperpyrexia Hypertension Hypotonia Nausea	Consumer Health Professional	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890080-4Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-M0101028

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Atrioventricular Block	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890082-8Report Type:Periodic Company Report #001-0945-M0101050
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
PER ORAL							

Date:02/28/02ISR Number: 3890084-1Report Type:Periodic Company Report #001-0945-M0101057
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Overdose Suicide Attempt	Health Professional	Neurontin (Gabapentin) Alcohol (Ethanol)	PS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3890086-5Report Type:Periodic
 Age: Gender: I/FU:I

Company Report #001-0945-M0101059

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Thrombocytopenia	Health Professional	Neurontin (Gabapentin)	PS		
				Four Unspecified Medicatoinis	C		

Date:02/28/02ISR Number: 3890091-9Report Type:Periodic
 Age:37 YR Gender:Male I/FU:I

Company Report #001-0945-M0101081

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error Overdose	Health Professional	Neurontin (Gabapentin)	PS		ORAL
3600 MG		Pneumonia					
(DAILY), PER							
ORAL				Clonazepam	C		
				Benztropeine	C		
				Fluphenazine Hydrochloride	C		
				Lorazepam	C		
				Perphenazine	C		
				Phenytoin Sodium	C		
				Risperidone	C		

Date:02/28/02ISR Number: 3890106-8Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-M0101089

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia Communication Disorder	Consumer Health	Neurontin (Gabapentin)	PS		
1200 MG		Loss Of Consciousness	Professional				
(THREE TIMES		Mental Impairment					
DAILY),							

Movement Disorder

Estrogens Conjugated C
Telmisartan C
Hydrochlorothiazide C
Paracetamol C

Date:02/28/02ISR Number: 3890110-XReport Type:Periodic Company Report #001-0945-M0101093
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema Urticaria	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890111-1Report Type:Periodic Company Report #001-0945-M0101112
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema Urticaria	Health Professional	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3890112-3Report Type:Periodic
Age:57 YR Gender:Female I/FU:I

Company Report #001-0945-M0101161

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Neurontin			
1600 MG		Feeling Abnormal	Professional	(Gabapentin)	PS		
(QID),		Non-Hodgkin'S Lymphoma					
				Carbatrol			
				(Carbamazepine)	C		
				Chemotherapy	C		

Date:02/28/02ISR Number: 3890113-5Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-M0101171

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Consumer	Neurontin			
		Hypersensitivity	Health	(Gabapentin)	PS		
		Tachycardia	Professional	Klonopin			
				(Clonazepam)	SS		
				Nexium	SS		
				Betapace (Sotalol			
				Hydrochloride)	SS		
				Prednisone			
				(Prednisone)	SS		
				Trazodone			
				(Trazodone)	SS		

Date:02/28/02ISR Number: 3890115-9Report Type:Periodic
Age:76 YR Gender:Female I/FU:I

Company Report #001-0945-M0101205

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200 MG		Headache	Consumer	Neurontin			
(DAILY), PER		Initial Insomnia		(Gabapentin)	PS		ORAL
ORAL		Pain					
		Somnolence					

Weight Increased

Zolpidem Tartrate	C
Atorvastatin	C
Metformin	
Hydrochloride	C
Moexipril	
Hydrochloride	C

Date:02/28/02ISR Number: 3890116-0Report Type:Periodic
 Age:72 YR Gender:Male I/FU:I

Company Report #001-0945-M0101211

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (300 MG, TID), PER ORAL	Abdominal Pain Upper Atrial Fibrillation Constipation Diarrhoea	Consumer	Neurontin (Gabapentin)	PS		ORAL
1000 MCG (500 MCG, BID), PER ORAL			Tikosyn (Dofetilide)	SS		ORAL
PER ORAL			Methadone (Methadone)	SS		ORAL
			Coumadin (Warfarin Sodium) Tylenol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Paracetamol) C
 Altace (Ramipril) C
 Coreg (Carvedilol) C

Date:02/28/02ISR Number: 3890117-2Report Type:Periodic
 Age:40 YR Gender:Male I/FU:I

Company Report #001-0945-M0101292

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG (THREE TIMES A DAY), PER ORAL		Nephrolithiasis	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Oxycontin (Oxycodone
 Hydrochloride) C

Date:02/28/02ISR Number: 3890119-6Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-M0101294

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (TID), PER ORAL		Disorientation Dizziness Neuropathy Peripheral Syncope	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Valsartan Diltiazem Hydrochloride Hydrochlorothiazide Pravastatin Sodium Theophylline Salmeterol Xinafoate Triamcinolone Acetonide Timolol Maleate	C C C C C C C C C C		

Date:02/28/02ISR Number: 3890123-8Report Type:Periodic Company Report #001-0945-M0101342

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890128-7Report Type:Periodic Company Report #001-0945-M0101343

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890130-5Report Type:Periodic Company Report #001-0945-M0101384

Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Somnolence	Consumer Health	Neurontin (Gabapentin)	PS		ORAL

900 MG (300

MG, TID), PER

ORAL

Paxil (Paroxetine

22-Aug-2005 10:40 AM

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Freedom Of Information (FOI) Report

Hydrochloride) C

Date:02/28/02ISR Number: 3890132-9Report Type:Periodic
Age:65 YR Gender:Female I/FU:I

Company Report #001-0945-M0101300

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG DAILY PER ORAL		Blood Pressure Increased Burning Sensation Cataract Diplopia Dizziness Dysarthria Electrocardiogram Abnormal Liver Function Test Abnormal Motion Sickness Photophobia Photopsia Retinoschisis Visual Disturbance	Consumer	Neurontin (Gabapentin) 1) (Nabumetone)	PS C		ORAL

Date:02/28/02ISR Number: 3890134-2Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-M0101394

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 1800 MG (THREE TIMES DAILY), PER ORAL		Deafness	Health Professional	Neurontin (Gabapentin) Aspirin (Acetylsalicyclic Acid) Darvocet (Paracetamol,	PS C		ORAL

Dextropropoxyphene) C
 Prilosec
 (Omeprazole) C
 Vicodin
 (Paracetamol,
 Hydrocodone
 Bitartate) C
 Zocor (Simvastatin) C
 Flonase (Fluticasone
 Propionate) C
 Advil (Ibuprofen) C

Date:02/28/02ISR Number: 3890135-4Report Type:Periodic Company Report #001-0945-M0101314
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3890137-8Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-M0101325

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Deafness	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (THREE TIMES DAILY PER ORAL							

Serzone (Nefazodone Hydrochloride) C
Risperdal (Risperidone) C
(Benzatropine Mesilate) C

Date:02/28/02ISR Number: 3890138-XReport Type:Periodic
Age:5 YR Gender:Female I/FU:I

Company Report #001-0945-M0101430

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Pneumonia	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890141-XReport Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-M0101335

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890160-3Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-M0101435

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Myocardial Infarction	Health	Neurontin			

Other			Professional	(Gabapentin)	PS		
RESPIRATORY							
(INHALATION)	INHALATION			Remeron (Mirtazapine)	SS		

Date:02/28/02ISR Number: 3890163-9Report Type:Periodic Company Report #001-0945-M0101336
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890174-3Report Type:Periodic Company Report #001-0945-M0101337
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Death	Consumer	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3890175-5Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-M0200003

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890177-9Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-M0101338

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890179-2Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-M0101339

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890182-2Report Type:Periodic
 Age: Gender:Unknown I/FU:I

Company Report #001-0945-M0101340

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890184-6Report Type:Periodic
 Age: Gender: I/FU:I

Company Report #001-0945-M0200004

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890186-XReport Type:Periodic Company Report #001-0945-M0101341
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890190-1Report Type:Periodic Company Report #001-0945-M0200005
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890194-9Report Type:Periodic Company Report #001-0945-M0200006
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3890205-0Report Type:Periodic Company Report #001-0945-M0200007

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890238-4Report Type:Periodic Company Report #001-0945-M0200008

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890239-6Report Type:Periodic Company Report #001-0945-M0200009

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890243-8Report Type:Periodic Company Report #001-0945-M0200010

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890245-1Report Type:Periodic Company Report #001-0945-M0200011

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890250-5Report Type:Periodic Company Report #001-0945-M0200012
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890251-7Report Type:Periodic Company Report #001-0945-M0200189
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer Other	Neurontin (Gabapentin)	PS		
300 MG				Sertraline Hydrochloride	SS		
50 MG				Tocopherol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3890253-0Report Type:Periodic
 Age:46 YR Gender:Female I/FU:F

Company Report #001-0945-M0000102

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract	Health	Neurontin			
		Refraction Disorder	Professional	(Gabapentin)	PS		ORAL
PER ORAL				Klonopin			
				(Clonazepam)	C		
				Ultram (Tramadol			
				Hydrochloride)	C		
				Imitrex (Sumatriptan			
				Succinate)	C		

Date:02/28/02ISR Number: 3890255-4Report Type:Periodic
 Age:43 YR Gender:Female I/FU:F

Company Report #001-0945-M0000132

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Visual Disturbance	Health	Neurontin			
			Professional	(Gabapentin)	PS		ORAL
400 MG (100MG							
IN AM; 300MG							
IN PM); 100							
MG (DAILY),							
PER ORAL				Ativan (Lorazepam)	C		

Date:02/28/02ISR Number: 3890258-XReport Type:Periodic
 Age:45 YR Gender:Male I/FU:F

Company Report #001-0945-M0000149

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Liver Disorder	Health	Neurontin			
		Oedema Peripheral	Professional	(Gabapentin)	PS		ORAL
PER ORAL				Oxycontin (Oxycodone			
		Swelling		Hydrochloride)	C		

Date:02/28/02ISR Number: 3890265-7Report Type:Periodic Company Report #001-0945-M0001268
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Health	Neurontin Capsules			
		Death	Professional	300 Mg (Gabapentin)	PS		
300 MG (TID)							

Date:02/28/02ISR Number: 3890275-XReport Type:Periodic Company Report #001-0945-M0100030
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Anaemia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Prevacid (Lansoprazole)	C		
				Coumadin (Warfarin Sodium)	C		
				(Insulin)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Lasix (Furosemide)	C		
				(Potassium)	C		
				Zaroxolyn			

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Freedom Of Information (FOI) Report

(Metolazone) C
 (Sotalol) C
 Colace (Docusate Sodium) C
 (Digoxin) C

Date:02/28/02ISR Number: 3890279-7Report Type:Periodic Company Report #001-0945-M0100034
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angioneurotic Oedema Urticaria	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890293-1Report Type:Periodic Company Report #001-0945-M0100036
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Tremor Unevaluable Event	Health Professional Company Representative	Neurontin (Gabapentin) Unspecified Medications	PS C		

Date:02/28/02ISR Number: 3890306-7Report Type:Periodic Company Report #001-0945-M0100060
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Agitation Confusional State Hallucination Unevaluable Event	Health Professional	Neurontin (Gabapentin) Flexeril (Cyclobenzaprine Hydrochloride) Norco (Paracetamol, Hydrocodone Bitartrate) Senna S (Senna) Cytotec (Misoprostol) Vioxx (Rofecoxib) Flomax	PS SS C C C C		

(Morniflumate) C
 Zoloft (Sertraline
 Hydrochloride) C
 Cardura (Doxazosin
 Mesilate) C

Date:02/28/02ISR Number: 3890311-0Report Type:Periodic Company Report #001-0945-M0100095
 Age:65 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged SEE IMAGE	Alopecia Drug Ineffective	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Professional	Glucophage (Metformin Hydrochloride) Glyburide (Glibenclamide) (Etodolac)	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3890314-6Report Type:Periodic
Age:40 YR Gender:Female I/FU:I

Company Report #001-0945-M0100096

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1600 MG (400 MG, QID), PER ORAL		Depression Disease Recurrence Oedema Peripheral Post-Traumatic Stress Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Zoloft (Sertraline Hydrochloride) Vicodin (Paracetamol, Hydrocodone Bitrtrate) Colace (Docusate Sodium) Seroquel (Quetiapine)	C C C C		

Date:02/28/02ISR Number: 3890317-1Report Type:Periodic
Age:54 YR Gender:Male I/FU:I

Company Report #001-0945-M0100120

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/28/02ISR Number: 3890318-3Report Type:Periodic
Age:60 YR Gender:Male I/FU:I

Company Report #001-0945-M0100183

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other SEE IMAGE		Deafness	Consumer Health Professional	Neurontin (Gabapentin) Topamax (Topiramate) Tiazac (Diltiazem Hydrochloride) Lipitor	PS C C		ORAL

(Atorvastatin) C
 Paxil (Proxetine Hydrochloride) C
 Flomax (Tamsulosin Hydrochloride) C
 Ambien (Zolpidem Tartrate) C
 Imitrex (Sumatriptan Succinate) C

Date:02/28/02ISR Number: 3890320-1Report Type:Periodic Company Report #001-0945-M0100199
 Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		White Blood Cell Count Decreased	Health Professional	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/02ISR Number: 3890324-9Report Type:Periodic
Age:10 YR Gender:Male I/FU:F

Company Report #001-0945-M0000181

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged SEE IMAGE	Abnormal Behaviour Anorexia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other	Anxiety Aphasia Attention Deficit/Hyperactivity Disorder Bruxism Cognitive Disorder Confusional State Conversion Disorder Convulsion Cough Euphoric Mood Hallucination Incontinence Insomnia Jaw Disorder Mania Memory Impairment Psychotic Disorder Stupor	Professional	Tegretol(Carbamazepi ne) Dexedrine (Dexamfetamine Sulfate) Risperdal (Risperidone)	C C C		

Date:02/28/02ISR Number: 3890328-6Report Type:Periodic
Age:50 YR Gender:Female I/FU:F

Company Report #001-0945-M0000334

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other SEE IMAGE	Hypoaesthesia Tunnel Vision	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Glucophage (Metformin Hydrochloride) Diabeta (Glibenclamide)	C C		

Date:02/28/02ISR Number: 3890334-1Report Type:Periodic Company Report #001-0945-M0000897
Age:57 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged SEE IMAGE	Suicide Attempt	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Professional	Sinemet (Levodopa, Cabidopa) Digoxin	C C		

Date:02/28/02ISR Number: 3890340-7Report Type:Periodic Company Report #001-0945-M0001030
Age:11 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Anorexia Blood Albumin Decreased Blood Glucose Increased Conjunctival Hyperaemia Convulsion Cushingoid

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Diarrhoea Dyspepsia Face Oedema	Health Professional	Neurontin Capsules 300 Mg (Gabapentin)	PS		ORAL
SEE IMAGE		Headache Hypersensitivity	Company Representative	Lamictal (Lamotrigine)	SS		
SEE IMAGE		Malaise Mouth Ulceration		Prednisone	SS		
SEE IMAGE		Nocturia		Ritali N (Methylphenidate Hydrochloride)	C		
		Pharyngolaryngeal Pain Pollakiuria Pruritus Stomach Discomfort Urticaria Vision Blurred Vomiting Weight Increased White Blood Cell Count Increased		Dextrostat (Dexamfetamine Sulfate) Clonidine Benadryl (Diphenhydramine Hydrochloride) Atarax (Hydroxyzine Hydrochloride) Topical Steroid Tylenol (Paracetamol) Zantac (Ranitidine Hydrochloride)	C C C C C C C C C		

Date:02/28/02ISR Number: 3890663-1Report Type:Periodic Company Report #001-0073-M0100450
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Headache Liver Function Test Abnormal	Consumer	Dilantin (Phenytoin Sodium) Neurontin (Gabapentin)	PS SS		ORAL
900 MG (300 MG, TID), PER ORAL				Sudafed			

(Pseudoephedrine Hydrochloride)	SS
Caffeine (Caffeine Anhydrous)	SS
Nutrasweet (Aspartame)	SS
Vicotuss	C
Neurontin (Gabapentin)	C

Date:03/01/02ISR Number: 3877333-0Report Type:Expedited (15-DaCompany Report #034-0945-M0200002
 Age:21 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Foreign	Gabapentin			
1200 MG		Drug Interaction	Health	(Gabapentin)	PS		ORAL
(TID), PER		Neutropenia	Professional				
ORAL			Company				
PER ORAL			Representative	(Clozapine)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/02ISR Number: 3877598-5Report Type:Expedited (15-DaCompany Report #A200378

Age:50 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - INTRAMUSCULAR 40.00 MG	Bradycardia	Foreign	Feldene Capsules	PS		
Initial or Prolonged TOTAL:DAILY:I	Confusional State	Health				
	Drug Interaction	Professional				
NTRAMUSCULAR 2200.00 MG	Drug Level Above Therapeutic		Gabapentin	SS		ORAL
TOTAL:ORAL 12.00 MG	Dysarthria		Tizanidine	SS		ORAL
TOTAL:ORAL 20.00 MG	Dysphagia					
	Hallucination		Citalopram	SS		
TOTAL	Hypertension					
	Hypotension		Voltaren	C		
	Hypovolaemia		Vioxx	C		
	Medication Error					

Date:03/01/02ISR Number: 3877642-5Report Type:Expedited (15-DaCompany Report #044-0945-M0200031

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Disability 1200 MG (FOUR TIMES DAILY), PER ORAL	Ecchymosis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Professional				

Date:03/01/02ISR Number: 3877643-7Report Type:Expedited (15-DaCompany Report #033-0945-M0200017

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 100 MG		Aggression Agitation	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER		Delirium	Professional				
ORAL		Hallucinations, Mixed		(Amitriptyline Hydrochloride)	SS		ORAL
50 MG							
(DAILY), PER							
ORAL				(Clonazepam)	C		
				(Abacavir)	C		
				(Didanosine)	C		
				(Ritonavir)	C		
				(Lopinavir/Ritonavir)	C		

Date:03/01/02ISR Number: 3877645-0Report Type:Expedited (15-DaCompany Report #055-0945-M0200011
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (TWICE		Haemorrhoid Operation	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
DAILY), PER							
ORAL				(Osseine-Hydroxyapat ite Complex)	C		
				(Atorvastatin)	C		
				(Levothyroxine Sodium)	C		
				(Tibolone)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/02ISR Number: 3878753-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200247

Age:76 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged PER ORAL Disability	Cerebrovascular Accident Difficulty In Walking Fall Lumbar Vertebral Fracture Macular Degeneration	Health Professional	Neurontin (Gabapentin) Amitriptyline Estrogens (Conjugated) Nitrofurantoin Polysaccharide-Iron Complex Calcium Carbonate Calcitonin, Salmon Oxybutynin	PS SS C C C C C C		ORAL

Date:03/01/02ISR Number: 3878810-9Report Type:Expedited (15-DaCompany Report #A201293

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 100.00 MG Intervention to TOTAL:DAILY Prevent Permanent Impairment/Damage	Anxiety Appetite Disorder Balance Disorder Convulsion Decreased Appetite Depression Fatigue Headache Irritability Libido Decreased Palpitations Sedation Sleep Disorder Weight Decreased	Consumer Health Professional	Zoloft Tablets Neurontin Klonopin Keppra Gaba	PS SS SS C C		

Date:03/01/02ISR Number: 3878826-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200001

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 4500 MG (TID) Other		Drug Effect Decreased	Consumer Health Professional	Neurontin (Gabapentin)	PS		

Date:03/01/02ISR Number: 3878836-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200121
Age: Gender:Female I/FU:F

Outcome	PT
Other	Anxiety Balance Disorder Convulsion Decreased Appetite Depression Energy Increased Fatigue Headache Irritability Libido Decreased Palpitations Sedation

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Sleep Disorder Weight Decreased					
100 MG	(DAILY)		Consumer Health Professional	Neurontin (Gabapentin) Sertraline Hydrochloride	PS SS		
				Clonazepam Levetiracetam Gaba	SS C C		

Date:03/05/02ISR Number: 3879057-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200266
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (TID) Other		Blood Potassium Increased Drug Ineffective Haemorrhage Heart Rate Decreased Red Blood Cell Count Decreased	Consumer	Neurontin (Gabapentin)	PS		

Date:03/05/02ISR Number: 3879118-8Report Type:Expedited (15-DaCompany Report #NSADSS2002005632
Age:5 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG, 3 IN 1 DAY(S), GASTUBE 450 MG, 2 IN		Inappropriate Antidiuretic Hormone Secretion Medical Device Complication Oral Intake Reduced		Topamax (Tablet) (Topiramate) Trileptal (Oxcarbazepine)	PS SS		

1 DAY(S),
Pancreatitis
Urinary Tract Infection
GASTUBE :
Vomiting
300 MG, 1 IN

1 DAY(S) ,
Neurontin
(Gabapentin) SS
450 MG, 2 IN

1 DAY(S),
GASTUBE; 300
MG, 1 IN 1

DAY(S),
Provigil (Modafinil) C
Zantac (Ranitidine
Hydrochloride) C
Reglan
(Metoclopramide) C
Poly Vitamin With
Iron (Polyvitamin) C
Claritin
(Loratadine) C
Motrin (Ibuprofen) C
Colace (Docusate
Sodium) C
(Tums) (Tums) C
Morphine (Morphine) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/05/02ISR Number: 3879649-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200001

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged UNKNOWN Other (TID), UNKNOWN 1200 MG (BID), PER ORAL 200 MG (BID), PER ORAL	Convulsion	Consumer Health Professional	Neurontin (Gabapentin) (Carbamazepine) (Lamotrigine)	PS SS SS		ORAL
			(Novolin R) (Paracetamol, Oxycodone Hydrochloride) (Vancomycin) (Ceftriaxone Sodium) (Metformin Hydrochloride) (Glibenclamide) (Tamoxifene Citrate) (Oxybutynin) (Warfarin Sodium) (Paracetamol)	C C C C C C C C C C		

Date:03/05/02ISR Number: 3879668-4Report Type:Expedited (15-DaCompany Report #A204121

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required ORAL Intervention to ORAL	Alopecia Condition Aggravated	Consumer	Zoloft Tablets Gabapentin	PS SS		ORAL ORAL

Prevent Permanent
Impairment/Damage

Contusion
Crying
Depression
Dizziness
Drug Ineffective
Drug Withdrawal Syndrome
Face Oedema
Fatigue
Movement Disorder
Neoplasm Skin
Nervous System Disorder
Suicidal Ideation
Urticaria
Weight Increased

Soma
Prednisone

C
C

Date:03/05/02ISR Number: 3879671-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200241
Age: Gender:Female I/FU:I

Outcome
Other

PT
Alopecia
Contusion
Depression
Dizziness
Drug Ineffective
Drug Withdrawal Syndrome
Face Oedema
Fatigue

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FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Movement Disorder Neoplasm Skin Nervous System Disorder	Report Source				
PER ORAL		Suicidal Ideation Urticaria	Consumer	Neurontin (Gabapentin)	PS		ORAL
PER ORAL		Weight Increased		(Sertraline Hydrochloride)	SS		ORAL
				(Carisoprodol) (Prednisone)	C C		

Date:03/05/02ISR Number: 3879990-1Report Type:Expedited (15-DaCompany Report #041-0945-M0200004
Age:89 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG, PER Other ORAL		Agitation Clonic Convulsion Condition Aggravated	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
INTRAVENOUS	2 GM,	Confusional State Encephalopathy Pneumonia	Other	(Cefepime Hydrochloride)	SS		
INTRAVENOUS		Renal Impairment		(Clomethiazole Edisilate) (Lisinopril) (Acetylsalicylic Acid) (Glyceryl Trinitrate) (Molsidomine) (Paracetamol) (Folic Acid) (Torasemide) (Ipratropium Bromide) (Salbutamol)	C C C C C C C C C C		

Date:03/05/02ISR Number: 3879997-4Report Type:Expedited (15-DaCompany Report #002-0945-M0100118
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neuroleptic Malignant Syndrome	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
PER ORAL			Professional	(Haloperidol) (Unspecified Medications)	SS C		

Date:03/05/02ISR Number: 3880071-1Report Type:Expedited (15-DaCompany Report #044-0945-M0200024
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Dysaesthesia	Foreign Health	Neurontin Tablets (Gabapentin)	PS		ORAL
600 MG		Oedema Peripheral	Professional				
(DAILY), PER		Weight Increased					
ORAL				Amitriptyline Baclofen	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/06/02ISR Number: 3879674-XReport Type:Expedited (15-DaCompany Report #A203444

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Convulsion	Consumer	Zoloft Tablets	PS		
Intervention to Prevent Permanent Impairment/Damage		Movement Disorder Muscle Spasms		Neurontin	SS		

Date:03/08/02ISR Number: 3879799-9Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-11743036

Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose		Glucovance Tabs	PS	Bristol-Myers Squibb Company	ORAL
Other unknown							
quantity				Trazodone Hcl Tabs	SS	Apothecon	ORAL
				Neurontin	SS		
				Xanax	SS		
				Darvocet	SS		
				Phenergan + Codeine	SS		

2 oz

Date:03/08/02ISR Number: 3880617-3Report Type:Direct

Company Report #CTU 163055

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Neurontin 100mg, 400 Mg Parke, Davis	PS	Parke, Davis	ORAL
3-100 MG-							
2-400MGX3DAA							
ORAL				Morphine Sulf 30 Mg Endo Labs	SS	Endo Labs	ORAL
MORP- 1-30MG							

X 2 DAY

ORAL

Date:03/08/02ISR Number: 3881482-0Report Type:Expedited (15-DaCompany Report #033-0945-M0200008

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (300 MG, QID), PER ORAL		Anti Factor Viii Antibody Positive	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Coagulation Factor Viii Level Decreased	Professional				
2000 MG (500 MG, QID), PER ORAL		Coagulation Time Prolonged		Paracetamol	SS		ORAL
		Disorientation Subdural Haematoma		Kaskadil	SS		
INTRAVENOUS MILLILITERS (DAILY), INTRAVENOUS	40						
				Heparin-Fraction, Sodium Salt	SS		
SUBCUTANEOUS ML (DAILY), SUBCUTANEOUS	2500 IU/0.2						
20 MG (DAILY), PER ORAL				Omeprazole	SS		ORAL
TRANSDERMAL TRANSDERMAL	5 MG DAILY			Glyceryl Trinitrate	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/08/02ISR Number: 3881491-1Report Type:Expedited (15-DaCompany Report #044-0945-M0200036

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia	Foreign	Neurontin	PS		ORAL
PER ORAL		Inflammation	Consumer	Vigabatrin	C		
		Liver Disorder		Clobazam	C		
		Lymphadenopathy					
		Rash					
		Sarcoidosis					
		Tuberculosis					

Date:03/08/02ISR Number: 3881732-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200282

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Neurontin			
1600 MG (FOUR		Diplopia		(Gabapentin)	PS		
TIMES)				Amfebutamone			
				Hydrochloride	C		
				Alprazolam	C		

Date:03/08/02ISR Number: 3881742-3Report Type:Expedited (15-DaCompany Report #HQ0633311FEB2002

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Coarctation Of The Aorta	Health	Effexor Xr			
Congenital Anomaly		Complications Of Maternal	Professional	(Venlafaxine			
		Exposure To Therapeutic		Hydrochloride,			
		Drugs		Capsule, Extended			
150 MG 1X PER		Congenital Anomaly		Release	PS		ORAL
1 DAY, ORAL		Maternal Drugs Affecting					
		Foetus		Insulin (Insulin,			
		Pregnancy		Injection)	SS		

Premature Baby

Neurontin
(Gabapentin,)

SS

ORAL

300 MG 1X PER

1 DAY, ORAL

Prilosec
(Omeprazole,)

SS

ORAL

20 MG 1X PER

1 DAY, ORAL

Reglan
(Metoclopramide
Hydrochloride,
Tablet)

SS

ORAL

ORAL

Date:03/08/02ISR Number: 3881746-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200283

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma	Consumer	Neurontin			
Other		Drug Level Increased		(Gabapentin)	PS		ORAL
PER ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/08/02ISR Number: 3881986-0Report Type:Expedited (15-DaCompany Report #HQ0633311FEB2002

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Congenital Anomaly		Coarctation Of The Aorta Congenital Anomaly Maternal Drugs Affecting Foetus Pregnancy	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
150 MG 1XPER							
1 DAY, ORAL							
				Insulin (Insulin, Injection) Neurontin (Gabapentin,)	SS SS		ORAL
300 MG 1X PER							
1 DAY, ORAL							
				Prilosec (Omeprazole,)	SS		ORAL
20 MG 2X PER							
1 DAY, ORAL							
				Reglan (Metoclopramide Hydrochloride, Tablet)	SS		ORAL
ORAL							

Date:03/12/02ISR Number: 3882323-8Report Type:Expedited (15-DaCompany Report #044-0945-M0200040

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Dependence	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:03/12/02ISR Number: 3882631-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200218
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (BID),		Anaemia Anorgasmia Condition Aggravated Hypothyroidism Parkinsonism Tardive Dyskinesia Urinary Incontinence	Health Professional	Neurontin (Gabapentin) (Lithium)	PS C		

Date:03/12/02ISR Number: 3882862-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200294
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Neoplasm Malignant	Consumer	Neurontin (Gabapentin)	PS		

Date:03/12/02ISR Number: 3883045-XReport Type:Expedited (15-DaCompany Report #033-0945-M0200021
Age:63 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
Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
PER ORAL		Acute Myeloid Leukaemia Gastrointestinal	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
PER ORAL		Haemorrhage	Professional	Valproate Sodium	SS		ORAL
PER ORAL		Neutrophil Count Abnormal Pancytopenia		Procarbazine Hydrochloride Bromazepam	SS SS		ORAL
				Vincristine Sulfate Lomustine	SS C		

Date:03/13/02ISR Number: 3883227-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200102
Age: Gender:Male I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG (800 MG, TID), PER ORAL		Erectile Dysfunction Skin Operation	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Morniflumate	C		

Date:03/13/02ISR Number: 3883350-7Report Type:Expedited (15-DaCompany Report #039-0945-M020002
Age:81 YR Gender:Female I/FU:F

Outcome Dose Hospitalization - Initial or Prolonged 300 MG (DAILY) PER ORAL	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dermatitis Atopic	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

200 MG
 (DAILY) PER
 ORAL

Celecoxib SS ORAL

Date:03/13/02ISR Number: 3883680-9Report Type:Expedited (15-DaCompany Report #033-0945-M0200012
 Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia	Foreign	Neurontin			
Other		Cardiac Disorder	Consumer	(Gabapentin)	PS		ORAL
800 MG (400		Confusional State	Health				
MG BID) PER		Face Oedema	Professional				
ORAL		Insomnia		(Morphine Sulfate)	SS		ORAL
60 MG (30 MG		Skin Discolouration					
BID) PER ORAL				(Diazepam)	SS		ORAL
5 MG (DAILY)							
PER ORAL				(Tetracosactide)	C		
				(Tetrazepam)	C		
				(Floctafenine)	C		

Date:03/14/02ISR Number: 3882689-9Report Type:Direct Company Report #CTU 163464
 Age:73 YR Gender:Male 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
400MG PO		Somnolence		Gabapentin	PS		ORAL
TID				Multivitamin/Minerals Cap/Tab	C		
				Clopidogrel Bisulfate	C		
				Paroxetine Hcl	C		
				Digoxin (Lanoxin)	C		
				Lisinopril	C		
				Spiroonolactone	C		
				Furosemide	C		
				Rabeprazole Na	C		

Date:03/15/02ISR Number: 3884683-0Report Type:Expedited (15-DaCompany Report #HQ0633311FEB2002
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Congenital Anomaly		Coarctation Of The Aorta Maternal Drugs Affecting Foetus Pregnancy Premature Baby	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
150 MG 1X PER 1 DAY, ORAL				Insulin (Insulin , Injection)	SS		
300 MG 1X PER 1 DAY, ORAL				Neurontin (Gabapentin,)	SS		ORAL
20 MG 2X PER				Prilosec (Omeprazole,)	SS		ORAL

1 DAY, ORAL

Reglan
(Metoclopramide
Hydrochloride,
Tablet)

SS

ORAL

ORAL

Date:03/18/02ISR Number: 3884754-9Report Type:Expedited (15-DaCompany Report #US0201300

Age:11 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Genital Disorder Female	Other	Carbatrol	PS		
SEE NARRATIVE							
SEE NARRATIVE				Neurontin	SS		

Date:03/18/02ISR Number: 3884928-7Report Type:Expedited (15-DaCompany Report #PHRM2001FR02891

Age:37 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Blood Pressure Decreased
Initial or Prolonged	C-Reactive Protein
Other	Increased
	Drug Toxicity
	Ecchymosis
	Hypersensitivity
	Idiopathic Capillaritis

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FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	Medication Error Purpura Pyrexia	Report Source	Product	Role	Manufacturer	Route
400 MG,			Foreign Health Professional	Tegretol Lp(Carbamazepine)	PS		ORAL
ONCE/SINGLE,				Alepsal(Belladonna Extract, Caffeine) Tablet	SS		ORAL
ORAL							
1 DF,				Neurontin(Gabapentin) Tablet	SS		ORAL
ONCE/SINGLE,							
ORAL							
1 DF,				Lepticur (Tropatepine Hydrochloride) Tablet	SS		ORAL
ONCE/SINGLE,							
ORAL							
1 DF,				Urbanyl(Clobazam)	SS		ORAL
ONCE/SINGLE,							
ORAL							
1 DF,				Tercian(Cyamemazine) Tablet	SS		ORAL
ONCE/SINGLE,							

ORAL

Date:03/18/02ISR Number: 3885267-0Report Type:Expedited (15-DaCompany Report #HQ0633311FEB2002

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Congenital Anomaly		Coarctation Of The Aorta Congenital Anomaly Pregnancy Premature Baby	Health Professional	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
150 MG 1X PER							
1 DAY, ORAL							
				Insulin (Insulin, Injection)	SS		
300 MG 1X PER				Neurontin (Gabapentin,)	SS		ORAL
1 DAY, ORAL							
				Prilosec (Omeprazole,)	SS		ORAL
20 MG 2X PER							
1 DAY, ORAL							
				Reglan (Metoclopramide Hydrochloride, Tablet)	SS		ORAL

ORAL

Date:03/19/02ISR Number: 3885422-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200295

Age: Gender: I/FU:I

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
		Arrhythmia	Health	Neurontin			
		Post Procedural Haematoma	Professional	(Gabapentin)	PS		
		Pulmonary Embolism	Company	Fosphenytoin Sodium	SS		
		Subdural Haematoma	Representative	Metronidazole	SS		
		Trigeminal Neuralgia		Carbamazepine	SS		
				Clavulanate			
				Potassium, Amoxicillin			
				Trihydrate	SS		

Date:03/19/02ISR Number: 3885429-2Report Type:Expedited (15-DaCompany Report #001-0945-M020097
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia	Health	Neurontin			
Hospitalization -		Post Procedural	Professional	(Gabapentin)	PS		
Initial or Prolonged		Complication	Company	Fosphenytoin Sodium	SS		
Other		Pulmonary Embolism	Representative	Metronidazole	SS		
		Subdural Haematoma		Carbamazepine	SS		
				Clavulanate			
				Potassium, Amoxicillin			
				Trihydrate	SS		

Date:03/19/02ISR Number: 3885431-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200033
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cataract	Consumer	Neurontin			
Other		Constipation		(Gabapentin)	PS		
700 MG		Drug Withdrawal Syndrome					
(DAILY)		Eye Pain					
		Vision Blurred					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Prostatic Specific	Health	Neurontin			
600 MG		Antigen Increased	Professional	(Gabapentin)	PS		ORAL
(DAILY), PER							
ORAL							
				Acetylsalicylic Acid	C		
				Ergocalciferol,			
				Ascorbic Acid, Folic			
				Acid, Retinol,			
				Riboflavin,			
				Nicotinamide	C		
				Tamsulosin			
				Hydrochloride	C		
				Fenofibrate	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/19/02ISR Number: 3885466-8Report Type:Expedited (15-DaCompany Report #045-0945-M0200006

Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Apnoea	Foreign	Gabapentin	PS		ORAL
600 MG (BID), Other PER ORAL		Blood Creatinine	Health				
		Increased Blood Lactate	Professional	Donepezil Hydrochloride	SS		ORAL
10 MG, PER ORAL		Dehydrogenase Increased					
		C-Reactive Protein Increased		Sertraline	C		
		Cardiac Arrest		Diazepam	C		
		Circulatory Collapse		Risperidone	C		
		Convulsion		Mianserin	C		
		Dizziness					
		Haemoglobin Decreased					
		Heart Rate Increased					
		Hypotension					
		Malaise					
		Myocardial Infarction					
		Pallor					
		Pulse Absent					
		Somnolence					

Date:03/19/02ISR Number: 3885539-XReport Type:Expedited (15-DaCompany Report #NSADSS2002007512

Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Dependence	Consumer	Duragesic (75 Mcg/Hr Patch) (Fentanyl)	PS		
Hospitalization - TRANSDERMAL	75 MCG/H, 1						
Initial or Prolonged IN 48 HOUR(S), TRANSD				Neurontin			

3000 , 3 IN	(Gabapentin)	SS	ORAL
24 HOUR(S),			
ORAL			
100, 1 IN 24	Trazodone (Trazadone)	SS	ORAL
HOUR(S), ORAL			
100 MG, 2 IN	Seroquel (Seroquel)	SS	ORAL
24 HOUR(S),			
ORAL			
ORAL	Effexor (Venlafaxine Hydrochloride)	SS	ORAL
	Pepcid (Famotidine)	C	
	Darvocet (Carvocet)	C	
	Ambien (Zolpidem Tartrate)	C	
	Zofran (Ondansetron Hydrochloride)	C	
	Provera (Medroxyprogesterone Acetate)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/19/02ISR Number: 3885728-4Report Type:Expedited (15-DaCompany Report #002#4#2002-00037 (0)

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Congenital Anomaly ORAL		Coarctation Of The Aorta Congenital Anomaly	Health Professional	Reglan-Dose Unknown (Metoclopramide Hcl)	PS		ORAL
150 MG, 1 IN		Maternal Drugs Affecting Foetus	Other	Venlafaxine	SS		ORAL
1 D, ORAL		Pregnancy		Gabapentin	SS		ORAL
300 MG, 1 IN		Premature Baby		Omeprazole	SS		ORAL
1 D, ORAL				Insulin	SS		
20 MG, 2 IN 1							
D, ORAL							

Date:03/19/02ISR Number: 3885922-2Report Type:Expedited (15-DaCompany Report #061-0945-M0200026

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident Cognitive Deterioration	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600		Coordination Abnormal	Professional				
MG, TID), PER		Difficulty In Walking	Company				
ORAL		Disturbance In Attention Memory Impairment Mental Impairment Sedation	Representative	Morphine Sulfate Atorvastatin Valproate Sodium Omeprazole Ramipril Docusate Sodium	C C C C C C		

Date:03/19/02ISR Number: 3885930-1Report Type:Expedited (15-DaCompany Report #049-0945-M0200004

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Abdominal Pain Upper Difficulty In Walking	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG, PER		Disease Progression					
ORAL		Fibromyalgia Joint Stiffness Migraine Muscle Spasms Polyarthritits Sleep Disorder					

Date:03/19/02ISR Number: 3885995-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200198
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300		Accidental Overdose	Health Professional	Neurontin (Gabapentin)	PS		ORAL
MG, TWICE A							
DAY), PER							
ORAL				(Sertraline Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/19/02ISR Number: 3886002-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200170

Age:85 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG Other (DAILY), PER ORAL PER ORAL	Convulsion Diabetes Mellitus Drug Toxicity Laboratory Test Abnormal Orthostatic Hypotension Urinary Tract Infection	Health Professional	Neurontin (Gabapentin) (Phenytoin) (Sulfamethoxazole, Trimethoprim)	PS SS SS		ORAL ORAL

Date:03/19/02ISR Number: 3886010-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200292

Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged PER ORAL	Confusional State Dizziness Fall Haematoma Somnolence	Health Professional	Neurontin (Gabapentin) Hydralazine Isosorbide Dinitrate Verapamil Trazodone Diphenhydramine Hydrochloride Folic Acid, Vitamins Nos Calcium Acetate Sodium Bicarbonate Lorazepam Morphine Paracetamol Warfarin Sodium	PS C C C C C C C C C C C C C C C		ORAL

Date:03/19/02ISR Number: 3886011-3Report Type:Expedited (15-DaCompany Report #033-0945-M0200025

Age: Gender: I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Hospitalization -		Liver Function Test	Foreign	Neurontin			
Initial or Prolonged		Abnormal	Health	(Gabapentin)	PS		
			Professional				
			Company				
			Representative				

Date:03/19/02ISR Number: 3886023-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200283
 Age:43 YR Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Coma	Consumer	Neurontin			
Other		Fall		(Gabapentin)	PS		ORAL
PER ORAL							

Date:03/19/02ISR Number: 3886032-0Report Type:Expedited (15-DaCompany Report #001-0719-M0100408
 Age: Gender:Male I/FU:F

Outcome		PT
Hospitalization -		Alanine Aminotransferase
Initial or Prolonged		Decreased
Other		Arteriosclerosis
		Arthralgia

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Urinary Incontinence
 Urinary Tract Infection
 Urine Ketone Body Present
 Varicose Vein
 Vasodilatation
 Very Low Density
 Lipoprotein Increased
 Vision Blurred

Date:03/19/02ISR Number: 3886091-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200293

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Thyroid Disorder	Consumer	Neurontin			
Initial or Prolonged	Weight Increased		(Gabapentin)	PS		
10 MG (5 MG,			Olanzapine	SS		
TWICE A DAY)			Dicycloverine			
			Hydrochloride	C		
			Folic Acid	C		
			Docusate Sodium			
			Casanthranol	C		
			Levothyroxine	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lisinopril	C
Multivitamins With	
Minerals	C
Thiamine	C
Verapamil	C
Verapamil	C
Benzatropine	
Mesilate	C
Loxapine	C
Bisacodyl	C
Lorazepam	C

Date:03/19/02ISR Number: 3886147-7Report Type:Expedited (15-DaCompany Report #033-0945-M0200027
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Hypomania	Foreign	Neurontin			
Initial or Prolonged		Health	(Gabapentin)	PS		
UNKNOWN	900 MG (TID),					
Other		Professional				
UNKNOWN			(Unspecified Anesthetic Drug)	SS		
UNKNOWN						

Date:03/20/02ISR Number: 3885857-5Report Type:Expedited (15-DaCompany Report #044-0945-M0200039
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Dysphagia	Foreign	Neurontin			
Initial or Prolonged	Neurological Examination	Consumer	(Gabapentin)	PS		
Other	Abnormal		(Cimetidine)	SS		
INTRAVENOUS	INTRAVENOUS					
	Tremor		(Salbutamol)	C		
			(Beclometasone)	C		
			(Fluticasone			
			Propionate)	C		
			(Lansoprazole)	C		
			(Tegremet)	C		
			(Clopidogrel)	C		
			(Fluvastatin Sodium)	C		

Date:03/21/02ISR Number: 3887005-4Report Type:Expedited (15-DaCompany Report #020-0945-M0200002

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 800 MG (BID), PER ORAL		Road Traffic Accident Somnolence	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:03/21/02ISR Number: 3887097-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200328

Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Other	Asthma Blood Potassium Decreased Cough Drug Ineffective Dyspnoea

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Haematoma Headache Hypokalaemia	Consumer	Neurontin (Gabapentin)	PS		ORAL
PER ORAL		Increased Appetite Muscle Spasms	Consumer	Quinine Carbamazepine	SS		ORAL
200 MG DAILY,		Petechiae Stevens-Johnson Syndrome		Levothyroxine Sodium	C		
PER ORAL		Weight Increased Wheezing					

Date:03/21/02ISR Number: 3887098-4Report Type:Expedited (15-DaCompany Report #001-0073-M0200120
Age: Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bilirubin Conjugated Increased	Health Professional	Phenytoin (Phenytoin)	PS		
Hospitalization - 400 MG (200 Initial or Prolonged MG, BID)		Blood Bilirubin Increased		Carbamazepine Gabapentin	SS SS		
Other		Drug Toxicity Encephalopathy Hepatocellular Damage Pyrexia					

Date:03/21/02ISR Number: 3887180-1Report Type:Expedited (15-DaCompany Report #039-0945-M0200005
Age: Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression Cognitive Disorder	Foreign Literature	Gabapentin (Gabapentin)	PS		
1200 MG, DAILY		Condition Aggravated	Consumer	Donepezil Hydrochloride	SS		
5 MG, DAILY		Confusional State Dementia Memory Impairment					

Mental Disorder
Nervous System Disorder
Paranoia

Date:03/21/02ISR Number: 3887182-5Report Type:Expedited (15-DaCompany Report #039-0945-M0200004

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression Cognitive Disorder	Foreign Literature	Gabapentin (Gabapentin)	PS		
900 MG, DAILY		Condition Aggravated Hallucination, Visual	Consumer	Donepezil Hydrochloride	SS		
5 MG, DAILY		Hypersomnia Ideas Of Reference Pneumonia					

Date:03/21/02ISR Number: 3887311-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200310

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Intervertebral Disc Operation	Consumer	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/21/02ISR Number: 3887355-1Report Type:Expedited (15-DaCompany Report #001-0073-M0200128

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Change Of Bowel Habit	Consumer	Dilantin (Phenytoin Sodium)	PS		ORAL
PER ORAL		Dyspnoea					
		Faecal Incontinence		Neurontin (Gabapentin)	SS		ORAL
3000 MG							
(DAILY), PER							
ORAL							

Date:03/21/02ISR Number: 3887357-5Report Type:Expedited (15-DaCompany Report #001-0945-M0101373

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blepharospasm	Consumer	Neurontin (Gabapentin)	PS		ORAL
Initial or Prolonged		Cerebrovascular Accident	Health				
1200 MG							
Disability		Fall	Professional				
(DAILY), PER							
ORAL		Muscle Spasms					

Date:03/21/02ISR Number: 3887371-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200313

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (TID),		Drug Ineffective					
PER ORAL		Infection					
		Prostatic Specific		Serenoa Repens	C		
		Antigen Increased		Flax Seed Oil	C		
		Trigeminal Nerve Disorder		"Gugul"	C		
				Omega 2,6 Fish Oil	C		

Date:03/22/02ISR Number: 3887960-2Report Type:Expedited (15-DaCompany Report #02P-056-0189541-00
 Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acute Myeloid Leukaemia	Foreign	Depakine Tablets			
Hospitalization -		Anisocytosis	Health	(Sodium Valproate)			
Initial or Prolonged		Gastrointestinal	Professional	(Sodium Valproate)			
PER ORAL		Haemorrhage	Other	(Sodium Valproate)	PS		ORAL
200 MG, 1 IN		Pancytopenia		Lomustine	SS		
1 D				Procarbazine			
100 MG, EVEN				Hydrochloride	SS		
DAYS; 50 MG,							
ON ODD DAYS;							
100 MG, ON							
EVEN DAYS				Vincristine Sulfate	SS		
2 MG, 1 IN 1							
D; 1 MG, 1 IN							
1 D				Bromazepam	SS		ORAL
PER ORAL				Gabapentin	SS		ORAL
2 UNIT, 1 IN							
1 D, PER ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/25/02ISR Number: 3888682-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200233

Age:51 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged PER ORAL	Condition Aggravated Decreased Appetite	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
	Dizziness Drug Withdrawal Syndrome Insomnia Stress Tremor Trigeminal Neuralgia Weight Decreased	Professional	(Medroxyprogesterone Acetate, Estrogens Conjugated) (Carbamazepine)	C C		

Date:03/25/02ISR Number: 3888684-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200098

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2700 MG (DAILY),	Feeling Abnormal Grand Mal Convulsion	Health Professional	Neurontin (Gabapentin)	PS		
	Medication Error Muscle Twitching		(Fluoxetine Hydrochloride) (Zolpidem Tartrate) (Oxycodone Hydrochloride)	C C C		

Date:03/25/02ISR Number: 3888786-6Report Type:Expedited (15-DaCompany Report #033-0945-M0200027

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG, TID Other	Hypomania	Foreign Health	Neurontin (Gabapentin)	PS		
		Professional	Unspecified Anesthetic Drug	SS		

Date:03/25/02ISR Number: 3888870-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200326
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Electrocardiogram Qt	Health	Neurontin			
Initial or Prolonged	Prolonged	Professional	(Gabapentin)	PS		
41600 MG						
Other	Overdose		Lamotrigine	C		
			Reperidal	C		

Date:03/25/02ISR Number: 3888872-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200325
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Electrocardiogram Qt	Health	Neurontin			
Initial or Prolonged	Prolonged	Professional	(Gabapentin)	PS		
Other	Overdose		Tricyclic			
			Antidepressant	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/25/02ISR Number: 3898845-XReport Type:Periodic
Age:67 YR Gender:Male I/FU:F

Company Report #01RUSWE0100245

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation Condition Aggravated	Health Professional	Welchol (Colesevelam Hcl) Tablet, 625 Mg	PS		ORAL
ORAL		Drug Interaction	Company Representative	Neurontin (Gabapentin)	SS		ORAL
ORAL							

Date:03/26/02ISR Number: 3889916-2Report Type:Expedited (15-DaCompany Report #2002CG00412
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hospitalization - 20 MG QD PO	Foreign	Mopral	PS		ORAL
Initial or Prolonged		Haemoglobin Decreased	Health	Lasilix	SS		ORAL
40 MG QD PO		Thrombocytopenia	Professional	Neurontin	SS		ORAL
500 MG QD PO		White Blood Cell Count Decreased	Other	Augmentin	SS		
1G+200 MG							

Date:03/27/02ISR Number: 3890351-1Report Type:Direct
Age:50 YR Gender:Male I/FU:I

Company Report #CTU 164291

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hospitalization - Initial or Prolonged		Neurontin 300 Mg Parke-Davis	PS	Parke-Davis	ORAL
1T TID PO		Pancreatitis					

Date:03/27/02ISR Number: 3890949-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200345
Age:92 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Confusional State Depressed Level Of Consciousness	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), PER ORAL		Feeling Abnormal Haemorrhage Intracranial Hemianopia Hypersomnia Mobility Decreased Nervous System Disorder Nuclear Magnetic Resonance Imaging Abnormal Somnolence Visual Disturbance		(Lisinopril) (Rosiglitazone) (Metoprolol Tartrate) (Furosemide) (Magnesium Oxide)	C C C C C		

Date:03/28/02ISR Number: 3891930-8Report Type:Direct Company Report #CTU 164499
Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 400 PO / Initial or Prolonged LIGUID		Respiratory Depression		Gabapentin	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/02ISR Number: 3892342-3Report Type:Expedited (15-DaCompany Report #002-0945-M0200023

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Adrenocortical	Foreign	Gabapentin			
PER ORAL		Insufficiency Chronic	Health	(Gabapentin)	PS		ORAL
		Blood Pressure Decreased	Professional				
		Syncope					

Date:03/29/02ISR Number: 3892369-1Report Type:Expedited (15-DaCompany Report #049-0945-M0200039

Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Antibody Test Positive	Foreign	Neurontin			
Initial or Prolonged		Haematoma	Health	(Gabapentin)	PS		ORAL
900 MG		Thrombocytopenia	Professional				
(DAILY), PER			Company				
ORAL			Representative				

Date:03/29/02ISR Number: 3892466-0Report Type:Expedited (15-DaCompany Report #033-0945-M0200031

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Angioneurotic Oedema	Foreign	Neurontin			
Life-Threatening		Aphasia	Health	(Gabapentin)	PS		ORAL
300 MG (100		Chest Pain	Professional				
Hospitalization -		Respiratory Distress					
MG, TID), PER		Torsade De Pointes		(Carbocisteine)	SS		ORAL
Initial or Prolonged		Ventricular Fibrillation		Ibuprofen	C		
ORAL							
PER ORAL							

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2400 MG (DAILY), PER ORAL	Clonic Convulsion Fall Nervous System Disorder Somnolence	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Outcome	PT
Hospitalization - Initial or Prolonged Other	Abdominal Pain Acral Lentiginous Melanoma Stage Unspecified Animal Scratch Arthralgia Arthritis Asthenia Atrial Hypertrophy Back Injury Back Pain Bacteria Urine Identified Benign Prostatic Hyperplasia Blood Cholesterol

Freedom Of Information (FOI) Report

Increased
Blood Glucose Increased
Blood Sodium Increased
Blood Triglycerides
Increased
Blood Urine Present
Brain Scan Abnormal
Cardiovascular Disorder
Carotid Artery Stenosis
Cellulitis
Cerebral Atrophy
Cerebrovascular Accident
Chest Pain
Condition Aggravated
Cystitis
Diabetes Mellitus
Difficulty In Walking
Dilatation Ventricular
Disorientation
Dizziness
Drug Ineffective
Dry Mouth
Dysarthria
Electrocardiogram
Abnormal
Erythema
Eye Disorder
Eye Swelling
Face Oedema
Haematuria
Headache
Hepatic Steatosis
Hyperlipidaemia
Hypertension
Hypoaesthesia
Intervertebral Disc
Protrusion
Ischaemia
Limb Injury
Micturition Urgency
Muscle Disorder
Myalgia
Myocardial Ischaemia
Neuropathy Peripheral
Nocturia
Oedema Peripheral
Osteoarthritis
Pain

Pain In Extremity
Pco2 Increased
Pharyngitis
Phlebitis
Pitting Oedema
Pollakiuria
Primary
Hyperaldosteronism
Pruritus
Pyrexia
Radiculopathy
Rash
Sinusitis
Skin Discolouration

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Sleep Apnoea Syndrome Spinal Column Stenosis Spinal Osteoarthritis	Consumer				
1200 MG		Stasis Dermatitis	Health	Lopid (Gemfibrozil)	PS		ORAL
(BID), PER		Urinary Incontinence	Professional				
ORAL		Urinary Retention					
PER ORAL		Urinary Tract Infection		Gabapentin	SS		ORAL
5 MG (DAILY),		Urine Flow Decreased		Amlodipine Besilate	SS		ORAL
PER ORAL		Urine Ketone Body Present					
PER ORAL		Urticaria		Glipizide	SS		ORAL
SUBLINGUAL	10 MG,	Varicose Vein		Nifedipine	SS		
SUBLINGUAL		Very Low Density					
		Lipoprotein Increased		Cholesterol Lowering			
		Vision Blurred		Drug	SS		
		Weight Increased		Furosemide	C		
		White Blood Cell Disorder		Acetylsalicylic Acid	C		
		White Blood Cells Urine		Ergocalciferol,			
		Positive		Ascorbic Acid, Folic			
				Acid, Thiamine			
				Hydrochloride,			
				Retinol, Riboflavin,	C		

Date:03/29/02ISR Number: 3892663-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200359
Age:35 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Contusion	Consumer	Neurontin			
900 (300,		Platelet Count Decreased		(Gabapentin)	PS		ORAL
TID) PER ORAL							
150 MG				(Venlafaxine			
				Hydrochloride)	SS		ORAL

(DAILY), PER

ORAL

Date:03/29/02ISR Number: 3892671-3Report Type:Expedited (15-DaCompany Report #001-0073-M0200138

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Dizziness	Consumer	Dilantin (Phenytoin Sodium)	PS		ORAL
500 MG		Drug Withdrawal Syndrome					
(DAILY), PER		Fatigue					
ORAL		Feeling Abnormal		Gabapentin	SS		ORAL
900 MG (TID),		Gingival Disorder					
PER ORAL		Gingival Hyperplasia Hypersomnia Insomnia Myalgia Psychomotor Hyperactivity Staring					

Date:03/29/02ISR Number: 3892754-8Report Type:Direct

Company Report #CTU 164639

Age:55 YR Gender:Female I/FU:I

Outcome
Other
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
		Pharmaceutical Product Complaint		Neurontin All Strengths Parke-Davis, Of Pfizer	PS	Park-Davis/Pfizer	ORAL
3X/DAY TABS							
CAPS ORAL							

Date:03/29/02ISR Number: 3892765-2Report Type:Expedited (15-DaCompany Report #001-0719-M0100407
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Distension
Initial or Prolonged	Abdominal Mass
Disability	Abdominal Pain Upper
Other	Angina Pectoris
	Anhedonia
	Aphasia
	Areflexia
	Arteriosclerosis
	Arthralgia
	Arthritis
	Asthenia
	Back Pain
	Benign Prostatic
	Hyperplasia
	Blindness
	Blood Albumin Decreased
	Bone Pain
	Cerebrovascular Accident
	Chest Pain
	Cholelithiasis
	Condition Aggravated
	Condyloma Acuminatum
	Coronary Artery
	Atherosclerosis
	Decreased Activity
	Depression
	Diabetes Mellitus
	Dizziness
	Drug Interaction

Dyspepsia
Ear Pain
Emphysema
Extrasystoles
Fatigue
Feeling Abnormal
Food Poisoning
Gastritis
Haematocrit Decreased
Haemoglobin Decreased
Headache
Hearing Impaired
Herpes Zoster
Hyperlipidaemia
Hypertension
Hypothyroidism
Insomnia
Intermittent Claudication

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1200 MG (600 MG, BID) PER ORAL 1600 MG (400 MG 2 CAPSULES IN AM ONE EVERY 5 MIN., MAX: 3 0.4 MG, DAILY	Joint Swelling	Consumer Health Professional	Lopid (Gemfibrozil)	PS		ORAL
	Lumbar Spinal Stenosis					
	Lymphocyte Count					
	Decreased					
	Macular Degeneration					
	Memory Impairment					
	Migraine		Gabapentin	SS		
	Myocardial Infarction					
	Myocardial Ischaemia					
	Nausea			Glyceryl Trinitrate	SS	
Neuralgia						
Neuropathy Peripheral			Cerivastatin Sodium	SS		
Neutrophil Count						
Increased						
Oedema Peripheral						
Osteoarthritis						
Pain In Extremity						
Palpitations						
Peptic Ulcer						
Peripheral Vascular Disorder						
Pharyngolaryngeal Pain						
Productive Cough						
Pruritus						
Pyrexia						
Red Blood Cell Count						
Decreased						
Rhabdomyolysis						
Sensory Disturbance						
Sinusitis						
Speech Disorder						
Supraventricular Extrasystoles						
Telangiectasia						
Tenderness						
Tinnitus						
Tongue Neoplasm Malignant						

Stage Unspecified
Tooth Abscess
Tremor
Upper Respiratory Tract
Infection
Ventricular Extrasystoles
Visual Acuity Reduced
Vomiting

Date:04/01/02ISR Number: 3892559-8Report Type:Expedited (15-DaCompany Report #033-0945-M0200035
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Epilepsy	Foreign Consumer	Neurontin (Gabapentin)	PS		
2200 MG (DAILY),							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/01/02ISR Number: 3892817-7Report Type:Direct
Age:49 YR Gender:Female I/FU:I

Company Report #CTU 164643

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required	2AM AND 3 HS	Coordination Abnormal		Neurontin 400mg	PS		
Intervention to Prevent Permanent Impairment/Damage		Diplopia		Lamictal	C		
		Disorientation		Seroquel	C		
		Tremor		Prozac	C		
				Synthroid	C		
				Verapamil	C		
				Glucophage	C		

Date:04/02/02ISR Number: 3893800-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200369
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	900 MG (300 MG, THREE TIMES A DAY, PER ORAL	Muscle Spasms Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Weight Decreased		Oxycodone Hydrochloride	C		
				Levothyroxine Sodium	C		
				Glipizide	C		
				Insulin	C		
				Atorvastatin	C		

Date:04/02/02ISR Number: 3893866-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200370
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	600 MG	Arterial Disorder Arteriosclerosis	Consumer	Neurontin (Gabapentin)	PS		ORAL

(DAILY), PER
 ORAL
 Calcinosi
 Inguinal Hernia
 Intervertebral Disc Degeneration
 Intervertebral Disc Protrusion
 Prostatic Disorder
 Scoliosis
 Unspecified Narcotics
 Diazepam

Date:04/02/02ISR Number: 3893867-7Report Type:Expedited (15-DaCompany Report #001-0945-M0101466
 Age:17 YR Gender:Male I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG		Agranulocytosis Fatigue	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Feeling Cold					
		Platelet Count Decreased					
15 MG		White Blood Cell Count Decreased		Mirtazapine	SS		ORAL
				Risperidone Venlafaxine Hydrochloride Cetirizine Hydrochloride	C C C C		

Freedom Of Information (FOI) Report

Salbutamol C

Date:04/02/02ISR Number: 3893870-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200361
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other (TWICE DAILY), PER ORAL		Benign Ovarian Tumour Memory Impairment	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Captopril Furosemide Glyceryl Trinitrate Hydrocone	C C C C		

Date:04/02/02ISR Number: 3894002-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200368
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG BID, PER ORAL 2 TABLETS (DAILY)		Bladder Spasm Condition Aggravated Faeces Discoloured Gastrointestinal Haemorrhage Haematocrit Decreased Iron Deficiency Anaemia	Consumer	Neurontin (Gabapentin) (Iron) (Acebutolol Hydrochloride) (Amlodipine Besilate) (Levothyroxine Sodium) (Omeprazole) (Hydrochlorothiazide , Triamterene) (Trandolapril)	PS SS C C C C C C		ORAL

Date:04/02/02ISR Number: 3894004-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200371
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged PER ORAL	Drug Effect Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:04/02/02ISR Number: 3894078-1Report Type:Expedited (15-DaCompany Report #049-0945-M0200040
Age:88 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death Hospitalization - 300 MG Initial or Prolonged (DAILY), PER Other ORAL	Cardiac Failure Condition Aggravated Dizziness Fall Head Injury Shock Subdural Haematoma	Foreign Health Professional	Neurontin (Gabapentin) Phenprocoumon	PS C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/02/02ISR Number: 3894765-5Report Type:Expedited (15-DaCompany Report #044-0945-M0200031

Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	1200 MG (300 MG, QID), PER ORAL	Ecchymosis	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				(Warfarin)	C		
				(Amiodarone)	C		
				(Furosemide,Amiloride Hydrochloride)	C		

Date:04/03/02ISR Number: 3893509-0Report Type:Expedited (15-DaCompany Report #B0263843A

Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hepatitis Cholestatic Pancreatic Disorder Rash Maculo-Papular		Combivir Kaletra Neurontin Anti Toxoplasmosis Therapy	PS SS SS C	Glaxo Wellcome	ORAL ORAL ORAL

UNKNOWN

Date:04/03/02ISR Number: 3894514-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200295

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 2400 MG (600 MG, QID) Other 400 MG (100 MG, Q68		Arrhythmia Cardiac Arrest Coordination Abnormal Deep Vein Thrombosis Fall	Health Professional Company Representative	Neurontin (Gabapentin)	PS		
				Fosphenytoin Sodium)	SS		

800 MG (200
 MG, QID)
 Peripheral Embolism (Carbamazepine) SS
 Post Procedural Haematoma
 Pulmonary Embolism (Clavulanate
 Pyrexia Potassium,
 Subdural Haematoma Amoxicillin
 Vomiting Trihydrate) SS
 (Metronidazole) C

2000 MG (500
 MG, QID)

Date:04/03/02ISR Number: 3895459-2Report Type:Expedited (15-DaCompany Report #033-0945-M0200035
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Epilepsy	Foreign	Neurontin			
		Loss Of Consciousness	Consumer	(Gabapentin)	PS		

2200 MG
 (DAILY),
 UNKNOWN

Date:04/05/02ISR Number: 3896107-8Report Type:Expedited (15-DaCompany Report #A204838
 Age:44 YR Gender:Female 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
INTRAVENOUS	400.00	Abdominal Distension MG; Agitation	Consumer	Diffucan Injection	PS		
TOTAL;DAILY;I							
NTRAVENOUS		Ankle Fracture					
INTRAVENOUS	400.00	Atrial Fibrillation MG		Phenytoin Sodium	SS		
TOTAL;QID;INT		Bladder Disorder					
RAVENOUS		Blood Cholesterol					
900.00 MG		Increased		Gabapentin	SS		
TOTAL;TID		Blood Glucose Increased					
		Blood Triglycerides		Mannitol	C		
		Increased		Dexamethasone	C		
		Bradycardia		Famotidine	C		
		Carotid Artery Aneurysm		Nitroprusside Sodium	C		
		Cerebral Haemorrhage		Atropine	C		
		Coma		Pseudoephedrine			
		Confusional State		Hydrochloride	C		
		Constipation		Potassium Chloride	C		
		Convulsion		Vancomycin	C		
		Depression		Cefotaxime Sodium	C		
		Disorientation		Fresh Frozen Plasma	C		
		Dizziness Postural		Dopamine	C		
		Eye Pain		Pentobarbital	C		
		Face Oedema		Metoclopramide	C		
		Fall		Nystatin	C		
		Fatigue		Morphine	C		
		Flat Affect		Haloperidol	C		
		Haematoma					
		Headache					
		Hemiparesis					
		Hypertonic Bladder					
		Hypothyroidism					
		Insomnia					
		Laceration					
		Lethargy					
		Memory Impairment					
		Mental Status Changes					

Micturition Urgency
Obsessive-Compulsive
Disorder
Oligomenorrhoea
Pharyngitis
Pharyngolaryngeal Pain
Pollakiuria
Seborrheic Dermatitis
Soft Tissue Injury
Speech Disorder
Urinary Retention
Urinary Tract Infection
Weight Increased
White Blood Cell Count
Increased

Date:04/05/02ISR Number: 3896678-1Report Type:Expedited (15-DaCompany Report #055-0945-M0200024
Age:72 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Cardio-Respiratory Arrest
Initial or Prolonged	Coordination Abnormal

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema Visual Disturbance	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
PER ORAL				Pantoprazole	C		
				Folic Acid + Ferrous Hydroxide	C		
				Amitriptyline	C		

Date:04/05/02ISR Number: 3896779-8Report Type:Expedited (15-DaCompany Report #A202900
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 200.00 MG		Amenorrhoea	Health	Zoloft Tablets	PS		ORAL
Intervention to TOTAL;DAILY;O		Anxiety	Professional				
Prevent Permanent RAL		Depression					
Impairment/Damage 1600.00 MG		Mood Altered		Neurontin	SS		ORAL
TOTAL;BID;ORA		Neutrophil Count					
L		Decreased					
2.00 MG		Red Blood Cell Count		Risperdal	SS		
TOTAL;DAILY		Decreased					
		Tearfulness		Xanax	C		
		White Blood Cell Count		Depakote	C		
		Decreased		Benadryl	C		

Date:04/05/02ISR Number: 3897053-6Report Type:Expedited (15-DaCompany Report #001-0073-M0200107
Age:44 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Abdominal Distension Agitation

Disability Alanine Aminotransferase
Other Increased
Amnesia
Anaemia
Anger
Ankle Fracture
Arthralgia
Aspartate
Aminotransferase
Increased
Asthenia
Atelectasis
Bladder Disorder
Blood Cholesterol
Increased
Blood Fibrinogen
Increased
Blood Glucose Increased
Blood Lactate
Dehydrogenase Increased
Blood Osmolarity Abnormal
Blood Sodium Decreased
Blood Triglycerides
Increased
Carotid Aneurysm Rupture
Carotid Artery Aneurysm
Catheter Related

Freedom Of Information (FOI) Report

Infection
Chest X-Ray Abnormal
Constipation
Contusion
Coordination Abnormal
Crying
Depression
Dizziness Postural
Drug Hypersensitivity
Drug Level Increased
Dysphagia
Electrocardiogram T Wave
Abnormal
Encephalomalacia
Face Oedema
Fall
Flat Affect
Fungal Infection
Gamma-Glutamyltransferase
Increased
Granuloma
Hallucination, Visual
Head Injury
Headache Postoperative
Hemiplegia
Hyperlipidaemia
Hyperphagia
Hypersensitivity
Hypothyroidism
Iiird Nerve Paralysis
Intracranial Pressure
Increased
Joint Dislocation
Laceration
Lethargy
Memory Impairment
Obsessive-Compulsive
Disorder
Oesophagitis
Oligomenorrhoea
Optic Ischaemic
Neuropathy
Pneumonia Klebsiella
Pollakiuria
Prothrombin Level
Increased
Psychotic Disorder
Rash Pruritic

Respiratory Failure
Seborrhoeic Dermatitis
Sputum Culture Positive
Staphylococcal
Bacteraemia
Staphylococcal Infection
Strabismus
Tachycardia
Thinking Abnormal
Tracheobronchitis
Urge Incontinence
Urinary Tract Infection
Vaginal Mycosis
Visual Field Defect

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Weight Increased White Blood Cell Count Abnormal	Report Source	Product	Role	Manufacturer	Route
INTRAVENOUS	400 MG (100		Consumer	Dilantin (Phenytoin Sodium)	PS		
MG, Q6H),							
INTRAVENOUS				Gabapentin	SS		
900 MG (300							
MG, TID) PER							
TUBE				Fluconazole	SS		
INTRAVENOUS	400 MG						
(DAILY)							
INTRAVENOUS				Dexamethasone	C		
				Famotidine	C		
				Nitroprusside Sodium	C		
				Haloperidol	C		
				Nimodipine	C		
				D5 1/2 Ns With 20			
				Meq Kcl	C		
				Morphine Sulfate	C		
				Pancuronium Bromide	C		
				Mannitol	C		

Date:04/08/02ISR Number: 3896775-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200387
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Dizziness	Consumer	Neurontin			
Initial or Prolonged		Fatigue		(Gabapentin)	PS		ORAL
PER ORAL				Lamotrigine	SS		

Date:04/08/02ISR Number: 3896809-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200408
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bronchitis	Consumer	Neurontin			
300 MG (100		Drug Effect Decreased		(Gabapentin)	PS		ORAL
MG, TID), PER		Intervertebral Disc					
ORAL		Disorder					
		Renal Disorder		(Citalopram			
				Hydrobromide)	C		
				(Paracetamol,			
				Hydrocodone			
				Bitartrate)	C		
				(Lorazepam)	C		

Date:04/08/02ISR Number: 3896816-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200394
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cardiac Disorder	Consumer	Neurontin			
Initial or Prolonged		Chest Pain		(Gabapentin)	PS		ORAL
900 MG (TID),		Mitral Valve Incompetence					
PER ORAL		Swelling		(Folic Acid,			
				Vitamins)	C		
				(Ascorbic Acid)	C		
				(Ferrous Sulfate)	C		

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(Calcium Acetate) C
 Vitamins C

Date:04/08/02ISR Number: 3896818-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200393
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dialysis	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), PER ORAL							

(Folic Acid,
 Vitamins) C
 (Ascorbic Acid) C
 (Ferrous Sulfate) C
 (Calcium Acetate) C

Date:04/08/02ISR Number: 3896822-6Report Type:Expedited (15-DaCompany Report #001-0719-M0100408
 Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Alanine Aminotransferase
Other	Abnormal
	Angiopathy
	Animal Scratch
	Arterial Disorder
	Arteriosclerosis
	Arthralgia
	Asthenia
	Atrial Hypertrophy
	Back Injury
	Back Pain
	Bacteria Urine Identified
	Benign Prostatic
	Hyperplasia
	Bladder Disorder
	Blood Bilirubin Increased
	Blood Cholesterol
	Increased
	Blood Sodium Increased

Blood Triglycerides
Increased
Blood Urea Increased
Blood Urine Present
Brain Scan Abnormal
Cardiac Disorder
Cardiovascular Disorder
Carotid Artery Stenosis
Cellulitis
Cerebral Atrophy
Cerebral Ischaemia
Cerebrovascular Accident
Cerebrovascular Disorder
Chest Pain
Chest X-Ray Abnormal
Computerised Tomogram
Abnormal
Condition Aggravated
Cystitis

Freedom Of Information (FOI) Report

Deep Vein Thrombosis
Diabetes Mellitus
Difficulty In Walking
Disorientation
Dizziness
Drug Ineffective
Dry Mouth
Dysarthria
Dysstasia
Electrocardiogram St-T
Change
Emphysema
Erythema Of Eyelid
Face Oedema
Haematuria
Headache
Hemiparesis
Hepatic Steatosis
Hypertension
Hypoaesthesia
Intervertebral Disc
Protrusion
Lentigo
Muscular Weakness
Myalgia
Nervous System Disorder
Neuropathy Peripheral
Nocturia
Obesity
Obstructive Uropathy
Oedema Peripheral
Osteoarthritis
Pain
Pain In Extremity
Pco2 Increased
Pelvic Pain
Pharyngitis
Phlebitis
Pitting Oedema
Pneumonia
Pollakiuria
Polyarthritits
Primary
Hyperaldosteronism
Protein Urine Present
Pruritus
Pulmonary Embolism

Pyelogram Retrograde
Abnormal
Pyrexia
Radiculopathy
Rash
Renal Disorder
Sciatica
Sinusitis
Skin Discolouration
Sleep Apnoea Syndrome
Spinal Column Stenosis
Spinal Osteoarthritis
Stasis Dermatitis
Tenderness

Date:04/08/02ISR Number: 3896825-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200392

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Health	Neurontin			
		Drug Level Above	Professional	(Gabapentin)	PS		
		Therapeutic	Company				
		Haemodialysis	Representative				

Date:04/08/02ISR Number: 3896827-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200391

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Residue	Health	Neurontin			
			Professional	(Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:04/08/02ISR Number: 3896928-1Report Type:Expedited (15-DaCompany Report #034-0945-M0200003

Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Foreign	Gabapentin			
		Fall	Health	(Gabapentin)	PS		ORAL
1600 MG (QID)							
PER ORAL		Head Injury	Professional				
		Loss Of Consciousness	Company	(Sertraline)	SS		ORAL
150 MG (TID)							
PER ORAL			Representative				
				(Lormetazepam)	C		
				(Ranitidine			
				Hydrochloride)	C		

Date:04/08/02ISR Number: 3897262-6Report Type:Expedited (15-DaCompany Report #A206957

Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Dizziness	Foreign	Zoloft Tablets	PS		ORAL
150.00 MG							
Intervention to		Fall	Health				
TOTAL: DAILY:							
Prevent Permanent		Head Injury	Professional				
ORAL							
Impairment/Damage		Loss Of Consciousness	Company	Gabapentin	SS		ORAL
1600.00 MG							
			Representative				
TOTAL: DAILY:							
ORAL				Noctamid	C		
				Zantac (Ranitidine)	C		

Date:04/09/02ISR Number: 3897189-XReport Type:Direct

Company Report #CTU 165277

Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Life-Threatening
Other

Aggression
Anger
Antisocial Behaviour
Convulsion
Drug Dependence

Neurontin Parke
Davis

PS Parke Davis

Date:04/09/02ISR Number: 3898409-8Report Type:Direct
Age:61 YR Gender:Female I/FU:I

Company Report #CTU 165284

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Medication Error		Neurontin	PS		
Intervention to				Noroxin	SS		
460MG BID X							
Prevent Permanent							
10D							
Impairment/Damage							

Date:04/10/02ISR Number: 3898264-6Report Type:Expedited (15-DaCompany Report #049-0945-M0200044
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Hepatic Cirrhosis	Foreign	Neurontin			
Hospitalization -		Hepatocellular Damage	Health	(Gabapentin)	PS		ORAL
2400 MG (800							
Initial or Prolonged		Hepatotoxicity	Professional				
MG, TID), PER							
ORAL		Pruritus					
				Pravastatin Sodium	C		
				Silicur J	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/10/02ISR Number: 3898266-XReport Type:Expedited (15-DaCompany Report #044-0945-M0200047

Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression	Foreign	Neurontin			
PER ORAL		Drug Dependence	Consumer	(Gabapentin)	PS		ORAL
		Fatigue					
		Fluid Retention					
		Weight Increased					

Date:04/10/02ISR Number: 3898831-XReport Type:Direct

Company Report #CTU 165356

Age:11 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100 MG AM &		Depression		Seroquel	PS		
200 MG HS PO		Neutropenia					
800 MG TID PO				Neurontin	SS		ORAL
				Clonidine	C		
				Hemocyte Plus	C		

Date:04/10/02ISR Number: 3899456-2Report Type:Expedited (15-DaCompany Report #001-0719-M0100407

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Amnesia	Consumer	Lopid (Gemfibrozil)	PS		ORAL
1200 MG (600							
Initial or Prolonged		Aphasia	Health				
MG, BID), PER							
Disability		Arthralgia	Professional				
ORAL							
Other		Asthenia		Gabapentin	SS		
1600 MG (400							
MG, 2		Back Pain					
		Blood Cholesterol					
CAPSULES IN							

AM AND PM)	Increased		
ONE EVERY 5	Blood Pressure Increased	Glyceryl Trinitrate	SS
MIN., MAX: 3	Blood Triglycerides		
(PRN)	Increased		
0.4 MG	Coronary Artery Disease	Cerivastatin Sodium	SS
(DAILY),	Decreased Activity		
	Depression		
	Dizziness		
	Fatigue		
	Hyperlipidaemia		
	Hypoaesthesia		
	Infarction		
	Insomnia		
	Nausea		
	Neuralgia		
	Neuropathy Peripheral		
	Pain In Extremity		
	Paraesthesia		
	Pruritus		
	Speech Disorder		
	Tremor		

Date:04/10/02ISR Number: 3899459-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200210

Age: Gender:Female I/FU:I

Outcome	PT
Other	Brain Neoplasm
	Headache
	Hypoaesthesia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pituitary Enlargement Visual Disturbance	Report Source	Product	Role	Manufacturer	Route
3600 MG	(THREE TIMES DAILY), PER ORAL		Health Professional	Neurontin (Gabapentin)	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: 3899461-6Report Type:Expedited (15-DaCompany Report #001-0945-M0200409
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other PER ORAL		Arrhythmia Constipation	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Decreased Activity Dehydration Depression Dizziness Eye Disorder		...	SS		

Face Oedema
Fatigue
Fibromyalgia
Headache
Heart Rate Increased
Insomnia
Migraine
Sleep Walking
Tremor
Weight Increased

Date:04/10/02ISR Number: 3899462-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200160
Age: Gender:Male I/FU:F

Outcome PT
Other Abdominal Distension
Abnormal Faeces
Angioneurotic Oedema
Balance Disorder
Change Of Bowel Habit
Clostridium Colitis
Constipation
Culture Stool Positive
Discomfort
Dyskinesia
Facial Palsy

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2800 MG		Health Professional	Neurontin (Gabapentin)	PS		ORAL
(THREE TIMES DAILY), PER ORAL			Antibiotics	SS		
			Hydrochlorothiazide	C		
			Diltiazem			
			Hydrochloride	C		
			Doxazosin Mesilate	C		
			Atorvastatin	C		
			Rabeprazole Sodium	C		
			Domperidone	C		
			Clonazepam	C		

Date:04/11/02ISR Number: 3899330-1Report Type:Expedited (15-DaCompany Report #033-0945-M0200008
 Age:82 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1200 MG (300	Activated Partial Thromboplastin Time	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
MG, QID), PER ORAL		Shortened					
		Anti Factor Viii Antibody					
2000 MG (500		Positive		Paracetamol	SS		ORAL
MG, QID), PER ORAL		Coagulation Factor Viii					
		Level Decreased					
INTRAVENOUS	40	Coagulation Time Prolonged		Human Prothrombin Complex	SS		

MILLILITERS		Condition Aggravated			
(DAILY),		Disorientation			
INTRAVENOUS		Haematoma			
		Sensorimotor Disorder	Heparin-Fraction,	SS	
SUBCUTANEOUS	2500 IU/ 0.2	Subdural Haematoma	Sodium Salt		
ML (DAILY),					
SUBCUTANEOUS			Omeprazole	SS	ORAL
20 MG					
(DAILY), PER					
ORAL			Trinitrine	SS	
TRANSDERMAL	5 MG (DAILY),				
TRANSDERMAL					

Date:04/11/02ISR Number: 3899577-4Report Type:Expedited (15-DaCompany Report #001-0073-M0200175
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Consumer	Dilantin (Phenytoin			
Other		Gait Disturbance		Sodium)	PS		
		Headache		(Gabapentin)	SS		ORAL
1800 MG (600		Vertigo					
MG, TID) PER							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/12/02ISR Number: 3900158-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200416
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Consumer	Neurontin			
2400 MG		Diarrhoea		(Gabapentin)	PS		ORAL
(DAILY), PER		Hepatic Pain					
ORAL		Liver Function Test					
		Abnormal		(Buspirone			
		Weight Decreased		Hydrochloride)	C		
		Weight Increased		(Valproate			
				Semisodium)	C		
				(Citalopram			
				Hydrobromide)	C		

Date:04/12/02ISR Number: 3900198-5Report Type:Expedited (15-DaCompany Report #034-0945-M0200002
 Age:21 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Foreign	Gabapentin			
1200 MG(TID),		Drug Interaction	Health	(Gabapentin)	PS		ORAL
PER ORAL		Neutropenia	Professional				
350 MG (TWICE			Company	Clozapine	SS		ORAL
DAILY), PER			Representative				
ORAL							

Date:04/12/02ISR Number: 3900770-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200417
 Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Mouth	Consumer	Neurontin			

300 MG	Dysarthria	(Gabapentin)	PS	ORAL
(DAILY) PER	Eating Disorder			
ORAL	Sensory Loss			
	Weight Decreased	Amlodipine Besilate	C	
		Theophylline	C	
		Enalapril Maleate	C	
		Zafirlukast	C	

Date:04/12/02ISR Number: 3900771-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200426
 Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Neoplasm	Consumer	Neurontin			
		Neoplasm Recurrence		(Gabapentin)	PS		ORAL
900 MG (300		Weight Increased					
MG TID), PER							
ORAL				Calcium	C		

Date:04/12/02ISR Number: 3900877-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200218
 Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Anaemia
Initial or Prolonged	Anorgasmia
	Condition Aggravated
	Hypothyroidism
	Parkinsonism

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tardive Dyskinesia Urinary Incontinence	Health Professional	Neurontin (Gabapentin)	PS		
UNKNOWN	1200 MG						
(BID),							
UNKNOWN				Lithium	C		

Date:04/15/02ISR Number: 3900821-5Report Type:Expedited (15-DaCompany Report #2002UW05015
Age:47 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	100 MG HS PO		Drug Interaction	Health Professional	Seroquel	PS		ORAL
	50 MG BID				Luvox	SS		
	300 MG TID				Neurontin	SS		
	0.25 MG TID				Xanax	SS		
PRN								

Date:04/15/02ISR Number: 3901044-6Report Type:Expedited (15-DaCompany Report #033-0945-M0200040
Age:69 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL	150 PER ORAL		Pulmonary Embolism	Foreign Health Professional	Neurontin (Gabapentin) Topiramate	PS SS		ORAL ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 900 MG (300 Initial or Prolonged MG, TID) PER Other ORAL		Adrenal Neoplasm Anxiety Colitis Ischaemic Dizziness Nausea Orthostatic Hypotension Rosacea Syncope Vasovagal Weight Decreased	Consumer Health Professional	Neurontin (Gabapentin) (Levothyroxine Sodium)	PS C		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1200 MG Initial or Prolonged (BID), PER Other ORAL PER ORAL 5 MG (DAILY), PER ORAL 5 MG (DAILY), PER ORAL SUBLINGUAL 10 MG, SUBLINGUAL		Back Pain Blood Pressure Increased Confusional State Hyporeflexia Pain Spinal Osteoarthritis Transient Ischaemic Attack Vertigo	Consumer Health Professional	Lopid (Gemfibrozil) (Gabapentin) (Amlodipine Besilate) (Glipizide) (Nifedipine) (Cholesterol)	PS SS SS SS		ORAL ORAL ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lowering Drug)	SS
(Furosemide)	C
(Acetylsalicylic	
Acid)	C
(Ergocalciferol,	
Ascorbic Acid, Folic	
Acid, Thiamine	
Hydrochloride,	
Retinol, Riboflavin,	C
(Terazosin)	C
(Labetalol	
Hydrochloride)	C
(Sunastatin)	C
(Clopidogrel)	C
(Meloxicam)	C
(Pericad Plus	
Vitamins)	C
(L-Carnitine)	C
(Magnesium)	C
(Glucosamine	
Chondroitin)	C
(Msm)	C
(Celecoxib)	C
(Tamsulosin	
Hydrochloride)	C
(Tolterodine	
Tartrate)	C

Date:04/15/02ISR Number: 3901522-XReport Type:Expedited (15-DaCompany Report #001-0981-M0201960
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Atrovastatin (Atrovastatin)	PS		
				Gabapentin	SS		
				Rofecoxib	SS		

Date:04/16/02ISR Number: 3900242-5Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-11811908
 Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aortic Aneurysm		Buspirone Hcl	PS	Bristol-Myers Squibb	

			Company
Hospitalization - 15mg	Death		
Initial or Prolonged 5mg	Hepatic Cirrhosis	Diazepam	SS
	Multi-Organ Failure	Percocet Tabs	SS
		Loperamide Hcl	SS
2mg capsules		Propoxyphene Napsylate+Acetaminop hen	SS
100mg/650mg		Neurontin	SS
		Ambien	SS
		Oxycontin	SS
		Morphine	SS

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/16/02ISR Number: 3901008-2Report Type:Direct
 Age:53 YR Gender:Male I/FU:I

Company Report #CTU 165847

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300MG 3 CAP	Acidosis		Gabapentin 300mg	PS		ORAL
Initial or Prolonged TID ORAL	Blood Creatinine Abnormal					
15MG Q 6-8	Blood Potassium Increased		Ms Contin 15mg	SS		ORAL
HRS ORAL	Blood Urea Increased					
	Hypercapnia		Aquaphor Ointment	C		
	Somnolence		Aspirin Ec	C		
			Capsaicin Cream	C		
			Citalopram			
			Hydrobromide	C		
			Clonazepam	C		
			Clopidogrel			
			Bisulfate	C		
			Curasol Wound			
			Dressing Gel	C		
			Emulsion Top Cream	C		
			Gauze			
			Pad-Non-Sterile	C		
			Granulex	C		
			Insulin Novolin			
			70/30-Nph/Reg	C		
			Isosorbide Dinitrate	C		
			Lisinopril	C		
			Metoprolol Tartrate	C		
			Morphine So4	C		
			Olanzapine	C		
			Ranitidine Hcl	C		

Date:04/16/02ISR Number: 3901080-XReport Type:Direct
 Age:66 YR Gender:Male I/FU:I

Company Report #CTU 165809

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 700 MG PO QID 2 YR	Thrombocytopenia		Gabapentin	PS		ORAL
			Navane	C		
			Dilantin	C		

Provera C
 Ativan C
 Cardura C
 Amantadine C
 Folic Acid C
 Adalat Cc C
 Lisinopril C

Date:04/17/02ISR Number: 3901925-3Report Type:Expedited (15-DaCompany Report #C2002-0958.01
 Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aortic Aneurysm	Consumer	Buspirone			
Hospitalization -		Hepatic Cirrhosis		Hydrochloride			
Initial or Prolonged		Multi-Organ Failure		Tablets 15 Mg	PS		
				Diazepam Tablets 5mg	SS		
				Loperamide Hcl			
				Capsules 2 Mg	SS		
				Propoxyphene			
				Napsylate And Apap			
				Tablets 100mg/650mg	SS		
				Neurontin	SS	Parke Davis	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ambien SS Searle
 Oxycodone/Apap SS
 Oxycontin Tablets SS
 Morphine Tablets SS

Date:04/17/02ISR Number: 3902134-4Report Type:Expedited (15-DaCompany Report #02P-056-0190956-00
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Alkaline Phosphatase Increased Pancreatitis Rash Maculo-Papular	Foreign Health Professional	Kaletra Soft Gelatin Capsules (Kaletra) (Lopinavir/Ritonavir) (Lopinavir/Ritonavir	PS		ORAL
1 DOSAGE FORMS, 2 IN 1 D, PER ORAL				Gabapentin	SS		ORAL
500 MG, 3 IN 1 D, PER ORAL				Zidovudine W/Lamivudine	SS		ORAL
1 DOSAGE FORMS, 2 IN 1 D, PER ORAL				Calcium Folate Pyrimethamine Sulfadiazine	C C C		

Date:04/17/02ISR Number: 3902904-2Report Type:Direct Company Report #CTU 165984
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other SEE NOTE ORAL		Epistaxis		Gabapentin	PS		ORAL

Date:04/18/02ISR Number: 3902652-9Report Type:Expedited (15-DaCompany Report #061-0945-M0200047
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (UNKNOWN), PER ORAL		Abdominal Pain Upper Chest Pain Confusional State Hypertension	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Codeine Phosphate, Paracetamol)	PS C		ORAL

Date:04/18/02ISR Number: 3902726-2Report Type:Expedited (15-DaCompany Report #055-0945-M0200026
 Age:87 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 400 MG Initial or Prolonged (DAILY), PER ORAL		Drug Ineffective Malaise	Foreign Consumer	Gabapentin (Gabapentin) (Unspecified Medications)	PS SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/18/02ISR Number: 3903099-1Report Type:Expedited (15-DaCompany Report #039-0945-M0200002

Age:81 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (DAILY), PER ORAL		Dermatitis Atopic Rash Macular	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
200 MG (100 MG, BID) PER ORAL				Celecoxib	SS		ORAL

Date:04/19/02ISR Number: 3902178-2Report Type:Expedited (15-DaCompany Report #309551

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion Neonatal Feeding Problem In Newborn		Rivotril Neurontin Depakine	PS SS SS	Roche	

Date:04/19/02ISR Number: 3903079-6Report Type:Direct

Company Report #CTU 166058

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Required 600MG 4 TIMES Intervention to ORAL Prevent Permanent Impairment/Damage		Drug Withdrawal Syndrome		Neurontin 600mg Parke Davis	PS	Parke Davis	ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 200MG PO TID	Balance Disorder		Carbamazepine	PS		ORAL
Initial or Prolonged 0.5MG TID	Coordination Abnormal		Clonazepam	SS		
300-600 MG	Dysarthria		Gabapentin	SS		
TID	Visual Disturbance					
50-150 MG Q			Trazadone	SS		
HS			Aspirin	C		
			Fosinopril	C		
			Hydroxyzine	C		
			Ntg Sl	C		
			Simvastatin	C		
			Simethicone	C		
			Rabeprazole	C		
			Ibuprofen	C		
			Hctz	C		
			Gemfibrozil	C		
			Atenolol	C		
			Citalopram	C		
			Allopurinol	I		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/19/02ISR Number: 3903874-3Report Type:Direct
 Age:71 YR Gender:Male I/FU:I

Company Report #CTU 166188

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 400MG TID	Confusional State		Gabapentin	PS		
Initial or Prolonged 100MG TID	Fall		Phenytoin	SS		
5MG			Oxybutynin	SS		
5MG BID Q HR			Terazosin	SS		
			Clopidogrel	C		
			Beclomethasone	C		
			Donepezil	C		
			Metoprolol	C		
			Aspirin	C		
			Pseudoephedrine	C		

Date:04/19/02ISR Number: 3904198-0Report Type:Expedited (15-DaCompany Report #02P-056-0191091-00
 Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Nightmare	Foreign Health Professional	Isoptin Tablets (Isoptin) (Verapamil) (Verapamil)	PS		ORAL
3 IN 1 D, PER						
ORAL			Neurontin (Gabapentin)	SS		ORAL
600 MG, PER						
ORAL			Transipeg (Macrogol)	SS		ORAL
PER ORAL			Depakote	SS		
500 MG, 2 IN						
1 D			Lezonil	SS		ORAL
PER ORAL						

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
PER ORAL				Piportil (Pipotiazine)	SS		ORAL
Date:04/22/02ISR Number: 3904523-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200453 Age:30 YR Gender:Female I/FU:I							
Dose							
Other		Clonic Convulsion Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG		Difficulty In Walking					
(DAILY), PER		Insomnia					
ORAL		Somnolence Tremor		Phenytoin Sodium	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Date:04/22/02ISR Number: 3904561-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200371 Age: Gender:Female I/FU:F							
Hospitalization - Initial or Prolonged		Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL
PER ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Date:04/22/02ISR Number: 3904631-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200450 Age: Gender:Male I/FU:I							
Death							
Hospitalization -							

Freedom Of Information (FOI) Report

Initial or Prolonged

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Aortic Aneurysm	Health	Neurontin			
		Hepatic Cirrhosis	Professional	(Gabapentin)	PS		
		Multi-Organ Failure		(Buspirone			
				Hydrochloride)	SS		
				Diazepam	SS		
				(Loperamide			
				Hydrochloride)	SS		
				(Propoxyphene			
				Napsylate,			
				Acetaminophen)	SS		
				(Zolpidem Tartrate)	SS		
				(Oxycodone/Acetamino			
				phen)	SS		
				(Oxycodone			
				Hydrochloride)	SS		
				Morphine	SS		

Date:04/22/02ISR Number: 3904633-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200370
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arteriosclerosis	Consumer	Neurontin			
Other		Calcinosis		(Gabapentin)	PS		ORAL
600 MG		Inguinal Hernia					
(DAILY), PER		Intervertebral Disc					
ORAL		Degeneration		Unspecified			
		Intervertebral Disc		Narcotics	C		
		Disorder		Diazepam	C		
		Pain					
		Scoliosis					

Date:04/22/02ISR Number: 3904679-XReport Type:Expedited (15-DaCompany Report #2002-02-1107
 Age:48 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abortion Induced	Health	Rebetol (Ribavirin)			
Other		Multiple Congenital	Professional	Capsules	PS		ORAL
1200 MG QD		Abnormalities	Company				
ORAL		Pregnancy	Representative	Intron A (Interferon			
		Transmission Of Drug Via		Alfa-2b Recombinant)			
		Semen		Injectable Solution	SS		
SUBCUTANEOUS	3 MU TIW						
SUBCUTANEOUS				Gabapentin	SS		
				Olanzapine	SS		

Date:04/22/02ISR Number: 3904747-2Report Type:Expedited (15-DaCompany Report #200211316BCC
Age:26 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Alanine Aminotransferase
Initial or Prolonged	Increased
	Aspartate
	Aminotransferase
	Increased
	Blood Creatine

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Phosphokinase Increased Gamma-Glutamyltransferase Increased Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
			Health Professional Other	Aleve Tablets (Naproxen Sodium) Aspirin (Acetylsalicylic Acid) Acetaminophen Effexor (Venlafaxine Hydrochloride) Sudafed (Pseudoephedrine Hydrochloride) Neurontin (Gabapentin)	PS SS SS SS SS SS		

Date:04/22/02ISR Number: 3904985-9Report Type:Expedited (15-DaCompany Report #033-0945-M0200050

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Nightmare	Foreign Health Professional	Neurontin (Gabapentin) Macrogol Valproate Semisodium	PS SS SS		ORAL ORAL ORAL
1000 MG (500 MG, BID), PER ORAL (DAILY), PER ORAL 120 (TID), PER ORAL				Bromazepam Verapamil Hydrochloride Pipotiazine	SS SS C		ORAL ORAL ORAL

Date:04/22/02ISR Number: 3904986-0Report Type:Expedited (15-DaCompany Report #033-0945-M0200045
 Age:38 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1500 MG (TID), PER ORAL	Blood Alkaline Phosphatase Increased Pancreatitis Rash Rash Maculo-Papular	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG (BID) PER ORAL			Lamivudine, Zidovudine	SS		ORAL
800 MG (BID), PER ORAL			Lopinavir, Ritonavir	SS		ORAL
			Calcium Folate Pyrimethamine Sulfadiazine	C C C		

Date:04/22/02ISR Number: 3904988-4Report Type:Expedited (15-DaCompany Report #358-0945-M0200012
 Age:82 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 1200 MG DAILY, PER	Loss Of Consciousness	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

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Freedom Of Information (FOI) Report

ORAL

Oxycodone
 Hydrochloride C
 Acetylsalicylic Acid C
 Carvedilol C
 Isosorbide
 Mononitrate C
 Furosemide C
 Losartan Potassium C

Date:04/22/02ISR Number: 3904990-2Report Type:Expedited (15-DaCompany Report #351-0945-M0200002
 Age:52 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 3600 MG DAILY , PER ORAL	Hepatic Failure	Foreign Health Professional	Neurontin (Gabapentin) (Omeprazole) (Clonazepam)	PS C C		ORAL

Date:04/22/02ISR Number: 3904991-4Report Type:Expedited (15-DaCompany Report #044-0945-M0200058
 Age:43 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 100 MG DAILY , PER ORAL	Disorientation Face Oedema Neck Pain Speech Disorder Tenderness Tongue Oedema Vision Blurred	Foreign Health Professional	Gabapentin (Gabapentin) (Acetylsalicylic Acid) (Heparin-Fraction Sodium Salt) (Amoxicillin) (Flucloxacillin) (Metoclopramide) (Prochlorperazine) (Paracetamol) (Morphine Sulfate)	PS C C C C C C C		ORAL

(Nifedipine)

C

Date:04/22/02ISR Number: 3905025-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200447

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Health	Neurontin			
		Pain	Professional	(Gabapentin)	PS		ORAL

900 MG (300

MG, TID), PER

ORAL

Furosemide	C
Potassium	C
Paracetamol,	
Dextropropoxyphene	C
Compounded Ativan,	
Benadryl,	
Haloperidol	
Suppository	C
Oxycodone	
Hydrochloride	C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/22/02ISR Number: 3905026-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200454

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bundle Branch Block Left	Health Professional	Neurontin (Gabapentin)	PS		
				Amitriptyline	SS		

Date:04/22/02ISR Number: 3905050-7Report Type:Expedited (15-DaCompany Report #033-0945-M0200044

Age:1 DY Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Apgar Score Low Clonic Convulsion Feeding Problem In	Foreign Health Professional	Neurontin Valproic Acid, Valproate Sodium	PS SS		
1500 MG (BID)		Newborn		Clonazepam	SS		
0.5 MG DAILY		Foetal Distress Syndrome Maternal Drugs Affecting Foetus Neonatal Apnoeic Attack Neonatal Disorder Tonic Clonic Movements					

Date:04/23/02ISR Number: 3906075-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200456

Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Glossitis Psychotic Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
200 MG (100							

MG, BID), PER

ORAL

Quetiapine	C
Clonazepam	C
Lithium Carbonate	C
..	C

Date:04/23/02ISR Number: 3906078-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200460
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Renal Cell Carcinoma Stage Unspecified	Health Professional	Neurontin (Gabapentin)	PS		
2400 MG			Company				
(DAILY)			Representative				

Date:04/23/02ISR Number: 3906079-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200465
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Cholesterol Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
800 MG (BID),		Bone Scan Abnormal					
PER ORAL		Cardiac Valve Disease		(Atorvastatin)	SS		ORAL
40 MG		Impaired Gastric Emptying					
(DAILY), PER		Neuropathy Peripheral					
ORAL		Pain		Glipizide Metformin Hydrochloride	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Mirtazapine C
 Paracetamol,
 Hydrocodone
 Bitartrate C
 Bumetanide C
 Potassium Chloride C
 Acetylsalicylic Acid C
 Ascorbic Acid C
 Ergocalciferol,
 Calcium Phosphate,
 Calcium Sodium
 Lactate C
 Ascorbic Acid,
 Tocopheryl Acetate,
 Retinol, Zinc,
 Calcium, Vitamins
 Nos, Minerals Nos, C

Date:04/23/02ISR Number: 3906082-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200466

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
(BID), PER		Fall		(Gabapentin)	PS		ORAL
ORAL		Weight Increased					

Paracetamol,
 Oxycodone
 Hydrochloride,
 Oxycodone
 Terephthalate C
 Furosemide C
 Irbesartan C
 Glipizide C
 Acetylsalicylic Acid C

Date:04/23/02ISR Number: 3906106-5Report Type:Expedited (15-DaCompany Report #309551

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged 0.5 MG DAILY	Apgar Score Low Convulsion Neonatal	Foreign Other	Rivotril (Clonazepam)	PS	ORAL
ORAL	Feeding Disorder Of Infancy Or Early Childhood		Neurontin (Gabapentin)	SS	ORAL
3 DOSE FORM DAILY ORAL	Foetal Distress Syndrome				
750 MG 2 PER DAY ORAL	Maternal Drugs Affecting Foetus		Depakine (Valproate Sodium) 500 Mg	SS	ORAL
	Neonatal Apnoeic Attack Pregnancy				

Date:04/23/02ISR Number: 3906199-5Report Type:Expedited (15-DaCompany Report #044-0945-M0200059
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Hypertension	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
300 MG (100 MG, TID), PER ORAL			Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Celiprolol C
 Diltiazem
 Hydrochloride C
 Paracetamol
 Dextropropoxyphene
 Hydrochloride C

Date:04/23/02ISR Number: 3906221-6Report Type:Expedited (15-DaCompany Report #031-0945-M0200005
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Paramnesia	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:04/23/02ISR Number: 3906371-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200467
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Autoimmune Thyroiditis Oedema Peripheral	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, TID), PER ORAL							

Venlafaxine
 Hydrochloride) C
 Salbutamol C

Date:04/24/02ISR Number: 3906778-5Report Type:Expedited (15-DaCompany Report #031-0945-M0200006
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 4200 MG		Blood Pressure Decreased Gastrointestinal	Foreign Health	Neurontin(Gabapentin)	PS		ORAL

Haemorrhage Professional
 (DAILY), PER
 Loss Of Consciousness
 ORAL
 (Diazepam) C
 (Midazolam) C
 (Fentanyl) C

Date:04/24/02ISR Number: 3906780-3Report Type:Expedited (15-DaCompany Report #044-0945-M0200039
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (100 Other MG, TID), PER		Cerebrovascular Accident	Foreign Consumer Health Professional	Neurontin Tablets (Gabapentin)	PS		ORAL
ORAL				Cimetidine	SS		
INTRA VENOUS	400 MG (1						
DOSE),							
INTRA VENOUS				Salbutamol	C		
				Beclometasone	C		
				Fluticasone			
				Propionate	C		
				Lansoprazole	C		
				Tegremet	C		
				Clopidogrel	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Fluvastatin Sodium C
 Amitriptyline C
 Carbamazepine C

Date:04/24/02ISR Number: 3907353-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200477
 Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hemiplegia	Health	Neurontin			
		Muscle Twitching	Professional	(Gabapentin)	PS		ORAL
PER ORAL		Pain		Donepezil			
				Hydrochloride	C		
				Mirtazapine	C		
				Vitamins	C		

Date:04/25/02ISR Number: 3906111-9Report Type:Expedited (15-DaCompany Report #309551
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion Neonatal		Rivotril	PS	Roche	
Initial or Prolonged		Feeding Problem In		Neurontin	SS		
		Newborn		Depakine	SS		

Date:04/26/02ISR Number: 3907950-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200232
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin			
			Health	(Gabapentin)	PS		
900 MG (300			Professional				
MG, TID)							

Date:04/26/02ISR Number: 3908105-6Report Type:Expedited (15-DaCompany Report #02P-056-0191421-00
 Age:15 DY 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		Clonic Convulsion Convulsion Neonatal Drug Withdrawal Convulsions Dysphagia Foetal Distress Syndrome Maternal Drugs Affecting Foetus Neonatal Apnoeic Attack Neonatal Disorder Sleep Disorder	Foreign Health Professional Other	Depakine Chron Tablets (Depakene) (Sodium Valpraote/Valproic Acid) (Sodium Neurontin (Gabapentin) Rivotril (Clonazepam)	PS SS SS		

Date:04/26/02ISR Number: 3908167-6Report Type:Expedited (15-DaCompany Report #055-0945-M0200024
Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL		Cardio-Respiratory Arrest Coordination Abnormal Oedema Visual Disturbance	Foreign Consumer	Gabapentin (Gabapentin) Pantoprazole (Folic Acid + Ferrous	PS C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C
 Amitriptyline C

Date:04/26/02ISR Number: 3908344-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200283
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Alcoholism		Neurontin			
Other		Blood Alcohol Increased		(Gabapentin)	PS		ORAL
PER ORAL		Coma					
		Fall					
		Hallucination					
		Hypothermia					
		Medication Error					

Date:04/26/02ISR Number: 3908423-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200490
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abdominal Pain	Consumer	Neurontin			
		Completed Suicide		(Gabapentin)	PS		
		Drug Ineffective		Sertraline			
		Eating Disorder		Hydrochloride	SS		
		Gun Shot Wound		Unknown Narcotic	SS		
		Idiosyncratic Drug					
		Reaction					
		Oligodipsia					

Date:04/26/02ISR Number: 3908432-2Report Type:Expedited (15-DaCompany Report #A208483
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abdominal Pain	Consumer	Zoloft Tablets	PS		
		Completed Suicide		Gabapentin	SS		
		Drug Ineffective		Unknown Narcotic	SS		
		Eating Disorder					
		Gun Shot Wound					
		Heart Injury					

Idiosyncratic Drug
Reaction
Oligodipsia

Date:04/26/02ISR Number: 3908511-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200308
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Prostatic Specific Antigen Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG							

(DAILY), PER

ORAL

(Acetylsalicylic
Acid) C
(Ergocalciferol,
Ascorbic Acid, Folic
Acid, Thiamine
Hydrochloride,
Retinol, Riboflavin, C
(Tamsulosin

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride C
 (Fenofibrate) C
 (Olanzapine) C

Date:04/29/02ISR Number: 3909291-4Report Type:Expedited (15-DaCompany Report #A208992
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Coma	Health Professional	Zoloft Tablets	PS		
Intervention to Prevent Permanent Impairment/Damage		Drug Interaction Pneumonia	Company Representative	Gabapentin	SS		

Date:04/29/02ISR Number: 3909296-3Report Type:Expedited (15-DaCompany Report #A208993
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Coma	Health Professional	Zoloft Tablets	PS		
Intervention to Prevent Permanent Impairment/Damage		Drug Interaction Pneumonia	Company Representative	Gabapentin	SS		

Date:04/29/02ISR Number: 3909409-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200494
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Coma	Health Professional	Neurontin (Gabapentin)	PS		
		Drug Interaction Pneumonia	Company Representative	(Sertraline)	SS		

Date:04/29/02ISR Number: 3909411-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200495
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Coma	Health	Neurontin			

Drug Interaction
Pneumonia

Professional
Company
Representative

(Gabapentin)
(Sertraline)

PS
SS

Date:04/29/02ISR Number: 3909589-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200491
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Atherosclerosis	Health	Neurontin			
Other		Cardiomyopathy	Professional	(Gabapentin)	PS		ORAL
3600 MG (FOUR TIMES DAILY), PER ORAL		Drug Level Increased Myocardial Infarction					
				Hydrocodone	C		
				Clonazepam	C		
				Methadone	C		
				Cyclobenzaprine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/29/02ISR Number: 3909598-0Report Type:Expedited (15-DaCompany Report #351-0945-M0200002

Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3600 MG (DAILY), PER ORAL		Alanine Aminotransferase Increased Aspartate Aminotransferase Increased Liver Transplant	Foreign Health Professional	Neurontin (Gabapentin) Omeprazole Clonazepam	PS C C		ORAL

Date:04/29/02ISR Number: 3909634-1Report Type:Expedited (15-DaCompany Report #052-0945-M0200001

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Sudden Death	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Ethanol Celecoxib	PS SS C		ORAL ORAL

Date:04/29/02ISR Number: 3909906-0Report Type:Expedited (15-DaCompany Report #044-0945-M0200061

Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 1800 MG (THREE TIMES DAILY), PER ORAL		Skin Lesion Tongue Disorder	Foreign Health Professional	Gabapentin (Gabapentin) Fluconazole Lansoprazole Haloperidol	PS C C C		ORAL

Dantron Poloxamer C
Diclofenac C
Dexamethasone C
Oxycodone
Hydrochloride C

Date:04/29/02ISR Number: 3910340-8Report Type:Expedited (15-DaCompany Report #309551
Age: Gender:Male I/FU:F

Outcome PT
Hospitalization - Apgar Score Low
Initial or Prolonged Apnoea
Clonic Convulsion
Convulsion
Convulsion Neonatal
Drug Withdrawal Syndrome
Neonatal
Epilepsy
Feeding Problem In
Newborn
Foetal Distress Syndrome
Gastrointestinal Tube
Insertion
Grand Mal Convulsion
Maternal Drugs Affecting
Foetus

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Small For Dates Baby

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
0.5 MG DAILY		Foreign Other	Rivotril (Clonazepam)	PS		ORAL
ORAL			Neurontin (Gabapentin)	SS		ORAL
3 DOSE FORM						
DAILY ORAL						
750 MG 2 PER			Depakine (Valproate Sodium) 500 Mg	SS		ORAL
DAY ORAL						

Date:04/30/02ISR Number: 3910586-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200496
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Fall Hypersensitivity	Consumer	Neurontin (Gabapentin)	PS		ORAL
1800 MG		Iatrogenic Injury					
(THREE TIMES		Nerve Injury					
DAILY), PER		Oedema Peripheral					
ORAL		Pain In Extremity		Hyaluronate Sodium Tramadol Hydrochloride Paracetamol Ibuprofen Acetylsalicylic Acid, Magnesium Hydroxide, Aluminium Hydroxide	SS C C C C		

Date:04/30/02ISR Number: 3910588-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200532
Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain	Consumer	Neurontin			
		Fall		(Gabapentin)	PS		ORAL
1800 MG							
(TID), PER							
ORAL							
				Tramadol			
				Hydrochloride	C		
				Paracetamol	C		
				Ibuprofen	C		
				Acetylsalicylic			
				Acid, Magnesium			
				Hydroxide, Aluminum			
				Hydroxide	C		

Date:04/30/02ISR Number: 3910602-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200497
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Drug Level Decreased	Health	Neurontin			
Initial or Prolonged		Fall	Professional	(Gabapentin)	PS		ORAL
3600 MG							
Other		Head Injury					
(THREE TIMES							
DAILY), PER							
ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Phenytoin Sodium C

Date:04/30/02ISR Number: 3910617-6Report Type:Expedited (15-DaCompany Report #001-0981-M0201960
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Brain Neoplasm	Consumer	Atorvastatin			
Life-Threatening		Convulsion		(Atorvastatin)	PS		ORAL
PER ORAL							
Hospitalization -		Difficulty In Walking		Gabapentin	SS		ORAL
PER ORAL							
Initial or Prolonged		Lung Neoplasm Malignant		Rofecoxib	SS		ORAL
PER ORAL							
Other		Metastases To Bone		Phenobarbital	SS		
		Metastases To Liver		Valproate Semisodium	C		
		Metastases To Spine					
		Muscular Weakness					
		Myalgia					
		Rash					
		Somnolence					
		Tremor					

Date:04/30/02ISR Number: 3910955-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200535
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Diarrhoea	Consumer	Neurontin			
Initial or Prolonged		Pain		(Gabapentin)	PS		ORAL
300 MG							
Other		Weight Decreased					
(DAILY), PER							
ORAL							
				Oxycodone			
				Hydrochloride	SS		ORAL
PER ORAL							
				Hydrocodone/Acetamin			
				open	SS		
PER ORAL							
				Amitriptyline	C		

Date:04/30/02ISR Number: 3911384-2Report Type:Expedited (15-DaCompany Report #034-0945-M0200004
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (THREE TIMES DAILY), PER ORAL		Dyskinesia Haemodialysis Psychosexual Disorder	Foreign Health Professional	Gabapentin (Gabapentin) Sildenafil	PS C		ORAL

Date:05/01/02ISR Number: 3909994-1Report Type:Direct Company Report #CTU 167010
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Mania		Citalopram 20mg Q Am Olazapine Gabapentin	PS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/01/02ISR Number: 3910336-6Report Type:Direct
Age:69 YR Gender:Male I/FU:I

Company Report #CTU 167037

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG BID Initial or Prolonged		Oedema Peripheral		Gabapentin	PS		

Date:05/01/02ISR Number: 3910425-6Report Type:Direct
Age: Gender: I/FU:I

Company Report #CTU 167075

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100MG BID ORAL		Pruritus Rash		Neurontin	PS		ORAL
				Aspirin Zoloft Prilosec	C C C		

Date:05/01/02ISR Number: 3910535-3Report Type:Direct
Age:26 YR Gender:Female I/FU:I

Company Report #CTU 167115

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Required 225MG 1 Intervention to DAILY ORAL Prevent Permanent Impairment/Damage 300MG 3XDAILY ORAL		Anxiety Depression Drug Effect Decreased Panic Disorder Vision Blurred		Effexor Xr 225 Mg Wyeth-Ayerst Neurontin 300 Mg Pfizer	PS SS	Wyeth-Ayerst Pfizer	ORAL ORAL

Date:05/01/02ISR Number: 3910720-0Report Type:Expedited (15-DaCompany Report #200204-1548 (0)
Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Alanine Aminotransferase Increased Aspartate	Health Professional	Sudafed - Dosage Form Unspecified (Pseudoephedrine)	PS		ORAL
PER ORAL							
		Aminotransferase Increased		Aleve (Naproxen Sodium)	SS		ORAL
PER ORAL							
		Blood Creatine Phosphokinase Increased Suicide Attempt		Aspirin Bayer (Acetylsalicylic Acid)	SS		ORAL
PER ORAL							
				Acetaminophen (Paracetamol)	SS		ORAL
PER ORAL							
				Effexor (Venlafaxine)	SS		ORAL
PER ORAL							
				Neurontin (Gabapentin)	SS		ORAL
PER ORAL							

Date:05/01/02ISR Number: 3911171-5Report Type:Expedited (15-DaCompany Report #001-0719-M0100408
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Angiopathy
Other	Arteriosclerosis
	Arthralgia
	Asthenia
	Atrial Hypertrophy
	Back Pain

Freedom Of Information (FOI) Report

Benign Prostatic
Hyperplasia
Bladder Disorder
Blood Triglycerides
Increased
Cardiovascular Disorder
Carotid Artery Stenosis
Cellulitis
Cerebrovascular Accident
Cerebrovascular Disorder
Cervical Spinal Stenosis
Condition Aggravated
Confusional State
Cystitis
Depression
Diabetes Mellitus
Difficulty In Walking
Disorientation
Dizziness
Dry Mouth
Emphysema
Erythema Of Eyelid
Face Oedema
Fatigue
Feeling Abnormal
Flat Affect
Gait Disturbance
Headache
Hepatic Steatosis
Hypertension
Hypoaesthesia
Hyporeflexia
Hypotriglyceridaemia
Intervertebral Disc
Protrusion
Joint Stiffness
Lumbar Spinal Stenosis
Memory Impairment
Mobility Decreased
Muscle Spasms
Muscle Tightness
Myalgia
Neck Pain
Nocturia
Oedema Peripheral
Osteoarthritis
Pain

Pharyngitis
Phlebitis
Pitting Oedema
Pneumonia
Pollakiuria
Polyuria
Pruritus
Radicular Syndrome
Rash
Sciatica
Sinusitis
Skin Ulcer
Sleep Apnoea Syndrome
Spinal Osteoarthritis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Adverse Event	Report Source	Product	Role	Manufacturer	Route
		Stasis Dermatitis Transient Ischaemic Attack					
1200 MG (BID), PER ORAL PER ORAL 5 MG (DAILY), PER ORAL 5 MG (DAILY), PER ORAL SUBLINGUAL SUBLINGUAL	10 MG,	Urinary Tract Infection Urticaria Varicose Veins Pelvic Vision Blurred Weight Increased	Consumer Health Professional	Lopid (Gemfibrozil) (Gabapentin) (Amlodipine Besilate) (Glipizide) (Nifedipine) (Unspecified Cholesterol Lowering Drug) Furosemide Acetylsalicylic Acid Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Retinol, Riboflavin, Terazosin Labetalol Hydrochloride Sunastatin Clopidogrel Meloxicam Pericad Plus Vitamins L-Carnitine Magnesium Glucosamine Chondroitin	PS SS SS SS SS		ORAL ORAL ORAL ORAL

Msm C
 Celecoxib C
 Tamsulosin C
 Hydrochloride C
 Tolterodine Tartrate C

Date:05/02/02ISR Number: 3911462-8Report Type:Direct
 Age:69 YR Gender:Male I/FU:I

Company Report #CTU 167197

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300MG PO BID Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	Clonic Convulsion		Neurontin	PS		
			Prozac	C		
			Restoril	C		
			Ntg Patch	C		
			Tums	C		
			Epogen	C		
			Renagel	C		
			Pepcid	C		
			Xanax	C		
			Ciloxan	C		
			Zemplar	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/02/02ISR Number: 3911574-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200176

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arachnoiditis	Consumer	Neurontin			
1600 MG		Back Pain	Health	(Gabapentin)	PS		
(DAILY),		Intervertebral Disc	Professional				
		Disorder					
		Pain					
		Sensory Disturbance					

Date:05/02/02ISR Number: 3911789-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200504

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Creatine	Foreign	Neurontin			
Initial or Prolonged		Phosphokinase Increased	Health	(Gabapentin)	PS		
		Liver Function Test	Professional	(Naproxen Sodium)	SS		
		Abnormal		(Acetylsalicylic			
		Suicide Attempt		Acid)	SS		
				(Venlafaxine			
				Hydrochloride)	SS		
				(Paracetamol)	SS		
				(Pseudoephedrine			
				Hydrochloride)	SS		

Date:05/02/02ISR Number: 3911865-1Report Type:Expedited (15-DaCompany Report #055-0945-M0200033

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Asthenia	Foreign	Gabapentin			
Other		Bone Disorder	Consumer	(Gabapentin)	PS		ORAL
1200 MG							
(THREE TIMES		Condition Aggravated					
DAILY),PER		Movement Disorder					

ORAL

Date:05/02/02ISR Number: 3911925-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200245

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Consumer	Neurontin			
PER ORAL		Cognitive Disorder	Health	Gabapentin)	PS		ORAL
		Convulsion	Professional	(Unspecified			
		Electroencephalogram		Decongestant)	C		
		Abnormal					
		Learning Disorder					

Date:05/03/02ISR Number: 3910643-7Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-11811908

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Aortic Aneurysm		Buspirone Hcl	PS	Bristol-Myers Squibb	
Hospitalization -		Coagulopathy				Company	
15mg							
Initial or Prolonged		Death		Percocet Tabs	SS		
2mg capsules		Hepatic Cirrhosis		Loperamide Hcl	SS		
		Multi-Organ Failure		Neurontin	SS		
		Thrombocytopenia		Oxycontin	SS		
				Morphine	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

100mg/650mg
 5mg

Ambien	SS
Propoxyphene Napsylate+Acetaminop hen	SS
Diazepam	SS

Date:05/06/02ISR Number: 3911431-8Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-11843034
 Age:68 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - Initial or Prolonged SUBCUTANEOUS	Dialysis Hiv Wasting Syndrome Hypertension Pancytopenia	Health Professional	Videx T-20/Ro 29-9800	PS SS	Bristol-Myers Squibb Company	ORAL
not reported/inte rupted 15	Renal Failure Acute Urinary Tract Infection					

Feb 02,

Tenofovir	SS	ORAL
Amprenavir	SS	ORAL
Ritonavir	SS	ORAL
Bactrim	SS	
Ketoprofene	SS	

Start date
 unknown

Gabapentin	SS
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Date:05/06/02ISR Number: 3912226-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200529
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Drug Abuser	Health Professional	Neurontin (Gabapentin)	PS		

Date:05/06/02ISR Number: 3912256-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200527
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Breast Cancer					
600 MG (TWICE DAILY), PER ORAL							

Date:05/06/02ISR Number: 3912598-8Report Type:Expedited (15-DaCompany Report #2002103692FR
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Convulsion Hyponatraemia	Foreign Health	Xanax (Alprazolam) Tablet	PS		ORAL
ORAL			Professional Other	Gardenal (Phenobarbital)	SS		ORAL
ORAL				Tegretol (Carbamazepine)	SS		ORAL
ORAL				Neurontin (Gabapentin)	SS		ORAL
				Amlor (Amlodipine Besilate)	C		
				Celectol (Celiprolol)	C		

Freedom Of Information (FOI) Report

Agreal (Veralipride) C
 Calcium
 Carbonate/Colecalcif
 erol C

Date:05/06/02ISR Number: 3912957-3Report Type:Expedited (15-DaCompany Report #055-0945-M0200026
 Age:87 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dehydration	Foreign	Gabapentin			
Hospitalization - 400 MG		Drug Ineffective	Consumer	(Gabapentin)	PS		ORAL
Initial or Prolonged (DAILY), PER		Gallbladder Operation					
ORAL		Malaise					

Date:05/06/02ISR Number: 3912961-5Report Type:Expedited (15-DaCompany Report #044-0945-M0200047
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Depression	Foreign	Neurontin			
PER ORAL		Drug Dependence	Consumer	(Gabapentin)	PS		ORAL
		Fatigue	Health				
		Fluid Retention	Professional				
		Weight Increased					

Date:05/07/02ISR Number: 3913033-6Report Type:Expedited (15-DaCompany Report #044-0945-M0200065
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Blood Pressure Decreased	Foreign	Neurontin			
Initial or Prolonged		Dizziness	Health	(Gabapentin)	PS		
40 MG DAILY		Drug Interaction	Professional	(Lisinopril)	SS		

Date:05/07/02ISR Number: 3913231-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200528
Age:67 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Abdominal Pain	Consumer	Neurontin			
Initial or Prolonged	Back Pain		(Gabapentin)	PS		ORAL
PER ORAL	Difficulty In Walking					
	Nausea					
	Oral Intake Reduced					
	Pain					

Date:05/07/02ISR Number: 3913241-4Report Type:Expedited (15-DaCompany Report #001-0073-M0200194
Age: Gender:Male I/FU:I

Outcome	PT
Other	Agitation
	Akathisia
	Cognitive Disorder
	Deep Vein Thrombosis
	Delusion
	Depression
	Drug Ineffective
	Drug Interaction
	Drug Level Decreased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Oral Intake Reduced Protein Total Increased Tardive Dyskinesia Urine Calcium Increased	Report Source	Product	Role	Manufacturer	Route
400 MG (200 MG, BID), PER ORAL			Health Professional	Dilantin (Phenytoin Sodium)	PS		ORAL
				(Gabapentin)	SS		
				(Fluphenazine Deconoate)	SS		
INTRAMUSCULAR (EVERY 7 DAYS), INTRAMUSCULAR	17.75 MG						
				(Fluphenazine Hydrochloride)	SS		
35 MG (TID & HS)				Warfarin Sodium)	SS		
				(Fluoxetine Hydrochloride)	SS		
8 MG (4 MG, BID),				(Risperidone)	SS		

Date:05/07/02ISR Number: 3913662-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200525
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PER ORAL Other		Brain Neoplasm Diarrhoea	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Clonazepam	C		

Date:05/07/02ISR Number: 3913897-6Report Type:Expedited (15-DaCompany Report #001-0981-M0202246
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaemia Vitamin B12 Deficiency	Health Professional	Atorvastatin (Atorvastatin)	PS		ORAL
PER ORAL		Dyskinesia		Gabapentin	SS		ORAL
900 MG (THREE TIMES DAILY)		Ear Pain					
PER ORAL		Pain In Extremity					
		Parkinsonian Gait Restlessness Tardive Dyskinesia		(Glipizide)	C		

Date:05/07/02ISR Number: 3913963-5Report Type:Expedited (15-DaCompany Report #A209649
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Dizziness	Consumer	Lithane Tablets	PS		ORAL
TID:ORAL		Fatigue		Gabapentin	SS		ORAL
Intervention to 300.00 MG		Nausea					
Prevent Permanent TOTAL:DAILY:O		Osteoporosis					
Impairment/Damage RAL		Parkinson'S Disease		Prevacid Unknown Stool Softener	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/08/02ISR Number: 3912390-4Report Type:Expedited (15-DaCompany Report #B0267013A
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Dehydration	Health	Amprenavir	PS	Glaxo Wellcome	ORAL
Hospitalization - day	7	MON	Professional				
Initial or Prolonged SUBCUTANEOUS	90MG Twice	Diarrhoea		T-20	SS		
per day		Fatigue					
400MG per day	7	MON	Hiv Wasting Syndrome	Didanosine	SS		ORAL
300MG per day	7	MON	Hypertension	Tenofovir	SS		ORAL
200MG per day	5	MON	Pancytopenia	Ritonavir	SS		ORAL
13 YR			Renal Failure Acute	Bactrim	SS	Glaxo Wellcome	ORAL
2400MG per day			Urinary Tract Infection	Ibuprofene	SS	Glaxo Wellcome	ORAL
900MG per day	14	DAY	Vomiting	Gabapentin	SS		

Date:05/08/02ISR Number: 3913269-4Report Type:Direct Company Report #USP 54858
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Medication Error		Metoprolol Tartrate Neurontin (Carbopenbine)	PS SS	Teva Pfizer	

Date:05/08/02ISR Number: 3914276-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200453
 Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 100 MG (DAILY), PER ORAL	Convulsion Somnolence Tremor	Consumer Health Professional	Neurontin Phenytoin Sodium	PS C	ORAL
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Date:05/08/02ISR Number: 3914554-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200425
Age:46 YR Gender:Female I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 1200 MG Initial or Prolonged (DAILY), PER Other ORAL	Adrenal Neoplasm Anxiety Colitis Ischaemic Haematochezia Loss Of Consciousness Orthostatic Hypotension Pain Rosacea Syncope Syncope Vasovagal Weight Decreased	Consumer Health Professional	Neurontin(Gabapentin) Levothyroxine Sodium	PS C		ORAL

Date:05/09/02ISR Number: 3915343-5Report Type:Expedited (15-DaCompany Report #02P-056-0190595-00
Age:1 DY Gender:Male I/FU:F

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
		Clonic Convulsion	Foreign	Depakine Chrono			
		Convulsion Neonatal	Health	Tablets (Depakene)			
		Drug Withdrawal	Professional	(Sodium			
		Convulsions	Other	Valproate/Valproic			
		Dysphagia		Acid) (Sodium	PS		
		Epilepsy		Gabapentin	SS		
		Foetal Distress Syndrome		Clonazepam	SS		
		Maternal Drugs Affecting					
		Foetus					
		Neonatal Apnoeic Attack					
		Neonatal Disorder					
		Sleep Disorder					
		Tonic Clonic Movements					

Date:05/10/02ISR Number: 3914819-4Report Type:Expedited (15-DaCompany Report #033-0945-M0200064

Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Epilepsy	Foreign	Neurontin			
Initial or Prolonged		Hyponatraemia	Health	(Gabapentin)	PS		ORAL
PER ORAL			Professional	(Alprazolam)	SS		ORAL
PER ORAL				(Carbamazepine)	SS		ORAL
PER ORAL				(Phenobarbital)	SS		ORAL
PER ORAL				(Amlodipine			
				Besilate)	C		
				(Celiprolol)	C		
				(Verapride)	C		
				(Calcium Carbonate)	C		
				(Colecalciferol)	C		

Date:05/10/02ISR Number: 3915213-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200369

Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Muscle Spasms Pain	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
300 MG		Weight Decreased	Professional				
(DAILY), PER							
ORAL							

(Oxycodone
Hydrochloride) C
(Levothyroxine
Sodium) C
(Glipizide) C
(Insulin) C
(Atorvastatin) C

Date:05/10/02ISR Number: 3915834-7Report Type:Expedited (15-DaCompany Report #033-0945-M0200041
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dermatitis Bullous Purpura	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Hospitalization - 400 MG		Skin Necrosis	Professional				
Initial or Prolonged (UNKNOWN),							
PER ORAL							

(Hydrocortisone

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Aceponate) C
 (Folic Acid) C
 (Nicotinamide) C
 (Buflomedil
 Hydrochloride) C
 (Ramipril) C
 (Acetylsalicylate
 Lysine) C
 (Thiamine,
 Pyridoxine) C
 (Insulin Lispro) C

Date:05/13/02ISR Number: 3915659-2Report Type:Expedited (15-DaCompany Report #A208483
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abdominal Pain	Consumer	Zoloft Tablets	PS		
		Abdominal Pain Upper		Gabapentin	SS		
		Completed Suicide		Oxycontin	SS		
		Drug Ineffective					
		Gastrointestinal Disorder					
		Gun Shot Wound					
		Medication Error					
		Oral Intake Reduced					

Date:05/13/02ISR Number: 3915935-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200556
 Age:90 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, THREE TIMES DAILY)		Blood Creatinine Abnormal	Health	Neurontin			
		Blood Urea Increased	Professional	(Gabapentin)	PS		
		Cardiac Failure					
		Congestive					
		Creatinine Renal Clearance Decreased Myocardial Infarction		Warfarin Sodium	C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Muscle Spasticity Paraplegia	Health Professional	Neurontin (Gabapentin)	PS		
300 MG							
(DAILY)		Serotonin Syndrome					
				Baclofen	C		
				Ranitidine	C		
				Paroxetine			
				Hydrochloride	C		

Age: Gender:Male I/FU:F

Outcome	PT
Death	Abdominal Pain
Other	Completed Suicide
	Eating Disorder
	Gastrointestinal Disorder
	Gun Shot Wound

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Idiosyncratic Drug Reaction	Report Source	Product	Role	Manufacturer	Route
75 MG,			Consumer	Neurontin (Gabapentin)	PS		
(DAILY)				Sertraline Hydrochloride	SS		
				Oxycodone Hydrochloride	SS		

Date:05/13/02ISR Number: 3915944-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200581
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Neurontin (Gabapentin)	PS		
Other		Drug Interaction Rhabdomyolysis	Professional	(Unspecified Statin)	SS		

Date:05/13/02ISR Number: 3915946-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200580
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Neurontin (Gabapentin)	PS		
Other		Drug Interaction Rhabdomyolysis	Professional	(Pravastatin Sodium)	SS		

Date:05/13/02ISR Number: 3915947-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200579
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Neurontin (Gabapentin)	PS		
Other		Drug Interaction Rhabdomyolysis	Professional	(Pravastatin Sodium)	SS		

Date:05/13/02ISR Number: 3915948-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200578
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Neurontin			
Other		Drug Interaction	Professional	(Gabapentin)	PS		
		Rhabdomyolysis		(Pravastatin Sodium)	SS		

Date:05/13/02ISR Number: 3916232-2Report Type:Expedited (15-DaCompany Report #044-0945-M0200068
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign	Neurontin			
Disability		Movement Disorder	Health	(Gabapentin)	PS		ORAL
600 MG (200		Speech Disorder					
MG, THREE		Tremor	Professional				
TIMES DAILY),							
PER ORAL				(Morphine Sulfate)	C		
				(Phenylephrine			
				Hydrochloride,			
				Paracetamol,			

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Freedom Of Information (FOI) Report

Carbinoxamine
 Maleate) C
 (Imipramine) C
 (Simvastatin) C
 (Bisoprolol) C
 (Carbamazepine) C

Date:05/13/02ISR Number: 3916574-0Report Type:Expedited (15-DaCompany Report #02P-056-0192416-00
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged	200 MG, 1 IN	Dehydration Dialysis Diarrhoea	Foreign Health Professional	Ritonavir Soft Gelatin Capsules (Norvir) (Ritonavir)	PS		
	1 D	Hiv Wasting Syndrome	Other				
	300 MG, 1 IN	Hypertension		Tenofovir	SS		
	1 D	Oliguria					
	1200 MG, 1 IN	Pancytopenia		Amprenavir	SS		
	1 D	Renal Failure Acute					
	400 MG, 1 IN	Urinary Tract Infection		Didanosine	SS		
	1 D	Vomiting					
	1 CAPSULE, 1 IN			Bactrim	SS		
	1 D						
	2600 MG			Ibuprofen	SS		
	900 MG, 1 IN			Gabapentin	SS		
	1 D						
SUBCUTANEOUS	90 MG, 2 IN 1			Ro+29-9800 (T 20-Fusion Inhibitor)	SS		

SUBCUTANEOUS

Date:05/14/02ISR Number: 3916944-0Report Type:Expedited (15-DaCompany Report #A210098

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - ORAL	Accident	Health	Glucotrol Tablets	PS		ORAL
Initial or Prolonged 200 MG	Back Injury	Professional	Celebrex	SS		ORAL
Required	Blood Cholesterol					
TOTAL:DAILY:0	Increased					
Intervention to RAL						
Prevent Permanent 600.00 MG	Blood Glucose Decreased		Neurontin	SS		
Impairment/Damage TOTAL;BID	Coronary Artery Occlusion					
	Diabetic Neuropathy		Glucosamine	C		
	Dizziness		Chondroitin	C		
	Fall		Avandia	C		
	Gait Disturbance		Amitriptyline	C		
	Limb Injury		Zebeta	C		
	Myalgia		Atorvstatin	C		
	Somnolence					
	Vertigo					

Date:05/14/02ISR Number: 3917109-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200582

Age:33 YR Gender:Female I/FU:I

Outcome	PT
Other	Angiopathy
	Blood Cholesterol
	Increased
	Colour Vision Tests
	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
		Diabetes Mellitus					
		Diabetic Eye Disease					
		Diabetic Retinopathy					
		Eye Disorder	Literature	Gabapentin	PS		
		Papilloedema	Health	Bumetanide	SS		
		Pupillary Disorder	Professional	Insulin Human			
		Retinal Detachment		Injection, Isophane	SS		
		Splinter Haemorrhages		Nortriptyline	SS		
		Vision Blurred					
		Visual Acuity Reduced					
		Visual Field Defect					

Date:05/14/02ISR Number: 3917507-3Report Type:Expedited (15-DaCompany Report #033-0945-M0200041
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (UNKNOWN), PER ORAL		Dermatitis Bullous Necrosis Purpura	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				(Hydrocortisone Aceponate)	C		
				(Folic Acid)	C		
				(Nicotinamide)	C		
				(Buflomedil Hydrochloride)	C		
				(Ramipril)	C		
				(Acetylsalicylate Lysine)	C		
				(Thiamine, Pyridoxine)	C		
				(Insulin Lispro)	C		

Date:05/15/02ISR Number: 3917022-7Report Type:Direct Company Report #CTU 168090
 Age:11 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Life-Threatening
 Hospitalization -
 Initial or Prolonged
 Other
 Required
 Intervention to
 Prevent Permanent
 Impairment/Damage

Abnormal Behaviour
 Aggression
 Anger
 Condition Aggravated
 Drug Effect Decreased
 Suicidal Ideation
 Weight Increased

Neurontin PS
 Seroquel SS
 Serevent C
 Singulair C
 Zyrtec C
 Albuterol C

Date:05/15/02ISR Number: 3917119-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200577
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), Other PER ORAL		Muscle Rupture Neuropathy Peripheral Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
				(Levodopa, Carbidopa)	C		
				(Citalopram Hydrobromide)	C		
				(Celecoxib)	C		

Freedom Of Information (FOI) Report

(Levothyroxine Sodium) C
(Hydrochlorothiazide, Lisinopril) C

Date:05/15/02ISR Number: 3917131-2Report Type:Expedited (15-DaCompany Report #044-0945-M0200069
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1200 MG (400	Anorexia Arthralgia	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
MG,TID) PER		Chills	Professional				
ORAL		Fatigue					
		Headache		(Folic Acid, Ferrous Fumarate)	C		
		Joint Swelling		(Zinc Sulfate)	C		
		Musculoskeletal Stiffness		(Levothyroxine Sodium)	C		
		Nausea		(Aluminum Hydroxide)	C		
		Pain In Extremity		(Paracetamol, Dihydrocodeine Bitratrate)	C		
		Purpura		(Calcium Gluconate)	C		
				(Betaxolol Hydrochloride)	C		
				Paroxetine	C		
				(Zopiclone)	C		
				(Ergocalciferol)	C		

Date:05/15/02ISR Number: 3917132-4Report Type:Expedited (15-DaCompany Report #033-0945-M0200069
Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cyanosis	Foreign Health	Neurontin (Gabapentin)	PS		
Other			Professional	(Lamotrigine)	SS		
2400 MG (TID)							
300 MG							

Company
Representative

Date:05/15/02ISR Number: 3917676-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200576

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Cholesterol Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG		Vision Blurred					
(DAILY) PER							
ORAL				Tamoxifen	C		

Date:05/17/02ISR Number: 3918854-1Report Type:Expedited (15-DaCompany Report #034-0945-M0200004

Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged (THREE TIMES		Dyskinesia Haemodialysis	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
DAILY), PER							
ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sildenafil C

Date:05/17/02ISR Number: 3918990-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200592
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 500 MG (TID)		Clonic Convulsion Tremor	Health Professional	Neurontin (Gabapentin)	PS		
				Metoprolol Tartrate	C		
				Glipizide	C		
				Isosorbide			
				Mononitrate	C		
				Warfarin Sodium	C		
				Digoxin	C		
				Lisinopril	C		
				Indometacin	C		
				Fenofibrate	C		
				Simvastatin	C		
				Metolazone	C		
				Furosemide	C		

Date:05/17/02ISR Number: 3918992-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200375
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG (600 MG, TID), PER ORAL 1600 MG (BID)		Hypoaesthesia Muscle Spasms	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Cimetidine	SS		
				Metformin	C		
				Losartan Potassium	C		
				Atorvastatin	C		
				Diltiazem			
				Hydrochloride	C		
				Hydroxyprogesterone	C		

Estrogens Conjugated C
Hydrochlorothiazide C
Insulin Human
Injection, Isophane C
Glibenclamide C
Acetylsalicyclic
Acid C
Ergocalciferol,
Ascorbic Acid, Folic
Acid, Thiamine
Hydrochloride,
Retinol, Riboflavin, C
Ibuprofen C

Date:05/17/02ISR Number: 3918993-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200190
Age: Gender:Female I/FU:I

Outcome PT
Other Amnesia
Asthenia
Balance Disorder
Condition Aggravated

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression Drug Interaction Dysphemia	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Hydrochlorothiazide, Triamterene Levothyroxine Sodium	PS SS SS		
0.1 MG		Multiple Sclerosis Neck Pain Speech Disorder Thought Blocking					
(DAILY)				Desipramine	SS		
300 MG				Fluoxetine Hydrochloride	C		
(DAILY)							

Date:05/17/02ISR Number: 3919060-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200491
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 3600 MG (FOUR TIMES DAILY), PER		Atherosclerosis Cardiomyopathy Drug Level Increased	Consumer Health Professional	Neurontin (Gabapentin)	PS		
				Hydrocodone Clonazepam Methadone Cyclobenzaprine	C C C C		

Date:05/17/02ISR Number: 3919065-6Report Type:Expedited (15-DaCompany Report #001-0945-M0200590
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Diabetes Mellitus	Health	Neurontin			

1800 MG Professional (Gabapentin) PS ORAL
 (TID), PER
 ORAL
 Insulin C
 Amitriptyline C

Date:05/17/02ISR Number: 3919158-3Report Type:Expedited (15-DaCompany Report #032-0945-M0200002
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 300 MG Initial or Prolonged (DAILY),		Respiratory Depression Shock	Foreign Health Professional	Neurontin (Gabapentin)	PS		
				Clonazepam Amitriptyline Hydrochloride	C		
					C		

Date:05/20/02ISR Number: 3919453-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200594
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (THREE TIMES Other DAILY), PER		Rhabdomyolysis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
							ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Simvastatin	SS
Ciclosporin	C
Azathioprine	C
Prednisone	C
Lisinopril	C
Metoprolol	C
Insulin	C
Tocopherol	C
Ergocalciferol	C
Acetylsalicylic Acid	C
Amitriptyline	C

Date:05/20/02ISR Number: 3919454-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200596
 Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Pancreatitis	Health	Neurontin			
Hospitalization - 200 MG Initial or Prolonged (DAILY) Other		Renal Failure	Professional	(Gabapentin)	PS		
				Pravastatin	SS		
				Ciclosporin	C		
				Mycophenolate			
				Mofetil	C		
				Prednisone	C		
				Isosorbide			
				Mononitrate	C		
				Nifedipine	C		
				Famotidine	C		
				Sertraline	C		
				Temazepam	C		
				Acetylsalicylic Acid	C		
				Fexofenadine			
				Hydrochloride	C		

Date:05/20/02ISR Number: 3919599-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200465
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Bone Scan Abnormal	Consumer	Neurontin			

800 MG (BID),	Cardiac Valve Disease	Health	(Gabapentin)	PS	ORAL
PER ORAL	Drug Effect Decreased	Professional			
40 MG	Impaired Gastric Emptying		Atorvastatin	SS	ORAL
(DAILY), PER					
ORAL			Glipizide	C	
			Metformin		
			Hydrochloride	C	
			Mirtazapine	C	
			Paracetamol,		
			Hydrocodone		
			Bitartrate	C	
			Bumetanide	C	
			Potassium Chloride	C	
			Acetylsalicylic Acid	C	
			Ascorbic Acid	C	
			Ergocalciferol,		
			Calcium Phosphate,		
			Calcium Sodium		
			Lactate	C	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ascorbic Acid,
 Tocopheryl Acetate,
 Retinol, Zinc,
 Calcium, Vitamins
 Nos, Minerals Nos, C

Date:05/20/02ISR Number: 3919679-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200532
 Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain	Consumer	Neurontin			
		Bladder Operation	Health	(Gabapentin)	PS		ORAL
1800 MG (600		Fall	Professional				
MG, THREE							
TIMES A DAY),							
PER ORAL							

Tramadol
 Hydrochloride C
 Paracetamol C
 Ibuprofen C
 Acetylsalicylic
 Acid, Magnesium
 Hydroxide, Aluminium
 Hydroxide) C

Date:05/20/02ISR Number: 3919681-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200496
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fall	Consumer	Neurontin			
		Hypersensitivity	Health	(Gabapentin)	PS		ORAL
1800 MG (600		Injection Site Oedema	Professional				
MG, THREE		Injection Site Pain					
TIMES DAILY),		Oedema Peripheral					
PER ORAL							

Pain In Extremity
Peripheral Nerve Injury

Hyaluronate Sodium SS
Tramadol
Hydrochloride C
Paracetamol C
Ibuprofen C
Acetylsalicylic Aid,
Magnesium Hydroxide,
Aluminium Hydroxide) C

Date:05/20/02ISR Number: 3919683-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200456
Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 200 MG (100 MG, BID), PER ORAL	Glossitis Glossodynia Pharyngitis Psychotic Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Quetiapine Clonazepam Lithium Carbonate Benzatropine Mesilate Lorazepam	C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/20/02ISR Number: 3919686-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200326

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 41600 MG PER Other ORAL		Electrocardiogram Qt Corrected Interval Prolonged	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Electrocardiogram Qt Prolonged		Amitriptyline	SS		
		Medication Error		Lamotrigine	C		
		Mental Status Changes Overdose		Reperidal	C		

Date:05/20/02ISR Number: 3919703-8Report Type:Expedited (15-DaCompany Report #061-0945-M0200078

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG (800 MG, TID) PER ORAL		Chest Pain Renal Impairment	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Ramipril	C		
				Acetylsalicylic Acid	C		
				Clopidogrel	C		
				Cabergoline	C		
				Senoside A+B, Docusate	C		
				Pravastatin Sodium	C		
				Glyceryl Trinitrate	C		

Date:05/20/02ISR Number: 3919770-1Report Type:Expedited (15-DaCompany Report #055-0945-M0200038

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Foreign	Gabapentin			

800 MG
 (DAILY), PER
 ORAL

Depression
 Fear
 Tremor

Consumer
 (Gabapentin)

PS
 C

(Phenobarbital)

Date:05/21/02ISR Number: 3919210-2Report Type:Expedited (15-DaCompany Report #WAES 0205USA01611
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dizziness Drug Interaction Hypotension	Health Professional	Prinivil Prinivil Neurontin	PS SS SS	Merck & Co., Inc	ORAL ORAL

Date:05/21/02ISR Number: 3920411-8Report Type:Direct Company Report #CTU 168559
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 100MG PO BID Intervention to Prevent Permanent Impairment/Damage 400 MG/D		Iritis		Topiramate Citalopram Lithium Gabapentin	PS SS SS SS	Ortho-Mcneil	ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/22/02ISR Number: 3920926-2Report Type:Expedited (15-DaCompany Report #2002UW06742
Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	100 MG TID PO	Acidosis	Health	Seroquel	PS		ORAL
Hospitalization - Initial or Prolonged		Cardiotoxicity Grand Mal Convulsion Hypotension Neurotoxicity Overdose	Professional	Neurontin Thyroid Medication Celexa	SS C C		

Date:05/22/02ISR Number: 3921329-7Report Type:Expedited (15-DaCompany Report #001-0073-M0200138
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300 MG (DAILY), PER ORAL	Convulsion Dizziness Fatigue	Consumer Health Professional	Dilantin Suspension 125 Mg/5 Ml (Phenytoin Sodium)	PS		ORAL
		Feeling Abnormal Gingival Hyperplasia Insomnia		Gabapentin	SS		ORAL
	900 MG (TID), PER ORAL	Myalgia Psychomotor Hyperactivity		Phenobarbital	SS		ORAL
	90 MG (DAILY), PER ORAL	Staring					

Date:05/22/02ISR Number: 3921618-6Report Type:Expedited (15-DaCompany Report #001-0945-M0200598
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Arthralgia	Consumer	Neurontin			

Initial or Prolonged 900 MG (300 Other MG, TID)	Cerebrovascular Accident	(Gabapentin)	PS
	Drug Effect Decreased		
	Hypersomnia	(Levothyroxine	
	Hypoaesthesia	Sodium)	C
	Intervertebral Disc Protrusion	(Citalopram Hydrobromide)	C
	Movement Disorder	(Hydrochlorothiazide	
	Paraesthesia	, Lisinopril)	C
	Post Procedural Complication	(Furosemide)	C
	Postoperative Infection	(Potassium Chloride)	C
	Rotator Cuff Syndrome	(Celecoxib)	C
	Tendon Disorder	(Rofecoxib)	C
	Toe Deformity	(Cyanocobalamin)	C

Date:05/22/02ISR Number: 3922179-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200605
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (100 MG, TID), PER ORAL		Blood Pressure Decreased Dysarthria Hypoaesthesia Oral Masked Facies Movement Disorder Muscle Twitching	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Insulin Lispro Insulin Insulin Human Injection, Isophane Irbesartan	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lorazepam C
 Furosemide C
 Glyceryl Trinitrate C
 Magnesium Hydroxide,
 Aluminum Hydroxide
 Gel, Dried,
 Dimeticone,
 Activated) C

Date:05/22/02ISR Number: 3922346-3Report Type:Expedited (15-DaCompany Report #055-0945-M0200042
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness Dyspnoea	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
800 MG		Feeling Abnormal	Professional Company Representative				
(UNKNOWN), PER ORAL							

Date:05/22/02ISR Number: 3922651-0Report Type:Expedited (15-DaCompany Report #055-0945-M0200041
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness Dyspnoea	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
800 MG, PER ORAL			Professional Company Representative				

Date:05/23/02ISR Number: 3921809-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200633
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Abdominal Wall Anomaly	Consumer	Neurontin (Gabapentin)	PS
			Celecoxib	SS

Date:05/23/02ISR Number: 3922097-5Report Type:Expedited (15-DaCompany Report #PHNU2002DE01717
 Age:69 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG/DAY,	Alanine Aminotransferase Increased Aspartate	Foreign Health Professional	Trileptal(Oxcarbazep ine) Film-Coated Tablet	PS		
	Aminotransferase Increased Gamma-Glutamyltransferase Increased	Other	Neurontin(Gabapentin) Novalgin(Metamizole Sodium)	SS SS		ORAL
ORAL	Oedema Pulmonary Oedema		Zyprexa (Olanzapine) Remergil (Mirtazapine) Dytide (Benzthiazide) Dilatrend (Carvedilol)	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/23/02ISR Number: 3922101-4Report Type:Expedited (15-DaCompany Report #045-0945-M0200008

Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 3200 MG, PER Initial or Prolonged ORAL		Chromaturia	Foreign Health	Gabapentin	PS		ORAL
		Encephalitis	Professional	Aciclovir	C		
		Herpes Zoster		Valproate Sodium	C		
		Tetanus					

Date:05/23/02ISR Number: 3922112-9Report Type:Expedited (15-DaCompany Report #044-0945-M0200078

Age:8 DY Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2400 MG (800 MG, TID)		Convulsion Neonatal Drug Exposure Via Breast Milk	Foreign Health Professional	Neurontin (Gabapentin)	PS		
		Drug Withdrawal Syndrome Neonatal Maternal Drugs Affecting Foetus					

Date:05/23/02ISR Number: 3922248-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200613

Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 3600 MG (TID), PER ORAL		Drug Abuser	Health Professional	Neurontin(Gabapentin)	PS		ORAL
				Prevacid	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Angiogram Abnormal Angiogram Retina Abnormal Blindness	Literature	Mexiletine (Mexiletine Hydrochloride)	PS		ORAL
DAILY, PO		Conjunctival Hyperaemia Drug Toxicity		Clonidine (Clonidine)	SS		ORAL
PO		Eye Disorder		Morphine (Morphine)	SS		ORAL
PO		Fundoscopy Abnormal Maculopathy Retinogram Abnormal Retinopathy		Trazodine (Trazodine Hydrochloride) Pentosan Polysulfate (Pentosan Polysulfate) Gabapentin (Gabapentin) Lidocaine Hydrochloride (Lidocaine Hydrochloride)	SS SS SS SS		
INTRAVENOUS	385.7 MG						
(SINGLE DOSE INFUSION), IV							
PO				Baclofen (Baclofen)	SS		ORAL
PO				Methadone Hydrochloride (Methadone Hydrochloride)	SS		ORAL

Freedom Of Information (FOI) Report

Lorazepam
 (Lorazepam) SS
 Hydroxyzine
 Hydrochloride
 Hydroxyzine
 Hydrochloride) SS
 Glimepiride (Oral
 Antidiabetics) SS
 Promethazine
 Hydrochloride
 (Promethazine
 Hydrochloride) SS
 Fluoxetine
 Hydrochloride
 (Fluoxetine
 Hydrochloride) SS
 Sucralfate
 (Sucralfate) SS

Date:05/23/02ISR Number: 3922626-1Report Type:Expedited (15-DaCompany Report #C2002-0958.01
 Age:64 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Activated Partial	Consumer	Buspirone			
Hospitalization -	Thromboplastin Time		Hydrochloride			
Initial or Prolonged	Prolonged		Tablets 15 Mg	PS		
	Aortic Aneurysm		Diazepam Tablets 5			
	Blood Bilirubin Increased		Mg	SS		
	Blood Disorder		Loperamide Hcl			
	Cardiac Arrest		Capsules 2 Mg	SS		
	Coagulopathy		Propoxyphene			
	Diarrhoea		Napsylate And Apap			
	Fungal Infection		Tablets 100 Mg/650mg	SS		
	Gastrointestinal		Neurontin Capsules			
	Haemorrhage		100 Mg Parke Davis	SS	Parke Davis	
	Haematocrit Decreased		Ambien Tablets			
	Haemodialysis		Searle	SS		
	Hepatic Cirrhosis		Oxycodone/Apap			
	Hypoglycaemia		Tablets	SS		
	Hypotension		Oxycontin Tablets	SS		
	International Normalised		Morphine Tablets	SS		
	Ratio Increased		Acetaminphen	C		
	Liver Function Test		Penicillin	C		
	Abnormal		Amlodipine	C		

Low Cardiac Output
Syndrome
Multi-Organ Failure
Oedema
Splenomegaly
Thrombocytopenia
Vasodilatation

Nephrocaps (Qd) C
Lidocaine/Prilocaine C
Cream C
Folic Acid C
Omeprazole C
Cetirizine C
Capsules C
Sucralfate C

Date:05/24/02ISR Number: 3923114-9Report Type:Expedited (15-DaCompany Report #WAES 0205USA01611
Age: Gender:Male I/FU:I

Outcome PT
Hospitalization - Dizziness
Initial or Prolonged Drug Interaction

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hypotension

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
20 MG/DAILY		Foreign Other	Tab Prinivil (Lisinopril)	PS		ORAL
PO ; 40						
MG/DAILY PO			Neurontin (Gabapentin)	SS		

Date:05/24/02ISR Number: 3923157-5Report Type:Expedited (15-DaCompany Report #WAES 0205USA01611
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE IMAGE		Dizziness Drug Interaction	Foreign Health	Tab Prinivil (Lisinopril)	PS		ORAL
		Hypotension	Professional Other	Neurontin (Gabapentin)	SS		

Date:05/28/02ISR Number: 3923298-2Report Type:Expedited (15-DaCompany Report #B0267013A
Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 1200MG per Hospitalization - day 7 MON		Angiosclerosis	Health	Amprenavir	PS	Glaxo Wellcome	ORAL
Initial or Prolonged SUBCUTANEOUS 90MG Twice per day		Anuria Cachexia Confusional State	Professional	T-20	SS		
400MG per day 7 MON		Dehydration		Didanosine	SS		ORAL
300MG per day 7 MON		Dialysis		Tenofovir	SS		ORAL

200MG per day	5	MON	Diarrhoea	Ritonavir	SS		ORAL
13 YR			Fatigue	Bactrim	SS	Glaxo Wellcome	ORAL
2400MG per day			Food Intolerance	Ibuprofene	SS	Glaxo Wellcome	ORAL
			Gastroenteritis				
900MG per day	14	DAY	Hypertension	Gabapentin	SS		
			Pancytopenia				
			Pyrexia				
			Renal Failure Acute				
			Renal Tubular Necrosis				
			Staphylococcal Infection				
			Urinary Tract Infection				
			Vomiting				

Date:05/28/02ISR Number: 3924318-1Report Type:Expedited (15-DaCompany Report #A211461
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Diabetes Mellitus	Consumer	Glucotrol Tablets	PS		
Intervention to				Lipitor	SS		
Prevent Permanent				Neurontin	SS		
Impairment/Damage							

Date:05/28/02ISR Number: 3924332-6Report Type:Expedited (15-DaCompany Report #001-0945-M0200621
Age:72 YR Gender:Female I/FU:I

Outcome	PT
Other	Balance Disorder
	Clumsiness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Condition Aggravated Disturbance In Attention Drug Effect Decreased	Report Source	Product	Role	Manufacturer	Route
400 MG (FOUR TIMES DAILY) PER ORAL		Impaired Driving Ability Oedema Peripheral Visual Acuity Reduced	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				(Diclofenac Sodium, Misoprostol)	C		
				(Enalapril Maleate)	C		
				(Hydrochlorothiazide , Enalapril Maleate)	C		
				(Amlodipine Besilate)	C		
				(Estradiol)	C		
				(Cyanocobalamin)	C		
				(Chlorpheniramine)	C		
				(Citracel With Magensium And D)	C		
				(Glucosamine And Chondroitin With Msm)	C		
				(Flaxseed Oil)	C		
				(Evening Primrose Oil)	C		
				(Tocopherol)	C		
				(Garlicscon)	C		
				(Cetirizine Hydrochloride)	C		
				(Clonazepam)	C		

Date:05/28/02ISR Number: 3924336-3Report Type:Expedited (15-DaCompany Report #001-0945-M200195

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN, (UNKNOWN),		Parkinson'S Disease	Health Professional	Neurontin (Gabapentin)	PS		

UNKNOWN

(Sertraline
Hydrochloride) C

Date:05/28/02ISR Number: 3924340-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200491

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Atherosclerosis	Consumer	Neurontin			
Other		Myocardial Fibrosis	Health	(Gabapentin)	PS		ORAL
3600 MG (FOUR TIMES DAILY), PER ORAL		Myocardial Infarction	Professional				
		Overdose					

(Hydrocodone) C
 (Clonazepam) C
 (Methadone) C
 (Fluoxetine
 Hydrochloride) C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/02ISR Number: 3924413-7Report Type:Expedited (15-DaCompany Report #2002-DE-01032GD (0)

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blindness Retinal Degeneration Retinopathy Scotoma	Literature	Mexiletine (Mexiletine Hydrochloride) (Nr) (Mexiletine-Hcl)	PS		ORAL
PO		Toxicologic Test Abnormal		Clonidine (Clonidine) (Nr) (Clonidine-Hcl)	SS		ORAL
PO				Morphine (Morphine) (Nr) (Morphine-Hcl)	SS		ORAL
PO				Trazodone (Trazodone Hydrochloride) (Nr) (Trazodone-Hcl)	SS		
				Pentosan Polysulfate (Pentosan Polysulfate)	SS		
				Gabapentin (Gabapentin) (Nr)	SS		
				Lidocaine Hydrochloride (Lidocaine Hydrochloride)	SS		
INTRAVENOUS	385.27 MG						
(SINGLE DOSE							
INFUSION) IV							
PO				Baclofen (Baclofen) (Nr)	SS		ORAL
PO				Methadone Hydrochloride (Methadone Hydrochloride)	SS		ORAL
				Lorazepam (Lorazepam) (Nr)	SS		
				Hydroxyzine Hydrochloride (Hydroxyzine			

Hydrochloride) (Nr) SS
 Glimepiride (Oral
 Antidiabetics) (Nr) SS
 Promethazine
 Hydrochloride
 (Promethazine
 Hydrochloride) SS
 Sucralfate
 (Sucralfate) (Nr) SS

Date:05/29/02ISR Number: 3925901-XReport Type:Expedited (15-DaCompany Report #033-0945-M0200060

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG (600 MG, QID)		Urticaria	Foreign Health Professional	Neurontin (Gabapentin)	PS		
				Clonazepam	C		
				Cetirizine	C		
				Loratadine	C		
				Hydroxyzine	C		
				Beclometasone	C		
				Salmeterol	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Matelukast	C
Caffeine, Opium,	
Paracetamol	C
Sertraline	C
Medrogestone	C

Date:05/29/02ISR Number: 3926034-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200393
 Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dialysis	Consumer	Neurontin			
		Disease Progression	Health	(Gabapentin)	PS		ORAL
900 MG (TID),		Renal Failure	Professional				
PER ORAL							

(Folic Acid,	
Vitamins Nos)	C
(Ascorbic Acid)	C
(Ferrous Sulfate)	C
(Calcium Acetate)	C

Date:05/29/02ISR Number: 3926035-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200394
 Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Chest Pain	Consumer	Neurontin			
Initial or Prolonged		Mitral Valve Incompetence	Health	(Gabapentin)	PS		ORAL
900 MG (TID),		Pericarditis	Professional				
PER ORAL							

(Folic Acid,	
Vitamins Nos)	C
(Ascorbic Acid)	C
(Ferrous Sulfate)	C
(Calcium Acetate)	C
Vitamins Unspecified	C

Date:05/29/02ISR Number: 3926041-6Report Type:Expedited (15-DaCompany Report #001-0981-M0202719
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetes Mellitus Diabetic Complication	Consumer	Atorvastatin (Atorvastatin) Gabapentin Glipizide	PS SS SS		

Date:05/29/02ISR Number: 3926074-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200622
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Other		Oedema Peripheral Sensory Disturbance	Consumer	Neurontin (Gabapentin)	PS		

Date:05/29/02ISR Number: 3926286-5Report Type:Expedited (15-DaCompany Report #061-0945-M0200082
Age: Gender:Male I/FU:I

Outcome	PT	Report Source
Other	Somnolence	Foreign Health

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Professional Company Representative	Product	Role	Manufacturer	Route
300 MG	(DAILY), PER		Neurontin (Gabapentin)	PS		ORAL
300 MG	(DAILY), PER		Morphine	SS		ORAL

Date:05/29/02ISR Number: 3926287-7Report Type:Expedited (15-DaCompany Report #049-0945-M0200058
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG	(DAILY), PER	Fatigue Generalised Erythema Gouty Arthritis Headache	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Hyperaesthesia Hypertension Oedema Peripheral Osteoarthritis Overweight Pain In Extremity Weight Increased		Doxepin Hydrochloride	C		

Date:05/29/02ISR Number: 3926289-0Report Type:Expedited (15-DaCompany Report #044-0945-M0200071
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization -	Convulsion	Foreign	Neurontin		
Initial or Prolonged	Medication Error	Health	(Gabapentin)	PS	ORAL
PER ORAL					
Other	Tic	Professional	Piroxicam	C	
			Tramadol	C	

Date:05/30/02ISR Number: 3927551-8Report Type:Expedited (15-DaCompany Report #044-0945-M02000079
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cognitive Disorder	Foreign	Neurontin			
		Renal Failure	Health	(Gabapentin)	PS		
900 MG, DAILY			Professional				

Date:05/30/02ISR Number: 3927553-1Report Type:Expedited (15-DaCompany Report #031-0945-M0200011
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Epilepsy	Foreign	Neurontin			
Other			Health	(Gabapentin)	PS		
			Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/30/02ISR Number: 3927826-2Report Type:Expedited (15-DaCompany Report #044-0945-M0200065

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (DAILY)	Blood Pressure Decreased Dizziness	Foreign Health Professional	Neurontin (Gabapentin)	PS		
40 MG (DAILY)			Lisinopril	SS		

Date:05/30/02ISR Number: 3927953-XReport Type:Expedited (15-DaCompany Report #2002GB01238

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 40 MG DAILY Initial or Prolonged	Dizziness Drug Interaction Hypotension	Other	Lisinopril Neurontin	PS SS		

Date:06/03/02ISR Number: 3928317-5Report Type:Expedited (15-DaCompany Report #351-0945-M0200002

Age:53 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 3600 MG (DAILY), PER ORAL	Hepatic Failure	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Omeprazole Clonazepam	C C		

Date:06/03/02ISR Number: 3928319-9Report Type:Expedited (15-DaCompany Report #044-0945-M0200052

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1500 MG		Alpha-1 Anti-Trypsin Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER		Hepatitis	Professional				
ORAL		Hepatomegaly	Company				
		Hypoglycaemia Vomiting	Representative	Amitriptyline Levothyroxine Sodium Dihydrocodeine Paracetamol Insulin Human Insulin Human Zinc Suspension	C C C C C C		

Date:06/03/02ISR Number: 3928362-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200649
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Liver Function Test Abnormal	Health Professional	Neurontin (Gabapentin)	PS		

Date:06/03/02ISR Number: 3928386-2Report Type:Expedited (15-DaCompany Report #001-0719-M0100408
Age: Gender:Male I/FU:F

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
1200 MG		Actinic Keratosis	Consumer	Lopid (Gemfibrozil)	PS		ORAL
(BID), PER		Arthralgia	Health				
ORAL		Asthenia	Professional				
ORAL		Back Pain		Gabapentin	SS		ORAL
5 MG (DAILY),		Basal Cell Carcinoma		Amlodipine Besilate	SS		ORAL
PER ORAL		Cerebrovascular Accident					
5 MG (DAILY),		Difficulty In Walking		Glipizide	SS		ORAL
PER ORAL		Disorientation					
SUBLINGUAL	10 MG,	Dizziness		Nifedipine	SS		
SUBLINGUAL		Headache					
		Myalgia		Unspecified			
		Osteoarthritis		Cholesterol Lowering			
		Pain		Drug	SS		
		Rash Papular		Furosemide	C		
		Recurrent Cancer		Acetylsalicylic Acid	C		
		Scar		Ergocalciferol,			
		Skin Irritation		Ascorbic Acid, Folic			
		Skin Papilloma		Acid, Thiamine			
		Tenderness		Hydrochloride,			
		Varicose Vein		Retinol, Riboflavin,	C		
		Vein Disorder		Terazosin	C		
		Wound		Labetalol			
				Hydrochloride	C		
				Sunastatin	C		
				Clopidogrel	C		
				Meloxicam	C		
				Pericad Plus			
				Vitamins	C		
				L-Carnitine	C		
				Magnesium	C		
				Glucosamine			
				Chondroitin	C		
				Msm	C		

Celecoxib C
Tamsulosin C
Hydrochloride C
Tolterodine Tartrate C

Date:06/03/02ISR Number: 3928593-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200643
Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Consumer	Neurontin			
		Arthritis		(Gabapentin)	PS		ORAL
1200 MG							
(DAILY), PER		Cartilage Injury					
ORAL		Decreased Activity					
		Movement Disorder		Estrogens Conjugated	C		
		Pain		Levothyroxine Sodium	C		
		Pain In Extremity		Vitamin	C		

Date:06/03/02ISR Number: 3928680-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200641
Age: Gender:Male I/FU:I

Outcome	PT
Other	Affective Disorder
	Asthenia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2400 MG	(DAILY)	Consumer	Neurontin (Gabapentin)	PS		
			Sertraline Hydrochloride	SS		

Date:06/03/02ISR Number: 3928714-8Report Type:Expedited (15-DaCompany Report #A211484
Age: Gender:Male I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 2400.00 MG Prevent Permanent TOTAL:DAILY Impairment/Damage		Consumer	Zoloft Tablets Gabapentin	PS SS		

Date:06/04/02ISR Number: 3928796-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100872
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Other 200 MG (DAILY)		Health Professional	Neurontin (Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthma	Consumer	Neurontin			
Other		Back Pain		(Gabapentin)	PS		ORAL
1200 MG		Conversion Disorder					
(TID), PER		Cough					
ORAL		Nephritis		Topiramate	SS		ORAL
200 MG (BID),		Nocturia					
PER ORAL		Oliguria		Ramipril	SS		ORAL
PER ORAL		Pain		Salbutamol	C		
		Pollakiuria					
		Protein Urine Present					
		Renal Disorder					
		Urinary Incontinence					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/04/02ISR Number: 3929094-4Report Type:Expedited (15-DaCompany Report #A208219

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Balance Disorder	Consumer	Zyrtec Tablets	PS		
Initial or Prolonged	Buttock Pain		Neurontin	SS		ORAL
900.00 MG						
Required	Dizziness					
TOTAL: DAILY:						
Intervention to	Eyelid Ptosis					
ORAL						
Prevent Permanent	Fatigue					
Impairment/Damage	Memory Impairment					
	Road Traffic Accident					
	Spinal Fusion Acquired					

Date:06/04/02ISR Number: 3929100-7Report Type:Expedited (15-DaCompany Report #A209649

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Required	Balance Disorder	Consumer	Lithane Tablets	PS		ORAL
TID: ORAL						
Intervention to	Bone Disorder	Health	Gabapentin	SS		ORAL
300.00 MG						
Prevent Permanent	Confusional State	Professional				
TOTAL: DAILY:						
Impairment/Damage	Dizziness					
ORAL						
	Fatigue		Prevacid	C		
	Nausea		Unknown Stool			
	Osteoporosis		Softener	C		
	Parkinson'S Disease		Welbutrin Sr	C		
	Tremor					

Date:06/04/02ISR Number: 3929186-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200656

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Aggression	Consumer	Neurontin			

Initial or Prolonged	Anxiety	(Gabapentin)	PS	ORAL
2400 MG (TID)				
Other	Drug Ineffective			
PER ORAL				
		Paracetamol,		
		Hydrocodone		
		Bitartrate	C	
		Lorazepam	C	

Date:06/04/02ISR Number: 3929356-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Brain Damage	Consumer	Neurontin			
Other		Depression		(Gabapentin)	PS		ORAL
1800 MG							
(DAILY), PER		Drug Hypersensitivity					
ORAL		Drug Level Above					
		Therapeutic		Lithium Carbonate	C		
		Speech Disorder		Clonazepam	C		
		Tongue Oedema		Methylphenidate			
				Hydrochloride	C		
				Levothyroxine Sodium	C		
				Liothyroxine Sodium	C		
				Sertraline			
				Hydrochloride	C		
				Pilocarpine			
				Hydrochloride	C		
				Metoprolol Succinate	C		
				Lansoprazole	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hyoscyamine Sulfate C
 Diltiazem
 Hydrochloride C

Date:06/05/02ISR Number: 3928458-2Report Type:Direct Company Report #CTU 169562
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG PO TID		Abdominal Pain		Neurontin 300mg	PS		ORAL
Initial or Prolonged		Cholecystectomy Gastrointestinal Disorder		Tamoxifen	C		

Date:06/05/02ISR Number: 3929530-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200662
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cerebrovascular Accident	Consumer	Neurontin (Gabapentin)	PS		

Date:06/05/02ISR Number: 3930705-8Report Type:Expedited (15-DaCompany Report #001-0981-M0202875
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Crying Drug Interaction	Consumer	Atorvastatin (Atorvastatin)	PS		ORAL
10 MG (DAILY), PER ORAL		Feeling Abnormal Suicidal Ideation Tremor		Gabapentin Lithium Paroxetine Hydrochloride Trazadone Buspirone Enalapril	SS SS SS SS SS C		

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Blood Creatinine	Health	Neurontin			
Initial or Prolonged	Increased	Professional	(Gabapentin)	PS		
500 MG (TID)						
Other	Blood Urea Increased		Simvastatin	SS		
	Cardiac Failure		Fenofibrate	SS		
	Congestive		Metoprolol Tartrate	C		
	Clonic Convulsion		Glipizide	C		
	Condition Aggravated		Isosorbide			
	Hyperreflexia		Mononitrate	C		
	Hyporeflexia		Warfarin Sodium	C		
	Muscle Twitching		Digoxin	C		
	Sensory Disturbance		Lisinopril	C		
	Tremor		Indometacin	C		
			Metolazone	C		
			Furosemide	C		
			Acetylsalicylic Acid	C		
			Tocopherol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/05/02ISR Number: 3930719-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200500

Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Cholesterol Abnormal	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
1400 MG		Blood Follicle Stimulating Hormone	Professional				
(DAILY), PER							
ORAL		Abnormal		Carbamazepine	SS		
1200 MG		Drug Effect Decreased					
(DAILY)		Menstruation Irregular					

Date:06/06/02ISR Number: 3929784-3Report Type:Expedited (15-DaCompany Report #049-0945-M0200043

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Visual Acuity Reduced	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		Visual Field Defect	Professional				
1800 MG,							
DAILY, PER							
ORAL				Carbamazepine	SS		ORAL
DAILY, PER							
ORAL							

Date:06/06/02ISR Number: 3929786-7Report Type:Expedited (15-DaCompany Report #041-0945-M0200013

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Convulsion	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
100 MG							

(DAILY), PER	Loss Of Consciousness	Professional			
ORAL					
20 MG			Clomipramine Hydrochloride	SS	ORAL
(DAILY), PER					
ORAL					
20 MG			Fluoxetine Hydrochloride	SS	ORAL
(DAILY), PER					
ORAL					
			Alprazolam	C	
			Propranolol Hydrochloride	C	
			Perindopril	C	

Date:06/06/02ISR Number: 3929788-0Report Type:Expedited (15-DaCompany Report #044-0945-M0200083
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neuroleptic Malignant Syndrome	Foreign Health	Neurontin (Gabapentin)	PS		
1200 MG			Professional				
(DAILY),							
1200 MG				Lithium	SS		
(DAILY),				...	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/06/02ISR Number: 3930394-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200428

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	1800 MG	Arthralgia	Health	Neurontin			
		Balance Disorder	Professional	(Gabapentin)	PS		ORAL
		Insomnia					
(TID), PER		Joint Stiffness					
ORAL		Memory Impairment		Medroxyprogesterone			
		Oedema Peripheral		Acetate, Estrogens			
		Sedation		Conjugated)	C		
				Medroxyprogesterone			
				Acetate	C		
				Paracetamol,			
				Hydrocodone			
				Bitartrate	C		
				Paracetamol,			
				Oxycodone			
				Hydrochloride	C		

Date:06/06/02ISR Number: 3930400-5Report Type:Expedited (15-DaCompany Report #A211484

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required	2400.00 MG	Asthenia	Consumer	Zoloft Tablets	PS		
Intervention to		Balance Disorder		Gabapentin	SS		ORAL
Prevent Permanent		Coordination Abnormal					
TOTAL:DAILY:0		Dizziness					
Impairment/Damage		Eye Irritation					
RAL		Eye Movement Disorder					
		Fall					
		Fatigue					
		Frequent Bowel Movements					
		Haematochezia					
		Incoherent					
		Muscle Atrophy					
		Muscle Spasms					

Nervous System Disorder
Thinking Abnormal
Vision Blurred

Date:06/06/02ISR Number: 3930436-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200641
Age: Gender:Male I/FU:F

Outcome PT
Other Asthenia
Central Nervous System
Neoplasm
Dizziness
Drug Ineffective
Eye Irritation
Eye Movement Disorder
Fall
Fatigue
Frequent Bowel Movements
Haematochezia
Haemorrhoids
Incoherent
Muscle Spasms
Thinking Abnormal

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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
2400 MG		Consumer	Neurontin (Gabapentin)	PS		ORAL
(BID), PER						
ORAL			Sertraline Hydrochloride	SS		

Date:06/06/02ISR Number: 3948074-6Report Type:Periodic Company Report #254373
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Accutane Capsules (Isotretinoin)	PS		ORAL
Other		Abdominal Distension Anorexia					
1 PER DAY		Constipation					
ORAL		Dry Skin Inflammatory Bowel Disease Lip Dry Weight Decreased		Neurontin (Gabapentin) Wellbutrin (Bupropion Hydrochloride)	SS C		

Date:06/07/02ISR Number: 3929383-3Report Type:Expedited (15-DaCompany Report #314251
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Saquinavir	PS	Roche	
Congenital Anomaly		Atrial Septal Defect Congenital Cardiovascular Anomaly		Norvir Epivir Ziagen Diflucan Neurontin	SS SS SS SS SS		

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Activated Partial Thromboplastin Time Prolonged	Health Professional	Ambien - Zolpidem Tartrate - Tablet - Unit Dose : Unknown	PS		ORAL
ORAL		Aortic Aneurysm Cardiac Arrest Cardiomegaly		Buspirone Hydrochloride - Unknown - 15 Mg	SS		
UNKNOWN		Coagulopathy Hypoglycaemia Hypotension Intestinal Ischaemia Lung Disorder Multi-Organ Failure Oedema Overdose Pneumonia Renal Failure Chronic Splenomegaly Thrombocytopenia Vasodilatation		Diazepam - Form : Unknown - 5 Mg Loperamide Hydrochloride - Form : Unknown - 2 Mg Propoxyphene Napsylate W/ Acetaminophene - Form: Unknown - 100 Mg Propoxyphene Napsylate W/ Acetaminophene - Form: Unknown - 650	SS SS		

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Freedom Of Information (FOI) Report

Mg SS
 Neurontin -
 Gabapentin - Form:
 Unknown - Unit Dose
 : Unknown SS
 Percocet - Form :
 Unknown - Unit Dose
 : Unknown SS
 Oxycontin -
 Oxycodone
 Hydrochloride - Dose
 : Ni SS
 Morphine - Form :
 Unknown - Unit Dose
 : Unknown SS

Date:06/07/02ISR Number: 3930780-0Report Type:Expedited (15-DaCompany Report #001-0073-M0200250
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Asthenia Chest Pain	Consumer	Dilantin Infatabs 50 Mg (Phenytoin Sodium)	PS		ORAL
300 MG (THREE TIMES DAILY) PER ORAL		Convulsion Crying					
400 MG (DAILY), PER ORAL		Feeling Abnormal Headache Insomnia		Gabapentin	SS		ORAL
PER ORAL		Irritable Bowel Syndrome		Lorazepam	SS		ORAL
1200 MG (DAILY) PER ORAL		Myocardial Infarction Nausea Nervousness		Vitamin C With D	SS		ORAL
PER ORAL		Osteoporosis		Alendronate Sodium	SS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
PER ORAL				Lamotrigine	SS		ORAL
				Lorazepam	C		
Date:06/07/02ISR Number: 3930783-6Report Type:Expedited (15-DaCompany Report #001-0945-M0200668							
Age: Gender:Male I/FU:I							
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Drug Abuser	Consumer	Neurontin			
Initial or Prolonged		Toxicologic Test Abnormal		(Gabapentin)	PS		ORAL
PER ORAL							
Other				Phenytoin Sodium	SS		ORAL
PER ORAL							
Date:06/07/02ISR Number: 3930787-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200466							
Age: Gender:Female I/FU:F							
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
		Fall	Health	(Gabapentin)	PS		ORAL
900 MG (BID),		Hypertension	Professional				
PER ORAL		Weight Increased		Paracetamol,			
				Oxycodone			
				Hydrochloride,			
				Oxycodone			
				Terephthalate	C		
				Furosemide	C		
				Irbesartan	C		
				Glipizide	C		
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Acetylsalicylic Acid C
 Clopidogrel C
 Chlordiazepoxide
 Hydrochloride C
 Atorvastatin C
 Potassium C
 Metoclopramide C

Date:06/07/02ISR Number: 3931074-XReport Type:Expedited (15-DaCompany Report #049-0945-M0200043
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Eye Pain	Foreign	Neurontin			
Other		Headache	Health	(Gabapentin)	PS		ORAL
1800 MG							
(DAILY), PER		Liver Function Test	Professional				
ORAL		Abnormal					
(DAILY), PER		Post Procedural Pain		(Carbamazepine)	SS		ORAL
ORAL		Trigeminal Neuralgia					
		Visual Acuity Reduced					
		Visual Disturbance					
		Visual Field Defect					

Date:06/10/02ISR Number: 3930921-5Report Type:Direct Company Report #CTU 169797
 Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Tremor		Gabapentin	PS		

Date:06/10/02ISR Number: 3931225-7Report Type:Expedited (15-DaCompany Report #02P-056-0190956-00
 Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abdominal Pain Upper	Foreign	Kaletra Soft Gelatin			

Initial or Prolonged	Alanine Aminotransferase	Health	Capsules (Kaletra)	
	Increased	Professional	(Lopinavir/Ritonavir	
	Aspartate)	
	Aminotransferase		(Lopinavir/Ritonavir	PS
1 DOSAGE				ORAL
	Increased			
FORMS, 2 IN 1				
	Blood Alkaline			
D, PER ORAL				
	Phosphatase Increased		Gabapentin	SS
500 MG, 3 IN				ORAL
	Cholestasis			
11D, PER ORAL				
	Gamma-Glutamyltransferase		Zidovudine	
	Increased		W/Lamivudine	SS
1 DOSAGE				ORAL
	Ichthyosis Acquired			
FORMS, 2 IN 1				
	Jaundice			
D, PER ORAL				
	Pancreatitis		Calcium Folate	C
	Rash Maculo-Papular		Pyrimethamine	C
			Sulfadiazine	C

Date:06/10/02ISR Number: 3931338-XReport Type:Expedited (15-DaCompany Report #044-0945-M0200079

Age: Gender:Male I/FU:F

Outcome PT
Other Agitation
Cognitive Disorder
Confusional State
Hallucination

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Freedom Of Information (FOI) Report

Renal Failure

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG (TID), PER ORAL		Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
(DAILY), PER ORAL			(Paroxetine Hydrochloridie)	SS		ORAL
			(Prednisolone)	C		
			(Unspecified Beta Blockers)	C		
			(Warfarin)	C		
			(Ranitidine)	C		
			(Temazepam)	C		

Date:06/10/02ISR Number: 3931355-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200676
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG (DAILY), PER ORAL		Condition Aggravated Difficulty In Walking Fatigue Lymphoedema	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Pain Swelling		Metaxalone Doxazosin Mesilate Alprazolam	C C C		

Date:06/10/02ISR Number: 3931358-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200667
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Fall	Health	Neurontin			

Initial or Prolonged Head Injury Professional (Gabapentin) PS ORAL
100 MG

(DAILY), PER

ORAL

Nifedipine C
Captopril C
Labetalol C

Date:06/11/02ISR Number: 3933474-0Report Type:Expedited (15-DaCompany Report #02P-163-0193776-00
Age:1 DY Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Atrial Septal Defect	Health	Norvir (Ritonavir)			
		Cardiac Aneurysm	Professional	(Ritonavir)	PS		
		Maternal Drugs Affecting Foetus	Other	Saquinavir	SS		
				Lamivudine	SS		
				Abacavir Sulfate	SS		
				Fluconazole	SS		
				Gabapentin	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/11/02ISR Number: 3933489-2Report Type:Expedited (15-DaCompany Report #314251

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Atrial Septal Defect Cardiac Aneurysm Complications Of Maternal	Health Professional	Saquinavir (Saquinavir Mesylate)	PS		ORAL
800 MG DAILY		Exposure To Therapeutic					
ORAL		Drugs		Norvir (Ritonavir)	SS		ORAL
800 MG DAILY		Maternal Drugs Affecting					
ORAL		Foetus		Epivir (Lamivudine)	SS		ORAL
300 MG DAILY		Pregnancy					
ORAL				Ziagen (Abacavir Sulfate)	SS		ORAL
600 MG DAILY							
ORAL				Diflucan (Fluconazole)	SS		ORAL
200 MG 1 PER							
PRN ORAL				Neurontin (Gabapentin)	SS		ORAL
300 MG DAILY							
ORAL							

Date:06/11/02ISR Number: 3933978-0Report Type:Expedited (15-DaCompany Report #044-0945-M0200085

Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Angiopathy Drug Interaction	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
PER ORAL		Pallor	Professional	Perindopril	SS		ORAL
PER ORAL							

Shock
Syncope

Insulin

C

Date:06/12/02ISR Number: 3932851-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200681
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Pain In Extremity	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG TID) PER ORAL				(Naproxen)	C		

Date:06/13/02ISR Number: 3935035-6Report Type:Expedited (15-DaCompany Report #033-0945-M0200069
Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cyanosis	Foreign	Neurontin			
Other		Electroencephalogram	Health	(Gabapentin)	PS		ORAL
2400MG (TID), PER ORAL		Abnormal	Professional				
350 MG (BID), PER ORAL		Epilepsy	Company Representative	Lamotrigine	SS		ORAL

Date:06/13/02ISR Number: 3935039-3Report Type:Expedited (15-DaCompany Report #046-0945-M0200011
Age: Gender:Male I/FU:I

Outcome	PT
Other	Atrial Fibrillation Cardiac Flutter

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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
1200 MG		Haemoglobin Decreased	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
(TID), PER							
ORAL							

				Citalopram Hydrobromide	C		
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Date:06/13/02ISR Number: 3935059-9Report Type:Expedited (15-DaCompany Report #045-0945-M0200008
 Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	3200 MG, PER	Chromaturia Crystal Urine Present	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
ORAL		Encephalitis					
		Herpes Zoster Infection Neurological		Valproate Sodium	C		
				Fentanyl	C		
				Propofol	C		
				Aciclovir	C		

Date:06/14/02ISR Number: 3934539-XReport Type:Expedited (15-DaCompany Report #A213023
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 900.00 MG		Unevaluable Event	Consumer	Lithane	PS		
Initial or Prolonged							
TOTAL							
Other 600.00 MG				Gabapentin	SS		

TOTAL:DAILY

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain Convulsion	Consumer	Neurontin(Gabapentin)	PS		ORAL
1200		Disease Recurrence					
MG(DAILY),		Gait Disturbance					
PER ORAL		Oral Infection Pleural Infection Sepsis		Solumedrol Ceftriaxone Sodium Imipenem, Ciclastatin Sodium Cetirizine Hydrochloride Fexofenadine Hydrochloride Salbutamol Advair Diskus Triamcinolone Acetonide Nasaryl Rhinacort Trazodone Pantoprazole Lansoprazole Celecoxib Tessalon Pearls	C C C C C C C C C C C C C C C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Plaquinil C
 Ultracet C
 Paracetamol C
 Estradiol C
 Medroxyprogesterone
 Acetate C
 Cefprozil C
 Levoquin C
 Immuneoglobulin G C
 Unspecified Over The
 Counter Migraine
 Medication) C

Date:06/17/02ISR Number: 3934821-6Report Type:Expedited (15-DaCompany Report #001-0719-M0100408
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1200 MG Initial or Prolonged (BID), PER Other ORAL	Acrochordon	Consumer	Lopid	PS		ORAL
PER ORAL	Arthralgia	Health				
PER ORAL	Arthritis	Professional				
5 MG (DAILY), PER ORAL	Basal Cell Carcinoma		Gabapentin	SS		ORAL
5 MG (DAILY), PER ORAL	Blood Pressure Systolic Increased		Amlodipine Besilate	SS		ORAL
5 MG (DAILY), PER ORAL	Bone Pain		Glipizide	SS		ORAL
PER ORAL	Ear Discomfort					
SUBLINGUAL SUBLINGUAL	10 MG Hyporeflexia Insomnia		Nifedipine	SS		
	Joint Swelling		Unspecified			
	Lichen Planus		Cholesterol Lowering Drug	SS		
	Otitis Externa		Furosemide	C		
	Pharyngolaryngeal Pain		Acetylsalicylic Acid	C		
	Scar		Ergocalciferol, Ascorbic Acid, Folic			
	Sensory Disturbance					
	Skin Atrophy					

Skin Irritation	Acid, Thiamine	
Skin Papilloma	Hydrochloride,	
Sleep Apnoea Syndrome	Retinol, Riboflavin,	C
Tenderness	Terazosin	C
Thermal Burn	Labetalol	
Upper Respiratory Tract	Hydrochloride	C
Infection	Sunasatin	C
Varicose Vein	Clopidogrel	C
Vein Disorder	Meloxicam	C
Viral Infection	Pericad Plus	
Weight Increased	Vitamins	C
	L-Carnitine	C
	Magnesium	C
	Glucosamine	
	Chondroitin	C
	Msm	C
	Celecoxib	C
	Tamsulosin	
	Hydrochloride	C
	Tolterodine Tartrate	C
	Amlodipine Besylate,	
	Benazepril	
	Hydrochloride)	C
	Metoprolol Succinate	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/17/02ISR Number: 3935412-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200698

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pancreatic Carcinoma	Health Professional	Neurontin (Gabapentin)	PS		

Date:06/17/02ISR Number: 3935492-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200696

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Blood Thyroid Stimulating Hormone Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other		Malaise					
ORAL		Thyroxine Decreased		Levothyroxine	SS		ORAL
PER ORAL				Clonidine Unspecified Medication	C		

Date:06/17/02ISR Number: 3935552-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200692

Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Drug Effect Decreased Fibroma	Consumer	Neurontin (Gabapentin)	PS		ORAL
3600 MG (1800		Medication Error					
MG, BID), PER		Nausea					
ORAL							

Date:06/17/02ISR Number: 3935615-8Report Type:Expedited (15-DaCompany Report #351-0945-M0200002

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

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3600 MG (DAILY), PER ORAL			Anorexia Asthenia Hepatic Failure Jaundice	Foreign Health Professional Other	Neurontin (Gabapentin)	PS		ORAL
					Omeprazole Clonazepam Ciclosporine Azatioprine	C C C C		

Date:06/17/02ISR Number: 3936182-5Report Type:Expedited (15-DaCompany Report #055-0945-M0200050
Age:60 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 900 MG (TID), Initial or Prolonged OTHER			Lung Infection	Foreign Consumer	Gabapentin (Gabapentin)	PS		OTHER

Date:06/17/02ISR Number: 3936192-8Report Type:Expedited (15-DaCompany Report #033-0945-M0200045
Age:38 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1500 MG TID PER ORAL			Ichthyosis Acquired Liver Disorder Pancreatitis Rash Maculo-Papular	Foreign Health Professional	Neurontin (Gabapentin) (Lamivudine,	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

600 MG BID			Zidovudine)	SS		ORAL
PER ORAL						
800 MG BID			(Lopinavir, Ritonavir)	SS		ORAL
PER ORAL						
			(Clobazam)	C		
			(Calcium Folate)	C		
			(Pyrimethamine)	C		
			(Sulfadiazine)	C		

Date:06/17/02ISR Number: 3936194-1Report Type:Expedited (15-DaCompany Report #033-0945-M0200088
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hyperaesthesia Major Depression	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
PER ORAL							
Other		Oedema Peripheral	Professional	(Zopiclone)	SS		ORAL
PER ORAL							
SUBCUTANEOUS	SUBCUTANEOUS	Peripheral Coldness		(Insulin)	SS		
		Peripheral Sensory		(Imatinib)	SS		ORAL
400 MG (100 MG QID) PER		Neuropathy					
ORAL							
PER ORAL				(Ramipril)	SS		ORAL
PER ORAL				(Paroxetine)	SS		ORAL

Date:06/18/02ISR Number: 3935054-XReport Type:Expedited (15-DaCompany Report #031-0945-M0200006
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Gastrointestinal	Foreign	Neurontin			

Initial or Prolonged 4200 MG (DAILY), PER ORAL	Haemorrhage Loss Of Consciousness Shock	Health Professional	(Gabapentin)	PS	ORAL
			Diazepam Midazolam Fentanyl	C C C	

Date:06/18/02ISR Number: 3935884-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200460
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other (DAILY)		Medication Error Overdose	Health Professional	Neurontin (Gabapentin)	PS		
		Renal Cell Carcinoma Stage Unspecified	Company Representative	(Oxycodone Hydrochloride) (Zolpidem Tartrate) (Rabeprazole Sodium) (Atorvastatin) (Citalopram Hydrobromide) (Fentanyl) (Pseudoephedrine Sulfate, Loratadine)	C C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/18/02ISR Number: 3936068-6Report Type:Expedited (15-DaCompany Report #001-0981-M0202719

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetes Mellitus	Consumer	Atorvastatin			
10 MG		Diabetic Complication		(Atorvastatin)	PS		
600 MG				Gabapentin	SS		
				Glipizide	SS		
				Insulin Human,			
				Insulin Human			
				Injection, Isophane	C		
				(Enalapril Maleate)	C		
				Furosemide	C		
				Metformin			
				Hydrochloride	C		
				Ranitidine	C		
				Rosiglitazone	C		
				Diltiazem	C		

Date:06/18/02ISR Number: 3936089-3Report Type:Expedited (15-DaCompany Report #A211461

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Diabetes Mellitus	Consumer	Glucotrol Tablets	PS		
Intervention to		Diabetic Complication		Lipitor	SS		
10.00 MG							
Prevent Permanent							
TOTAL							
Impairment/Damage				Neurontin	SS		
600.00 MG							
TOTAL							
				Novolin 70/30	C		
				Vasotec	C		
				Lasix	C		
				Glucophage Xr	C		
				Ranitidine	C		
				Avandia	C		
				Diltiazem	C		

Date:06/20/02ISR Number: 3937263-2Report Type:Expedited (15-DaCompany Report #PERI00202001538
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Foreign	Coversyl			
		Shock	Health	(Perindopril)	PS		ORAL
2 MG QD PO							
		Syncope	Professional	Gabapentin			
			Other	(Gabapentin)	SS		ORAL
900 MG DAILY							
PO, 300 MG							
DAILY PO							
				Insulin (Insulin)	C		

Date:06/21/02ISR Number: 3937681-2Report Type:Expedited (15-DaCompany Report #C2002-0958.01
Age:64 YR Gender:Male I/FU:F

Outcome	PT
Death	Activated Partial
Hospitalization -	Thromboplastin Time
Initial or Prolonged	Prolonged
	Aortic Aneurysm
	Arterial Occlusive
	Disease
	Blood Bilirubin Increased

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Freedom Of Information (FOI) Report

Dose	Duration	Cardiac Arrest Coagulopathy Diarrhoea	Report Source	Product	Role	Manufacturer	Route
		Fistula	Consumer	Buspirone			
		Fungal Infection	Health	Hydrochloride			
		Haematemesis	Professional	Tablets 15 Mg	PS		
		Haematochezia		Diazepam Tablets 5			
		Haematocrit Decreased		Mg	SS		
		Haemodialysis		Loperamide Hcl			
		Hepatic Cirrhosis		Capsule 2 Mg	SS		
		Hypersplenism		Propoxyphene			
		Hypoglycaemia		Napsulate And Apap			
		Hypotension		Tablets 100mg/650 Mg	SS		
		Injury		Neurontin Capsules			
		International Normalised		100 Mg Parke Davis	SS		
		Ratio Increased		Ambien Tablets			
		Liver Function Test		Searle	SS		
		Abnormal		Oxycodone/Apap			
		Multi-Organ Failure		Tablets	SS		
		Oedema		Oxycontin Tablets	SS		
		Thrombocytopenia		Morphine Tablets	SS		
		Vasodilatation		Acetaminophen	C		
				Penicillin	C		
				Amlodipine	C		
				Nephrocaps (Qd)	C		
				Lidocaine/Prilocaine			
				Cream	C		
				Folic Acid	C		
				Omeprazole	C		
				Cetirizine	C		
				Sevelamer Capsules	C		
				Propranolol	C		
				Sucralfate	C		

Date:06/21/02ISR Number: 3937860-4Report Type:Expedited (15-DaCompany Report #001-0073-M0200260

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Adrenal Insufficiency	Consumer	Dilantin (Phenytoin			
Other		Brain Neoplasm		Sodium)	PS		ORAL
PER ORAL		Disease Recurrence		Gabapentin	SS		ORAL
900 MG (300							

MG, TID), PER

Muscle Twitching

ORAL

Oedema

Rash

Vomiting

Methylphenidate

Hydrochloride SS

Dexamethasone SS

Carbamazepine SS

Esomeprazole C

Sulfamethoxazole,

Trimethoprim C

Celecoxib C

Date:06/21/02ISR Number: 3937935-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200718

Age: Gender:Male I/FU:I

Outcome

PT

Other

Back Pain

Fluid Retention

Hypoaesthesia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pharmaceutical Product	Report Source	Product	Role	Manufacturer	Route
600 MG (BID), PER ORAL		Nausea Oedema Peripheral Complaint	Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG (DAILY), PER ORAL				Celecoxib	SS		ORAL

Date:06/21/02ISR Number: 3937971-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200710
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (DAILY)		Unevaluable Event	Consumer	Neurontin (Gabapentin)	PS		
900 MG				Lithium	SS		

Date:06/21/02ISR Number: 3937974-9Report Type:Expedited (15-DaCompany Report #001-0981-M0201657
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other Required Intervention to 1200 MG Prevent Permanent (QID), PER Impairment/Damage ORAL		Aortic Aneurysm Coronary Artery Occlusion Diarrhoea Memory Impairment Muscular Weakness Pain	Consumer	Atorvastatin (Atorvastatin) (Gabapentin) (Sertraline)	PS SS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
(DAILY), PER		Vascular Access		Hydrochloride)	SS		ORAL
ORAL		Complication		(Atenolol)	C		
				(Terazosin			
				Hydrochloride)	C		
				(Rofecoxib)	C		
Date:06/21/02ISR Number: 3938006-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200447							
Age: Gender:Male I/FU:F							
Death		Aphagia	Health	Neurontin			
Other		Condition Aggravated	Professional	(Gabapentin)	PS		ORAL
300 MG (TID),		Dizziness					
PER ORAL		Neuralgia		Oxycodone			
		Oral Intake Reduced		Hydrochloride	SS		ORAL
PER ORAL				Oxycodone	SS		ORAL
PER ORAL				Morphine Sulfate	SS		ORAL
PER ORAL				Fentanyl	SS		
TOPICAL	TOPICAL			Furosemide	C		
				Potassium	C		
				Paracetamol,			
				Dextropropoxyphene	C		
				Compounded Ativan,			
				Bendaryl,			
				Haloperidol			
				Suppository	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/21/02ISR Number: 3938133-6Report Type:Expedited (15-DaCompany Report #041-0945-M0200015

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bile Duct Obstruction	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
PER ORAL		Liver Function Test					
		Abnormal	Professional Company Representative				

Date:06/21/02ISR Number: 3938134-8Report Type:Expedited (15-DaCompany Report #033-0945-M0200091

Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Respiratory Failure	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
300 MG (100							
MG, TID), PER			Professional				
ORAL							

Baclofen	C
Cascara Dry Extract	C
Amitriptyline Hydrochloride	C

Date:06/21/02ISR Number: 3938266-4Report Type:Expedited (15-DaCompany Report #044-0945-M0200071

Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Convulsion Medication Error	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
2400 MG (1200							
Other		Tic	Professional				
MG, DAILY)							

PER ORAL

Tramadol	SS
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INTRAVENOUS INTRAVENOUS

Clonidine C
Piroxicam Betadex C

Date:06/21/02ISR Number: 3939274-XReport Type:Expedited (15-DaCompany Report #049-0945-M0200059
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Cardiac Failure	Foreign	Neurontin			
Initial or Prolonged	Dyspnoea Exertional	Health	(Gabapentin)	PS		ORAL
(DAILY), PER						
Other	Hepatic Steatosis	Professional				
ORAL						
	Hepatitis		(Levodopa			
	Hepatomegaly		Bensarazide			
	Oedema Peripheral		Hydrochloride)	C		
	Weight Increased					

Date:06/24/02ISR Number: 3938400-6Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
Age:46 YR Gender:Female I/FU:F

Outcome	PT
Disability	Brain Damage
Other	Depression
	Drug Hypersensitivity
	Drug Toxicity
	Dyskinesia
	Intentional Misuse
	Medication Error

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Speech Disorder Tongue Oedema	Report Source	Product	Role	Manufacturer	Route
1800 MG	(DAILY), PER		Consumer Health	Neurontin (Gabapentin)	PS		ORAL
			Professional	(Lithium Carbonate)	C		
				(Clonazepam)	C		
				(Methylphenidate Hydrochloride)	C		
				(Levothyroxine Sodium)	C		
				(Liothyronine Sodium)	C		
				(Sertraline Hydrochloride)	C		
				(Pilocarpine Hydrochloride)	C		
				(Metoprolol Succinate)	C		
				(Lansoprazole)	C		
				(Hyoscyamine Sulfate)	C		
				(Diltiazem Hydrochloride)	C		

Date:06/24/02ISR Number: 3938788-6Report Type:Expedited (15-DaCompany Report #001-0945-M0200726

Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Accident Agitation	Consumer	Neurontin (Gabapentin)	PS		ORAL
PER ORAL		Anxiety Condition Aggravated		Sertraline Hydrochloride	SS		ORAL
50 MG	(DAILY), PER	Confusional State					

ORAL	Depression			
	Diarrhoea	Ziprasidone		
20 MG	Eating Disorder Symptom	Hydrochloride	SS	ORAL
(DAILY), PER	Fear			
ORAL	Feeling Hot And Cold			
25 MG	Headache	Topiramate	SS	ORAL
(DAILY), PER	Increased Appetite			
ORAL	Insomnia			
	Memory Impairment	Verapamil	C	
	Nervousness	Furosemide	C	
	Obsessive-Compulsive Disorder	Estropipate	C	
	Panic Disorder	Estradiol	C	
	Paraesthesia	Estradiol	C	
	Pruritus	Clonazepam	C	
	Psoriasis			
	Tremor			
	Weight Increased			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/24/02ISR Number: 3939280-5Report Type:Expedited (15-DaCompany Report #034-0945-M0200008

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatic Cirrhosis	Foreign	Gabapentin			
		Hepatic Steatosis	Health	(Gabapentin)	PS		ORAL
PER ORAL			Professional	Valproic Acid	SS		ORAL
PER ORAL							

Date:06/24/02ISR Number: 3939340-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200730

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia	Consumer	Neurontin			
		Chromaturia	Health	(Gabapentin)	PS		ORAL
3200 MG (FOUR		Oedema Peripheral	Professional				
TIMES DAILY),		Vision Blurred					
PER ORAL		Visual Disturbance		(Celecoxib)	SS		ORAL
400 MG (TWICE							
DAILY), PER				(Zolpidem Tartrate)	C		
ORAL							

Date:06/25/02ISR Number: 3939509-3Report Type:Expedited (15-DaCompany Report #2002CG00895

Age:91 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hepatitis Cholestatic	Foreign	Tenormine	PS		ORAL
25 MG DAILY			Health				
Initial or Prolonged		Pleural Effusion	Professional	Neurontin	SS		ORAL
PO							
600 MG DAILY							

Other

PO

Coumadin SS

Aldactone SS

25MG PO

Hemigoxine Nativelle SS ORAL

0.125 MG

DAILY PO

Lasilix SS ORAL

203G DAILY PO

Date:06/25/02ISR Number: 3939526-3Report Type:Expedited (15-DaCompany Report #002-0945-M0200070
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Gout	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
2400 MG (800 MG, TID),							
ORAL				Venlafaxine	SS		ORAL
112.5 MG							
(DAILY), ORAL							

Date:06/26/02ISR Number: 3940254-9Report Type:Expedited (15-DaCompany Report #001-0073-M0200250
 Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Anxiety Cerebrovascular Accident Convulsion Crying Feeling Abnormal Headache Insomnia Irritable Bowel Syndrome

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Myocardial Infarction Nausea Nervousness	Report Source	Product	Role	Manufacturer	Route
300 MG (THREE TIMES DAILY), ORAL		Osteoporosis Visual Disturbance	Consumer Health Professional	Dilantin Infatabs (Phenytoin Sodium)	PS		ORAL
400 MG (DAILY) ORAL ORAL				Gabapentin	SS		ORAL
1200 MG (DAILY), ORAL ORAL				Lorazepam (Vitamin C With D)	SS SS		ORAL ORAL
ORAL				Alendronate Sodium	SS		ORAL
				Lamotrigine	SS		ORAL
				Levetiracetam Lorazepam	SS C		

Date:06/26/02ISR Number: 3940433-0Report Type:Expedited (15-DaCompany Report #034-0945-M0200006
Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), Other ORAL		Blood Urea Increased Cardiac Failure Gastrointestinal	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
		Haemorrhage Hepatic Congestion Hepatomegaly International Normalised Ratio Increased Oedema Peripheral Pulmonary Hypertension		Anticoagulant Therapy Digoxin Furosemide	C C C		

Date:06/27/02ISR Number: 3943286-XReport Type:Periodic
Age:63 YR Gender:Female I/FU:I

Company Report #A203061

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 25.00 MG		Chest Pain	Consumer	Zoloft Tablets	PS		ORAL
Intervention to TOTAL:DAILY:0		Dysphagia					
Prevent Permanent RAL		Dyspnoea					
Impairment/Damage 2400.00 MG		Intentional Misuse		Gabapentin	SS		ORAL
TOTAL:QID:ORA		Medication Error					
L		Weight Increased					
				Vicodin	C		
				Synthroid	C		

Date:06/28/02ISR Number: 3941518-5Report Type:Direct
Age:79 YR Gender:Female I/FU:I

Company Report #CTU 171238

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia Blood Pressure Increased Coordination Abnormal		Neurontin 300 Mg=12 Tablets Daily=3600 Mg Daily	PS		ORAL
ORALLY =1200 MG ONCE X A		Syncope					
DAY	8	MON					
		Vertigo					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/28/02ISR Number: 3942035-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200517
Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Balance Disorder Blood Glucose Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
	Diplopia		Fexofenadine			
	Dysarthria		Hydrochloride	C		
	Feeling Abnormal		Levothyroxine Sodium	C		
	Hemiparesis		Acifex	C		
	Hypertension		Rofecoxib	C		
	Mental Impairment		Atenolol	C		
	Pain		Simvastatin	C		
	Paraesthesia		Amitriptyline			
	Somnolence		Hydrochloride	C		
			Alprazolam	C		
			Diazepam	C		

Date:06/28/02ISR Number: 3942164-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200747
Age:50 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG (TID), ORAL	Alopecia	Consumer	Neurontin (Gabapentin)	PS		ORAL
			(Steroids)	C		

Date:07/01/02ISR Number: 3942509-0Report Type:Expedited (15-DaCompany Report #200212411FR
Age:91 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 20 MG QD PO	Hepatitis Cholestatic Pleural Effusion	Foreign Health	Furosemide (Lasix Faible) Tablets	PS		ORAL
		Professional Other	Spirolactone (Aldactone) Coated			

25 MG QD PO		Tablets	SS	ORAL
		Digoxin (Hemigoxine Nativelle) Tablets	SS	ORAL
0.125 MG QD				
PO		Warfarin Sodium (Coumadine) Tablets	SS	ORAL
PO	722 DAY			
		Atenolol (Tenormine) Coated Tablets	SS	ORAL
25 MG QD PO	1079 DAY			
		Gabapentin (Neurontin) Coated Tablets	SS	ORAL
600 MG QD PO	17 DAY			

Date:07/01/02ISR Number: 3942623-XReport Type:Expedited (15-DaCompany Report #A208993
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Coma	Health	Zoloft Tablets	PS		ORAL
200.00 MG							
Hospitalization - TOTAL:DAILY:0		Drug Interaction	Professional				
Initial or Prolonged RAL		Pain					
Required ORAL		Pneumonia Streptococcal		Gabapentin	SS		ORAL
Intervention to Prevent Permanent Impairment/Damage				Elavil	C		
				Premarin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/01/02ISR Number: 3942627-7Report Type:Expedited (15-DaCompany Report #A208992

Age:51 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 100.00 MG	Coma	Health	Zoloft Tablets	PS		ORAL
Hospitalization - TOTAL:DAILY:0	Drug Interaction	Professional				
Initial or Prolonged RAL	Feeling Drunk					
Required 3600.00 MG	Gait Disturbance		Gabapentin	SS		ORAL
Intervention to TOTAL:TID:ORA	Hallucination					
Prevent Permanent L	Influenza					
Impairment/Damage	Lung Disorder		Anafranil	SS		
	Pneumonia		Slo-Bid Gyrocaps	C		
	Staphylococcal Infection		Q-Tuss	C		
			Klor-Con	C		
			Albuterol	C		
			Premarin	C		
			Lotensin	C		
			Baycol	C		
			Xanax	C		
			Furosemide	C		
			Prednisone	C		
			Trazodone	C		
			Prilosec	C		
			Metoclopramine	C		
			Atrovent	C		
			Intal	C		
			Pulmicort	C		
			Miacalcin	C		
			Nasonex	C		
			Gaviscon Type			
			Unknown	C		
			Darvocet	C		
			Lonox	C		
			Prochlorperazine	C		
			Carafate	C		
			Hydromet	C		
			Calcium With Vitamin			
			D	C		
			Serevent	C		
			Singulair	C		

Vitamin E	C
Ocuvite Extra	C
B-12	C
Osteo-Bi-Flex	C
Prinivil	C
Clonazepam	C
Protonix	C
Pepcid	C
Zyrtec	C
Mirapex	C

Date:07/01/02ISR Number: 3943680-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200447

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dizziness	Health	Neurontin			
Other		Dysphagia	Professional	(Gabapentin)	PS		ORAL
300 MG (TID),		Oral Intake Reduced					
ORAL		Pain		Oxycodone			
				Hydrochloride	SS		ORAL
ORAL				Oxycodone	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL				Morphine Sulfate	SS		ORAL
TOPICAL	TOPICAL			Fentanyl	SS		
				Furosemide	C		
				Potassium	C		
				Di-Gesic	C		
				Compounded Ativan, Benadryl, Haloperidol Suppository	C		

Date:07/01/02ISR Number: 3943682-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200754
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Arthralgia Disease Recurrence					
ORAL		Disturbance In Attention Dizziness Medication Error Memory Impairment Somnolence		Di-Gesic	C		

Date:07/01/02ISR Number: 3943684-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200391
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		
Other		Colonoscopy Medication Residue	Health Professional				

Date:07/01/02ISR Number: 3943733-3Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Brain Neoplasm	Health	Neurontin			

Initial or Prolonged	Cognitive Disorder	Professional	(Gabapentin)	PS	ORAL
ORAL					
Other	Convulsion	Company			
	Ependymoma	Representative			
	Fatigue				
	Memory Impairment				
	Somnolence				
	Speech Disorder				

Date:07/01/02ISR Number: 3943846-6Report Type:Expedited (15-DaCompany Report #032-0945-M0200002
Age:59 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Bronchospasm	Foreign	Neurontin			
Hospitalization -	Respiratory Depression	Health	(Gabapentin)	PS		
300 MG DAILY						
Initial or Prolonged	Respiratory Failure	Professional	Clonazepam	C		
	Shock		Amitriptyline			
			Hydrochloride	C		
			Acetylsalicylate			
			Lysine	C		
			Trimebutine Maleate	C		
			Baclofen	C		
			Dantrolene Sodium	C		

Freedom Of Information (FOI) Report

Vesirip C

Date:07/02/02ISR Number: 3942361-3Report Type:Expedited (15-DaCompany Report #309551
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Apgar Score Low Convulsion Neonatal Feeding Problem In Newborn Foetal Disorder Hypoglycaemia Neonatal Maternal Drugs Affecting Foetus Neonatal Apnoeic Attack Nuclear Magnetic Resonance Imaging Abnormal Pregnancy Small For Dates Baby Tonic Clonic Movements	Consumer	Rivotril Neurontin Depakine	PS SS SS	Roche	

Date:07/02/02ISR Number: 3943348-7Report Type:Expedited (15-DaCompany Report #1005340
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Bilirubin Increased	Foreign Health Professional Other	Keppra Gabapentin Valproate Carbamazepine	PS SS C C		

Date:07/03/02ISR Number: 3944158-7Report Type:Expedited (15-DaCompany Report #2002050012
Age:83 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200 MG (AFTER		Coma Drug Level Decreased	Literature Health	Gabapentin (Gabapentin)	PS		

EACH

Dyskinesia

Professional

Mental Impairment

Amiodarone	C
Calcium Carbonate	C
Docusate Sodium	C
Erythropoietin	C
Folic Acid	C
Metoprolol	C
Tamsulosin	C
Lactulose	C
Promethazine	C
Amitriptyline	C
Carbamazepine	C

Date:07/03/02ISR Number: 3944440-3Report Type:Expedited (15-DaCompany Report #2002050066
Age: Gender:Unknown I/FU:I

Outcome	PT
Congenital Anomaly	Anal Fissure
	Haemangioma
	Maternal Drugs Affecting

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Foetus

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:07/03/02ISR Number: 3944442-7Report Type:Expedited (15-DaCompany Report #2002050243
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Thrombosis	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:07/03/02ISR Number: 3944878-4Report Type:Expedited (15-DaCompany Report #2002050043
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Adrenal Adenoma	Consumer	Neurontin (Gabapentin)	PS		ORAL
(TWICE		Blood Arsenic Increased					
DAILY), ORAL		Blood Pressure Increased					
ORAL		Dyspnoea		Lamotrigine	SS		ORAL
		Feeling Abnormal		Clonazepam	C		
		Helicobacter Infection					
		Hunger					
		Laboratory Test Abnormal					
		Nausea					
		Weight Increased					

Date:07/03/02ISR Number: 3944880-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200698
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pancreatic Carcinoma	Health Professional	Neurontin (Gabapentin)	PS		

Date:07/03/02ISR Number: 3944898-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200641
Age: Gender:Male I/FU:F

Outcome	PT
Other	Asthenia
	Balance Disorder
	Burning Sensation
	Coordination Abnormal
	Depression
	Dizziness
	Drug Ineffective
	Eye Irritation
	Eye Movement Disorder
	Fatigue
	Frequent Bowel Movements
	Haematochezia
	Haemorrhoids
	Hallucination

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG		Incoherent Micturition Urgency Muscle Atrophy	Consumer	Neurontin (Gabapentin)	PS		ORAL
(BID), ORAL		Muscle Spasms Thinking Abnormal Vision Blurred		Zoloft (Sertraline)	SS		
Date:07/05/02ISR Number: 3944485-3Report Type:Expedited (15-DaCompany Report #001-0073-M0200194 Age: Gender:Male I/FU:F							
Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (100 Other MG BID), ORAL		Akathisia Cognitive Disorder Deep Vein Thrombosis Delusion	Health Professional	Dilantin Suspension (Phenytoin Sodium)	PS		ORAL
1200 MG (TID)		Depression Drug Ineffective		Gabapentin (Gabapentin)	SS		
35 MG (TID & HS), ORAL		Drug Interaction Drug Level Decreased Screaming		Fluphenazine Hydrochloride	SS		ORAL
8 MG (4 MG, BID), ORAL		Tardive Dyskinesia Urine Calcium Increased		Fluoxetine Hydrochloride Risperidone	SS SS		ORAL
17.75 MG				Warfarin Sodium Fluphenazine Decanoate	SS C		ORAL
(EVERY 7 DAYS), ORAL							

Sertraline
Hydrochloride C

Date:07/05/02ISR Number: 3944933-9Report Type:Expedited (15-DaCompany Report #2002050428
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arterial Occlusive Disease	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL							

Phenytoin Sodium C
Mitrazapine C
Pentoxifylline C
Acetylsalicylic Acid C
Tocopherol C
Potassium C

Date:07/05/02ISR Number: 3944934-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200633
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated Neoplasm Malignant	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other							
900 MG (300 MG, THREE TIMES DAILY) ORAL							

Celecoxib SS

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/05/02ISR Number: 3946323-1Report Type:Expedited (15-DaCompany Report #309551

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Apgar Score Low	Foreign	Rivotril			
Initial or Prolonged	Clonic Convulsion	Other	(Clonazepam)	PS		
0.5 MG DAILY						
	Convulsion Neonatal		Neurontin			
	Feeding Problem In		(Gabapentin)	SS		
3 DOSE FORM						
DAILY	Newborn					
	Foetal Distress Syndrome					
	Grand Mal Convulsion					
	Hypoglycaemia Neonatal					
	Maternal Drugs Affecting					
	Foetus					
	Neonatal Apnoeic Attack					
	Pregnancy					

Date:07/05/02ISR Number: 3946443-1Report Type:Expedited (15-DaCompany Report #033-0945-M0200007

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Hypokalaemia	Foreign	Neurontin			
Initial or Prolonged	Megacolon	Health	(Gabapentin)	PS		ORAL
2400 MG (800						
MG, TID),		Professional				
ORAL						
			Acetylsalicylate			
			Lysine	C		

Date:07/05/02ISR Number: 3946590-4Report Type:Expedited (15-DaCompany Report #2002050195

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Pruritus	Foreign	Neurontin			

800 MG (400
 MG, BID),
 ORAL

Health (Gabapentin) PS ORAL
 Professional

Oxacarbazepine C

Date:07/05/02ISR Number: 3946662-4Report Type:Expedited (15-DaCompany Report #B0263843A
 Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged TWICE PER DAY		Blood Alkaline Phosphatase Increased	Foreign	Combivir Tablet (Combivir)	PS		ORAL
ORAL		Hepatitis Cholestatic					
ORAL		Pancreatitis Rash Maculo-Papular		Kaletra Capsule (Kaletra)	SS		ORAL
				Gabapentin (Formulation Unknown) (Gabapentin)	SS		ORAL
500 MG THREE TIMES PER DAY							
ORAL				Pyrimethamine Sulphadiazine Calcium Folate	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/08/02ISR Number: 3944404-XReport Type:Expedited (15-DaCompany Report #A0373437A

Age:41 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - INTRAVENOUS 1.5MG Initial or Prolonged times per day	Blood Glucose Increased Nine Diabetic Ketoacidosis		Flolan	PS	Glaxo Wellcome	
	Pain In Extremity Pancreatitis		Neurontin	SS		ORAL

Date:07/09/02ISR Number: 3947547-XReport Type:Expedited (15-DaCompany Report #PHRM2002FR01755

Age:48 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 100 MG, QID, ORAL	Peripheral Sensory Neuropathy	Foreign Health Professional	Glivec (Imatinib) Capsule	PS		ORAL
7.5 MG, QD, ORAL		Other	Imovane (Zopiclone) Tablet	SS		ORAL
SUBCUTANEOUS 28 IU/D, SUBCUTANEOUS			Umuline (Insulin Human Zinc Suspension)	SS		
600 MG DAILY, ORAL : 1.2 G DAILY			Neurontin (Gabapentin)	SS		ORAL
5 MG, QD, ORAL			Triatec Capsule (Ramipril)	SS		ORAL

Deroxat (Paroxetine
Hydrochloride)
Tablet

SS

ORAL

20 MG, QD,

ORAL

Date:07/10/02ISR Number: 3947806-0Report Type:Expedited (15-DaCompany Report #044-0945-M0200083
Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neuroleptic Malignant Syndrome	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG							
(DAILY), ORAL				Lithium	C		

Date:07/10/02ISR Number: 3947995-8Report Type:Expedited (15-DaCompany Report #2002050494
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Hormone Level Abnormal Nerve Compression	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (TID), Other		Sleep Disorder					
ORAL		Tremor Weight Decreased		Sinemet	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/10/02ISR Number: 3948005-9Report Type:Expedited (15-DaCompany Report #2002050553

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Health	Neurontin			
		Skin Necrosis	Professional	(Gabapentin)	PS		

Date:07/10/02ISR Number: 3948015-1Report Type:Expedited (15-DaCompany Report #A210098

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Accident	Health	Glucotrol Tablets	PS		ORAL
ORAL							
Initial or Prolonged		Areflexia	Professional	Celebrex	SS		ORAL
200.00 MG							
Required		Back Injury					
TOTAL:DAILY:O							
Intervention to		Blood Cholesterol					
RAL							
Prevent Permanent		Increased		Neurontin	SS		
600.00 MG							
Impairment/Damage		Blood Glucose Decreased					
TOTAL:PID							
		Coronary Artery Occlusion		Oral Hypoglycemic			
		Diabetic Neuropathy		Agent	SS		
		Dizziness		Glucosamine	C		
		Fall		Chondroitin	C		
		Gait Disturbance		Avandia	C		
		Intervertebral Disc		Amitriptyline	C		
		Degeneration		Zebeta	C		
		Limb Injury		Atorvastatin	C		
		Myalgia					
		Osteoarthritis					
		Somnolence					
		Vertigo					

Date:07/11/02ISR Number: 3946725-3Report Type:Expedited (15-DaCompany Report #316676

Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -	Renal Failure	Rivotril	PS	Roche
17 DAY				
Initial or Prolonged		Bactrim	SS	Roche
1496 DAY				
		Neurontin	SS	
217 DAY		Viread	SS	
		Norvir	SS	
217 DAY				

Date:07/11/02ISR Number: 3948181-8Report Type:Expedited (15-DaCompany Report #2002050619
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abdominal Pain	Foreign	Neurontin			
Initial or Prolonged		Liver Function Test	Health	(Gabapentin)	PS		ORAL
		Abnormal	Professional				
		Pancreatic Enzymes					
		Increased					

Date:07/11/02ISR Number: 3948680-9Report Type:Expedited (15-DaCompany Report #A02200200709
Age:23 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Clonic Convulsion
Initial or Prolonged	Drug Level Below
	Therapeutic
	Epilepsy

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
5 MG/QD / 10 MG QD / 30 MG QD, ORAL	3 DAY	Fall Hallucination Intentional Misuse	Foreign Health Professional	Stilnox - (Zolpidem) - Tablet - 10 Mg / Tablet - 10 Mg / Tablet - 10 Mg	PS		ORAL
2 UNIT QD, ORAL	7 WK	Loss Of Consciousness Malaise		Neurontin - (Gabapentin)	SS		ORAL
ORAL	1 DAY			Gardenal - (Phenobarbital) - Tablet - 100 Mg	SS		ORAL

Date:07/11/02ISR Number: 3949064-XReport Type:Expedited (15-DaCompany Report #2002050568
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Encephalopathy	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:07/11/02ISR Number: 3949065-1Report Type:Expedited (15-DaCompany Report #A214294
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - DAILY:ORAL		Aneurysm	Consumer	Cardura	PS		ORAL
Initial or Prolonged 900.00 MG Disability TOTAL:		Blindness Cardiac Disorder		Neurontin	SS		ORAL

Required
 TID:ORAL
 Intervention to
 Prevent Permanent
 Impairment/Damage

Cardiac Failure
 Congestive
 Cerebral Haemorrhage
 Corneal Transplant
 Coronary Artery Occlusion
 Drug Effect Decreased
 Eye Disorder
 Intracranial Aneurysm
 Memory Impairment
 Muscle Disorder
 Neoplasm
 Renal Failure

Insulin C

Date:07/12/02ISR Number: 3948335-0Report Type:Expedited (15-DaCompany Report #2002050725
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other 600 MG (DAILY)		Drug Interaction Overdose Shock	Foreign Literature Health Professional	Neurontin (Gabapentin) Risperidone Zuclopendixol Decanoate Tropatepine Hydrochloride Lorazepam Fluphenazine Hydrochloride	PS SS C C C C		
6 MG (DAILY)							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Midazolam C
 Propofol C
 Remifentanyl C
 Atracurium C

Date:07/12/02ISR Number: 3948501-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
 Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Brain Damage	Consumer	Neurontin			
Other		Depression	Health	(Gabapentin)	PS		ORAL
1800 MG DAILY		Dysarthria	Professional				
ORAL		Gingival Disorder		Lamotrigine	C		
		Hypersensitivity		Lithium Carbonate	C		
		Intentional Misuse		Clonazepam	C		
		Medication Error		Methylphenidate			
		Nervous System Disorder		Hydrochloride	C		
		Oral Pain		Levothyroxine Sodium	C		
		Temperature Intolerance		Liothyronine Sodium	C		
		Tongue Oedema		Sertraline			
		Tongue Paralysis		Hydrochloride	C		
		Tooth Disorder		Pilocarpine			
				Hydrochloride	C		
				Metoprolol Succinate	C		
				Lansoprazole	C		
				Hyoscyamine Sulfate	C		
				Diltiazem			
				Hydrochloride	C		
				Lamotrigine	C		

Date:07/12/02ISR Number: 3948923-1Report Type:Expedited (15-DaCompany Report #2002050617
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion	Consumer	Neurontin			
Initial or Prolonged		Enamel Anomaly		(Gabapentin)	PS		
Other		Mental Disorder					
		Tooth Disorder					
		Tooth Loss					

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Akathisia
Other	Cognitive Disorder
	Convulsion
	Deep Vein Thrombosis
	Delusion
	Depression
	Drug Effect Decreased
	Drug Interaction
	Drug Level Decreased
	Fear
	Oral Intake Reduced
	Protein Total Increased
	Psychotic Disorder

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Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
200 MG (100 MG, BID),		Screaming Tardive Dyskinesia Thinking Abnormal	Health Professional	Dilantin Suspension (Phenytoin Sodium)	PS		ORAL
ORAL		Urine Calcium Increased Weight Increased		Gabapentin	SS		
1200 MG (TID)				Fluphenazine Decanoate	SS		ORAL
17.75 MG (EVERY 7 DAYS), ORAL				Fluphenazine Hydrochloride	SS		ORAL
35 MG (TID & HS), ORAL				Warfarin Sodium	SS		
				Fluoxetine Hydrochloride	SS		
8 MG (4 MG, BID), ORAL				Risperidone	SS		ORAL
				Sertraline Hydrochloride	C		

Date:07/12/02ISR Number: 3949477-6Report Type:Expedited (15-DaCompany Report #001-0719-M0100408

Age: Gender:Male I/FU:F

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
1200 MG (BID), ORAL		Hospitalization -	Consumer	Lopid (Gemfibrozil)	PS		ORAL
Other		Arthralgia	Health Professional	Gabapentin			
		Asthenia					

ORAL	Back Pain	(Gabapentin)	SS	ORAL
5 MG (DAILY),	Chest Wall Pain	Amlodipine		
ORAL	Confusional State	(Amlodipine)	SS	ORAL
	Depression			
5 MG (DAILY),	Difficulty In Walking	Glipizide		
ORAL	Disorientation	(Glipizide)	SS	ORAL
	Dizziness			
SUBLINGUAL	Headache	Nifedipine		
SUBLINGUAL	Hepatic Steatosis	(Nifedipine)	SS	
10 MG,	Hypoaesthesia			
	Joint Swelling	(Unspecified		
	Memory Impairment	Cholesterol Lowering		
	Myalgia	Drug)	SS	
	Osteoarthritis	Furosemide	C	
	Otitis Externa	Acetylsalicylic Acid	C	
	Polyuria	Multivitamins	C	
	Skin Papilloma	Terazosin	C	
	Varicose Vein	Labetalol		
	Vein Disorder	Hydrochloride	C	
	Viral Upper Respiratory	Sunastatin	C	
	Tract Infection	Clopidogrel	C	
	Vision Blurred	Meloxicam	C	
		Pericad Plus		
		Vitamins	C	
		(L-Carnitine)	C	
		Magnesium	C	
		(Glucosamine		
		Chondroitin)	C	
		(Msm)	C	
		Celecoxib	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tamsulosin	
Hydrochloride	C
Tolterodine	
L-Tartrate	C
Lotrel	C
Metoprolol Succinate	C
Simvastatin	C
Vicodin	C
Crataegus Extract	C
Ubidecarenone	C
Calcium	C
Garlic	C
(Unspecified Pain Medication)	C

Date:07/15/02ISR Number: 3948090-4Report Type:Expedited (15-DaCompany Report #316974
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Klonopin	PS	Roche	
Other		Dyspnoea	Professional	Celebrex	SS		
		Urticaria		Neurontin	SS		
				Topamax	SS		
				Zantac	SS		
				Progesterone	C		
				Salbutamol	C		
				Pentosane			
				Polysulphate	C		
				Hydroxyzine	C		
				Amitriptyline	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
Dose			Health	Zantac	PS	Glaxo Wellcome	
Other		Dyspnoea	Professional	Wellbutrin	SS	Glaxo Wellcome	
		Urticaria		Paxil	SS	Glaxo Wellcome	
				Celebrex	SS		
				Neurontin	SS		
				Topamax	SS		

Klonopin	SS	
Progesterone	C	
Salbutamol	C	Glaxo Wellcome
Pentosane		
Polysulfate	C	
Hydroxyzine	C	
Amitriptyline	C	
Oxycodone		
Hydrochloride	C	

Date:07/15/02ISR Number: 3948177-6Report Type:Expedited (15-DaCompany Report #B0273503A
Age:64 YR Gender:Male I/FU:I

Outcome	PT
Life-Threatening	Acute Myocardial
Hospitalization -	Infarction
Initial or Prolonged	Cardiac Arrest

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Freedom Of Information (FOI) Report

Pancreatitis

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1TAB Twice per day 600MG Per day			Combivir	PS	Glaxo Wellcome	ORAL
			Efavirenz	SS		ORAL
			Salmeterol	SS	Glaxo Wellcome	
			Gabapentin	SS		
			Amiodarone	C		ORAL
			Carvedilol	C	Glaxo Wellcome	
			Ramipril	C		
			Furosemide	C	Glaxo Wellcome	

Date:07/16/02ISR Number: 3949718-5Report Type:Expedited (15-DaCompany Report #045-0945-M0200008
Age:20 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3200 MG, PER ORAL		Chromaturia Crystal Urine Present Nervous System Disorder	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
				Valproate Sodium	C		
				Fentanyl	C		
				Propofol	C		
				Aciclovir	C		

Date:07/16/02ISR Number: 3950010-3Report Type:Expedited (15-DaCompany Report #02P-056-0190595-00
Age:1 DY Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Clonic Convulsion Drug Withdrawal Convulsions Dysphagia Epilepsy Foetal Distress Syndrome	Foreign Health Professional Other	Depakine Chrono Tablets (Depakine) (Sodium Valproate/Valproic Acid) (Sodium Gabapentin	PS SS		

Hypoglycaemia
Maternal Drugs Affecting
Foetus
Neonatal Apnoeic Attack
Neonatal Disorder

Clonazepam

SS

Date:07/16/02ISR Number: 3950026-7Report Type:Expedited (15-DaCompany Report #2002050950
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcoholism	Health	Neurontin			
Other		Convulsion	Professional	(Gabapentin)	PS		ORAL
900 MG (300		Drug Interaction					
MD, DAILY),		Dysarthria					
ORAL		Medication Error		Ethanol	SS		
		Sudden Death					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/16/02ISR Number: 3950034-6Report Type:Expedited (15-DaCompany Report #2002050787

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Unevaluable Event	Health	Neurontin			
Initial or Prolonged		Professional	(Gabapentin)	PS		
Other			Digoxin	C		
			Magnesium	C		
			Multivitamins	C		
			Furosemide	C		
			Metolazone	C		
			Acetylsalicylic Acid	C		
			Risperidone	C		
			Spironolactone	C		
			Nystatin	C		
			Fentanyl	C		
			Hydrobromide			
			Hydrochloride	C		
			Gatifloxacin	C		
			Heparin-Fraction,			
			Sodium Salt	C		
			Salbutamol	C		
			Ipratropium Bromide	C		
			Carvedilol	C		
			Lisinopril	C		

Date:07/16/02ISR Number: 3950049-8Report Type:Expedited (15-DaCompany Report #2002051018

Age:52 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Blood Potassium Increased	Health	Neurontin			
Initial or Prolonged	Drug Screen Positive	Professional	(Gabapentin)	PS		ORAL
900 MG (300						

MG, THREE

TIMES DAILY),

ORAL

Lisinopril	SS
Amlodipine Besilate	C
Furosemide	C
Clonidine	C

Lamivudine	C
Stavudine	C
Kaletra	C
Vicodin	C
Bactrim	C
Methadone	C

Date:07/16/02ISR Number: 3950096-6Report Type:Expedited (15-DaCompany Report #2002105777US
Age:70 YR Gender:Male I/FU:I

Outcome	PT
Other	Areflexia
	Arthropathy
	Diabetic Neuropathy
	Dizziness
	Fall
	Gait Disturbance
	Intervertebral Disc
	Degeneration
	Neuralgia
	Osteoarthritis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Sedation Somnolence Vertigo	Report Source	Product	Role	Manufacturer	Route
200 MG, QD, ORAL			Health Professional	Celebrex	PS		ORAL
ORAL				Glucotrol(Glipizide)	SS		ORAL
300 MG, BID, ORAL				Neurontin(Gabapentin)	SS		ORAL
				Amitriptyline(Amitri ptyline)	SS		
				Zebeta	C		
				Lipitor (Atorvastatin)	C		
				Glucosamine(Glucosam ine)	C		
				Chondroitin	C		

Date:07/16/02ISR Number: 3950289-8Report Type:Expedited (15-DaCompany Report #001-0981-M0202719
Age: Gender:Female I/FU:F

Outcome Dose Other	Duration	PT Diabetes Mellitus Medication Error	Report Source	Product	Role	Manufacturer	Route
10 MG (UNKNOWN), UNKNOWN			Consumer	Atorvastatin (Atorvastatin)	PS		
600 MG (UNKNOWN), UNKNOWN				Gabapentin (Gabapentin)	SS		
UNKNOWN UNKNOWN				Glipizide	SS		

(UNKNOWN),

UNKNOWN

Human Mixtard	C
Enalapril Maleate	C
Furosemide	C
Metformin	
Hydrochloride	C
Ranitidine	C
Rosiglitazone	C
Diltiazem	C

Date:07/16/02ISR Number: 3950300-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200641
Age: Gender:Male I/FU:F

Outcome	PT
Other	Anger
	Asthenia
	Atrophy
	Balance Disorder
	Brain Neoplasm
	Burning Sensation
	Coordination Abnormal
	Delusion
	Depression
	Dizziness
	Eye Irritation
	Eye Movement Disorder
	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
		Fatigue Frequent Bowel Movements Haematochezia	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG		Haemorrhoids Hallucination					
(BID), ORAL		Incoherent					
UNKNOWN		Muscle Spasms		Zoloft (Sertraline)	SS		
(UNKNOWN),		Pollakiuria					
UNKNOWN		Polydipsia					
		Thinking Abnormal Vision Blurred					

Date:07/17/02ISR Number: 3949474-0Report Type:Direct Company Report #CTU 172439
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia		Neurontin 100mg			
Other		Dysuria		Long'S Drugs??	PS	Long'S Drugs??	ORAL
Required		Speech Disorder					
200MG 3X ORAL							
Intervention to							
Prevent Permanent							
Impairment/Damage							

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
Dose		Dyspnoea	Health	Klonopin	PS	Roche	
Other		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

Date:07/17/02ISR Number: 3950401-0Report Type:Expedited (15-DaCompany Report #2002050824
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (DAILY), ORAL		Bone Marrow Disorder Diarrhoea Haemodialysis	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
200 MG (100 MG, TWICE DAILY), ORAL		Pancytopenia Renal Failure Acute Renal Tubular Disorder		Ritonavir	SS		ORAL
300 MG (DAILY), ORAL				Tenofovir Disoproxil Fumarate	SS		ORAL
10 DROP (DAILY), ORAL (DAILY), ORAL				Clonazepam	SS		ORAL
				Bactrim	SS		ORAL
				Didanosine Amprenavir	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pentafuside C
 Alfuzosin C
 Loperamide Oxide C

Date:07/17/02ISR Number: 3950685-9Report Type:Expedited (15-DaCompany Report #2002050790
 Age:37 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Foreign	Neurontin			
		Automatism	Health	(Gabapentin)	PS		ORAL
1800 MG							
(BID), ORAL		Convulsion	Professional				
				Cannabis	C		
				Ethanol	C		

Date:07/17/02ISR Number: 3950718-XReport Type:Expedited (15-DaCompany Report #2002050488
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Clonic Convulsion	Foreign	Neurontin			
Initial or Prolonged		Dizziness	Health	(Gabapentin)	PS		ORAL
ORAL							
		Fall	Professional	Zolpidem	SS		ORAL
ORAL							
		Hallucination		Phenobarbital	SS		ORAL
ORAL							
		Loss Of Consciousness					

Date:07/17/02ISR Number: 3950722-1Report Type:Expedited (15-DaCompany Report #033-0945-M0200044
 Age:1 DY Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Apgar Score Low	Foreign	Neurontin			
Initial or Prolonged		Convulsion Neonatal	Health	(Gabapentin)	PS		
		Drug Withdrawal Syndrome	Professional	Ergenyl Chrono	SS		
1500 MG							

(BID), Neonatal
Feeding Problem In Clonazepam SS
0.5 MG
Newborn
(DAILY),
Foetal Distress Syndrome
Hypoglycaemia Neonatal
Maternal Drugs Affecting
Foetus
Neonatal Apnoeic Attack
Neonatal Disorder
Nuclear Magnetic
Resonance Imaging
Abnormal

Date:07/17/02ISR Number: 3950733-6Report Type:Expedited (15-DaCompany Report #2002050066
Age: Gender:Female I/FU:F

Outcome PT
Congenital Anomaly Anal Fissure
Complications Of Maternal
Exposure To Therapeutic
Drugs
Convulsion
Dermoid Cyst
Haemangioma
Maternal Drugs Affecting
Foetus

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pregnancy

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
3200 MG	(BID),	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:07/17/02ISR Number: 3950743-9Report Type:Expedited (15-DaCompany Report #2002050621
 Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign Health Professional	Neurontin (Gabapentin)	PS		
Other		Complications Of Maternal Exposure To Therapeutic					
3200 MG	(BID),	Drugs					
		Epilepsy Pregnancy					

Date:07/17/02ISR Number: 3950762-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200598
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer Health Professional	Neurontin (Gabapentin)	PS		
Hospitalization - Initial or Prolonged		Arthralgia Cerebrovascular Accident					
900 MG (300							
Other		Hypersomnia					
MG, TID)		Hypoaesthesia		Levothyroxine Sodium	C		
		Intervertebral Disc Disorder		Citalopram			
		Localised Infection		Hydrobromide	C		
		Pain In Extremity		Prinzide	C		
		Paraesthesia		Furosemide	C		
		Tendon Rupture		Potassium Chloride	C		
				Celecoxib	C		
				Rofecoxib	C		

Date:07/17/02ISR Number: 3951339-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200621
 Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aspartate	Health	Neurontin			
Other		Aminotransferase	Professional	(Gabapentin)	PS		ORAL
100 MG		Increased					
(DAILY), ORAL		Balance Disorder		Arthrotec	C		
		Blood Creatinine		Enalapril Maleate	C		
		Increased		Vaseretic	C		
		Blood Urea Increased		Amlodipine Besilate	C		
		Disturbance In Attention		Estradiol	C		
		Oedema Peripheral		Cyanocobalamin	C		
		Visual Disturbance		Chlorpheniramine			
				Maleate	C		
				Citracel With			
				Magnesium And D	C		
				Glucosamine And			
				Chondroitin With Msm	C		
				Flaxseed Oil	C		
				Evening Primrose Oil	C		
				Tocopherol	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Garlicscon C
 Cetirizine
 Hydrochloride C
 Clonazepam C

Date:07/18/02ISR Number: 3950645-8Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 172555

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	400 MG TID PO	Musculoskeletal Stiffness		Gabapentin 400 Mg	PS		ORAL
		Neck Pain		Amiodarone	C		
				Ec Aspirin	C		
				Cozaar	C		
				Kcl	C		
				Verapamil Sr	C		
				Digoxin	C		
				Pericolace	C		
				Lasix	C		
				Micronase	C		
				Lescol	C		
				Novolin 70/30	C		
				Sorbitol	C		

Date:07/18/02ISR Number: 3951104-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200710
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG		Bipolar Disorder	Consumer	Neurontin (Gabapentin)	PS		
(DAILY)				Lithium (Lithium)	SS		
900 MG							

Date:07/18/02ISR Number: 3951300-0Report Type:Expedited (15-DaCompany Report #033-0945-M0200069
 Age:17 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cyanosis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		Electroencephalogram					
2400 MG		Abnormal	Professional				
(TID), ORAL		Sudden Death	Company	Lamotrigine	SS		ORAL
350 MG (BID),			Representative				
ORAL							

Date:07/18/02ISR Number: 3951311-5Report Type:Expedited (15-DaCompany Report #2002051131
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alanine Aminotransferase Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		Gamma-Glutamyltransferase	Professional				
1600 MG		Increased		Dihydrocodeine	C		
(QID), ORAL		Oedema Peripheral		Dosulepin	C		
				Nifedipine	C		
				Tylox	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/18/02ISR Number: 3951452-2Report Type:Expedited (15-DaCompany Report #2002050930

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 6400 MG	Gastrointestinal Haemorrhage	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other (QID), ORAL	Urinary Retention		Topiramate	C		
			Olanzapine	C		
			Oxcarbazepine	C		
			Pantoprazole	C		
			Domperidone	C		
			Seretide Mite	C		
			Paracetamol	C		

Date:07/19/02ISR Number: 3951698-3Report Type:Expedited (15-DaCompany Report #2002050806

Age:83 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 3200 MG, ORAL	Coma Confusional State	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other			Glyceryl Trinitrate	C		
			Allopurinol	C		
			Piribedil	C		
			Fluindione	C		
			Valsartan	C		
			Amiodarone	C		
			Furosemide	C		

Date:07/19/02ISR Number: 3951704-6Report Type:Expedited (15-DaCompany Report #A200833

Age:63 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Required	Alanine Aminotransferase Increased Ascites	Foreign Health Professional	Hydroxyzine Hydrochloride (Hydroxyzine			

Intervention to ORAL	Aspartate	Hydrochloride)	PS	ORAL
Prevent Permanent Impairment/Damage 900 MG (TID), ORAL	Aminotransferase Increased	Gabapentin (Gabapentin)	SS	ORAL
	Blood Alkaline			
ORAL	Phosphatase Increased Gamma-Glutamyltransferase	Ranitidine (Ranitidine)	SS	ORAL
ORAL	Increased Liver Disorder	Valproic Acid (Valproic Acid)	SS	ORAL
1000 MG (BID), ORAL	Liver Function Test			
	Abnormal Lung Disorder Prothrombin Level	Glyceryl Trinitrate (Glyceryl Trinitrate)	SS	
TRANSDERMAL (DAILY),	10 MG			
TRANSDERMAL	Abnormal Urinary Tract Infection			
10 MG (DAILY), ORAL		Noctran 10 (Noctran 10)	SS	ORAL
		Lactulose (Lactulose)	C	
		Omeprazole (Omeprazole)	C	
		Enoxaparine Sodique (Heparin-Fraction, Sodium Salt)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lamotrigine
 (Lamotrigine) C
 Ceftriazone
 (Ceftriaxone) C
 Spiramycine
 (Spiramycin) C

Date:07/23/02ISR Number: 3952869-2Report Type:Expedited (15-DaCompany Report #033-0945-M0200002
 Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Deep Vein Thrombosis Gout	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL Other ORAL		Middle Insomnia Oedema Peripheral Pain Pyrexia Synovitis	Professional	Clopidogrel	SS		ORAL

Date:07/23/02ISR Number: 3952870-9Report Type:Expedited (15-DaCompany Report #2002051352
 Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Tremor	Foreign Literature	Gabapentin (Gabapentin)	PS		
900 MG (DAILY)				Oxcarbamazepine	SS		
1200 MG (DAILY)							

Date:07/23/02ISR Number: 3952932-6Report Type:Expedited (15-DaCompany Report #2002051354
 Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Encephalopathy	Foreign Literature	Gabapentin (Gabapentin)	PS		
3600 MG			Consumer				
(DAILY)							

Date:07/23/02ISR Number: 3952933-8Report Type:Expedited (15-DaCompany Report #2002050488
 Age:23 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Clonic Convulsion	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Fall	Professional	Zolpidem	SS		ORAL
ORAL		Hallucination		Phenobarbital	SS		ORAL
ORAL		Loss Of Consciousness					

Date:07/23/02ISR Number: 3953000-XReport Type:Expedited (15-DaCompany Report #033-0945-M0200060
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Interaction	Foreign Health	Neurontin (Gabapentin)	PS		
2400 MG (600 MG, QID),		Urticaria	Professional				
				Clonazepam	SS		
				Beclometasone	SS		

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Salmeterol	SS
Metelukast	SS
Venlafaxine	SS
Tramadol	SS
Cetirizine	C
Loratadine	C
Hydroxyzine	C
(Caffeine, Opium, Paracetamol)	C
Sertraline	C
Medrogestone	C

Date:07/23/02ISR Number: 3953016-3Report Type:Expedited (15-DaCompany Report #034-0945-M0200003
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Foreign	Gabapentin			
Other		Fall	Health	(Gabapentin)	PS		ORAL
1600 MG		Head Injury	Professional				
(QID), ORAL		Loss Of Consciousness	Company	Sertraline			
			Representative	(Sertraline)	SS		ORAL
150 MG (TID),				Lormetazepam	C		
ORAL				Ranitidine	C		
				Hydrochloride	C		

Date:07/23/02ISR Number: 3953147-8Report Type:Expedited (15-DaCompany Report #A201293
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Zoloft Tablets	PS		
Required		Balance Disorder	Health				
100.00 MG		Convulsion	Professional	Neurontin	SS		
Intervention to		Decreased Appetite		Klonopin	SS		
TOTAL:DAILY		Depression		Keppra	C		
Prevent Permanent							
Impairment/Damage							

Drug Withdrawal Syndrome
Energy Increased
Fatigue
Headache
Hormone Level Abnormal
Irritability
Libido Decreased
Malaise
Palpitations
Sedation
Sleep Disorder
Weight Decreased

Gaba

C

Date:07/23/02ISR Number: 3953769-4Report Type:Expedited (15-DaCompany Report #2002050928
Age: Gender:Female I/FU:I

Outcome PT
Other Bedridden
Cluster Headache
Eating Disorder
Hypothyroidism
Mental Disorder
Pyrexia

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Dose	Duration	Therapeutic Agent Toxicity Tooth Disorder	Report Source	Product	Role	Manufacturer	Route
(QID), ORAL		Weight Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Venlafaxine Hydrochloride	C		
				Paroxetine Hydrochloride	C		
				Tramadol Hydrochloride	C		
				Clonazepam	C		
				Carisoprodol	C		

Date:07/23/02ISR Number: 3953771-2Report Type:Expedited (15-DaCompany Report #2002051379
Age: Gender:Female I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Balance Disorder	Health	Neurontin (Gabapentin)	PS		
		Chest Pain	Professional	Betaseron	SS		
SUBCUTANEOUS	8 I.U.	Condition Aggravated					
SUBCUTANEOUS		Liver Function Test					
		Abnormal		Nifedipine (Nifedipine)	SS		
		Multiple Sclerosis		Fluoxetine Hydrochloride	C		
		Oedema Peripheral		Bupropion			
		Pain In Extremity		Hydrochloride	C		
		Peripheral Vascular Disorder		Hydrochloride	C		
		Skin Discolouration		Tizanidine	C		
				Alprazolam	C		
				Di-Gesic	C		
				Estradiol	C		

Date:07/23/02ISR Number: 3953838-9Report Type:Expedited (15-DaCompany Report #2002051308
Age:46 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization -	Coma	Literature	Gabapentin	
Initial or Prolonged	Drug Toxicity	Health	(Gabapentin)	PS
Other	Haemodialysis	Professional		
	Hyporeflexia			
	Hypoxia			
	Mental Status Changes			
	Oxygen Saturation			
	Decreased			
	Somnolence			
	Tremor			

Date:07/23/02ISR Number: 3953840-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200241
Age: Gender:Female I/FU:F

Outcome	PT
Other	Alopecia
	Contusion
	Dizziness
	Drug Ineffective
	Drug Withdrawal Syndrome
	Face Oedema

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fatigue Urticaria Weight Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL				Sertraline Hydrochloride (Sertraline)	SS		ORAL
ORAL				Carisoprodol Prednisone	C C		

Date:07/23/02ISR Number: 3953846-8Report Type:Expedited (15-DaCompany Report #2002051331
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Aspartate Aminotransferase Abnormal Blood Albumin Abnormal Blood Alkaline Phosphatase Increased Blood Cholesterol Increased Gamma-Glutamyltransferase Increased Haemangioma Of Liver Intraocular Pressure Increased					

Date:07/24/02ISR Number: 3953361-1Report Type:Expedited (15-DaCompany Report #2002051326
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Other		Convulsion Tardive Dyskinesia	Consumer	Neurontin (Gabapentin) Sertraline	PS		

Hydrochloride

SS

Date:07/24/02ISR Number: 3953573-7Report Type:Expedited (15-DaCompany Report #2002051363
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dyspnoea	Foreign	Gabapentin			
Hospitalization - 400 MG		Hypertension	Consumer	(Gabapentin)	PS		ORAL
Initial or Prolonged (DAILY), ORAL		Malaise					
		Respiratory Failure		Cloxazolam	C		
				Amitriptyline			
				Hydrochloride	C		
				Ascorbic Acid	C		
				Tocopherol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/25/02ISR Number: 3954061-4Report Type:Expedited (15-DaCompany Report #001-0945-M0200552
 Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Consumer	Neurontin			
		Blood Pressure	Health	(Gabapentin)	PS		
1200 MG (300							
MG, FOUR		Fluctuation	Professional				
		Dizziness					
		Fatigue		Arthrotrec	C		
		Medication Error		Citalopram			
		Nausea		Hydrobromide	C		
		Pain		Tizanidine	C		
		Weight Increased		Rabeprazole	C		
				Loratadine	C		
				Acetylsalicylic Acid	C		
				(Multiple Vitamin)	C		
				Vicodin	C		

Date:07/25/02ISR Number: 3954062-6Report Type:Expedited (15-DaCompany Report #001-0945-M0200668
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Urine Analysis Abnormal	Consumer	Neurontin			
Initial or Prolonged				(Gabapentin)	PS		ORAL
ORAL							
Other				Phenytoin Sodium	SS		ORAL
ORAL							

Date:07/25/02ISR Number: 3954332-1Report Type:Expedited (15-DaCompany Report #2002051479
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Catheter Related	Consumer	Neurontin			
Initial or Prolonged		Infection		(Gabapentin)	PS		ORAL
1200 MG (300							
MG, FOUR		Convulsion					

TIMES DAILY),

Cyst

Drug Effect Decreased

ORAL

Pain In Extremity

Dilantin (Phenytoin Sodium)	SS
Morphine	SS
Unspecified	
Narcotics	SS
Clopidogrel	C
Venlafaxine	
Hydrochloride	C
Clonidine	C
Metoprolol Tartrate	C
Atorvastatin	C

Date:07/26/02ISR Number: 3956638-9Report Type:Expedited (15-DaCompany Report #2002050190

Age:71 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Cardiac Failure Hypertension Hypertensive Crisis Pulmonary Oedema	Foreign Health Professional	Neurontin (Gabapentin) Carbimazole Bisoprolol Fumarate Digoxin Pantoprazole Sodium Thioctic Acid	PS C C C C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/26/02ISR Number: 3956894-7Report Type:Expedited (15-DaCompany Report #2002051595

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser	Health Professional	Neurontin (Gabapentin)	PS		

Date:07/30/02ISR Number: 3955962-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200624

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain Condition Aggravated	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
1200 MG (400 MG, TID),		Cyst	Professional				
ORAL		Headache					
45 MG (15 MG, TID), ORAL		Neuralgia		Morphine	SS		ORAL
		Spinal Cord Disorder					
				Panadeine Co	C		
				Rofecoxib	C		
				Oxybutynin	C		

Date:07/30/02ISR Number: 3956001-0Report Type:Expedited (15-DaCompany Report #2002051523

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Alopecia	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG		Confusional State					
Other (DAILY), ORAL		Convulsion					
		Disease Recurrence		Topiramate	C		
		Dysarthria		Estrogen Nos	C		
		Electroencephalogram Abnormal		Levothyroxine Sodium	C		
		Fatigue		Amitriptyline	C		

Feeling Drunk
Gingival Pain
Hypoaesthesia
Vision Blurred

Date:07/30/02ISR Number: 3956002-2Report Type:Expedited (15-DaCompany Report #2002051524

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Neurontin (Gabapentin)	PS		

Date:07/30/02ISR Number: 3956003-4Report Type:Expedited (15-DaCompany Report #2002051526

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/30/02ISR Number: 3956004-6Report Type:Expedited (15-DaCompany Report #2002051530

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Neurontin (Gabapentin)	PS		

Date:07/30/02ISR Number: 3956005-8Report Type:Expedited (15-DaCompany Report #2002051532

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Neurontin (Gabapentin)	PS		

Date:07/30/02ISR Number: 3956006-XReport Type:Expedited (15-DaCompany Report #2002051533

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Neurontin (Gabapentin)	PS		

Date:07/30/02ISR Number: 3956007-1Report Type:Expedited (15-DaCompany Report #2002051535

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Neurontin (Gabapentin)	PS		

Date:07/30/02ISR Number: 3956008-3Report Type:Expedited (15-DaCompany Report #2002051537

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Neurontin (Gabapentin)	PS		

Date:07/30/02ISR Number: 3956009-5Report Type:Expedited (15-DaCompany Report #2002051538

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia Convulsion	Consumer	Neurontin(Gabapentin)	PS		

Date:07/30/02ISR Number: 3956610-9Report Type:Expedited (15-DaCompany Report #2002051718

Age:85 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Failure Hallucination	Foreign Health	Neurontin (Gabapentin)	PS		
100 MG (DAILY),		Renal Failure	Professional Company Representative	Tramadol Diltiazem Candesartan Glyceryl Trinitrate Furosemide Digoxin	C C C C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/31/02ISR Number: 3957199-0Report Type:Expedited (15-DaCompany Report #2002116944US

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Drug Interaction	Consumer	Bextra			
Initial or Prolonged	Hypertension		(Valdecoxib)	PS		ORAL
ORAL	7 DAY					
			Neurontin			
ORAL			(Gabapentin)	SS		ORAL
			Nexium	SS		ORAL
ORAL						
			Prilosec			
ORAL			(Omeprazole)	C		ORAL

Date:08/01/02ISR Number: 3958093-1Report Type:Expedited (15-DaCompany Report #A125321

Age:24 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Burning Sensation	Foreign	Doxepin(Doxepin)	PS		
Initial or Prolonged	Drug Abuser	Health	Neurontin(Gabapentin			
	Drug Toxicity	Professional)	SS		ORAL
600 MG, ORAL						
	General Physical Health		Heroin(Diamorphine)	SS		
	Deterioration		Cocaine (Cocaine)	SS		
	Hepatitis C Antibody		Benzodiazepines(Benz			
	Positive		odiazepine			
	Medication Error		Derivatives)	SS		
	Movement Disorder		Alcohol(Ethanol)	SS		
	Neuropathy Peripheral					
	Paraparesis					
	Rhabdomyolysis					
	Somnolence					
	Speech Disorder					

Date:08/01/02ISR Number: 3958181-XReport Type:Expedited (15-DaCompany Report #A202614

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					

Required	Blood Sodium Abnormal	Health	Zoloft Tablets	PS
Intervention to	Blood Testosterone	Professional	Neurontin	SS
Prevent Permanent	Decreased		Paxil	SS
Impairment/Damage			Wellbutrin	SS
			Ambien	C
			Albuterol Inhaler	C
			Atenolol	C
			Folic Acid	C
			Zocor	C
			Hydrochlorothiazide	C

Date:08/01/02ISR Number: 3958215-2Report Type:Expedited (15-DaCompany Report #2002051969
Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG(TID), Other ORAL		Blood Creatine Phosphokinase Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Risperidone	SS		ORAL
				Topiramate	C		
				Salbutamol	C		
				Pseudoephedrine			
				Hydrochloride	C		
				Paracetamol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/01/02ISR Number: 3958339-XReport Type:Expedited (15-DaCompany Report #2002051910

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphasia	Consumer	Neurontin			
		Asthenia		(Gabapentin)	PS		ORAL
	(THREE TIMES						
	DAILY), ORAL	Dizziness					
	ORAL	Drug Effect Decreased		Grapefruit	SS		ORAL
		Dysarthria		Oxycodone			
		Dysgraphia		Hydrochloride	C		
		Fall		Oxycocet	C		
		Food Interaction		Carisoprodol	C		
		Loss Of Consciousness		Hydrochlorothiazide	C		
		Memory Impairment					
		Photosensitivity Reaction					
		Somnolence					
		Stress					
		Vision Blurred					

Date:08/01/02ISR Number: 3958348-0Report Type:Expedited (15-DaCompany Report #2002051916

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis	Consumer	Neurontin			
		Back Pain		(Gabapentin)	PS		ORAL
	400 MG						
	(DAILY) ORAL	Constipation					
		Diarrhoea		Valdecoxib	SS		
		Dizziness		Applesauce	SS		
		Dysphagia		Atorvastatin	C		
		Foreign Body Trauma		Estrogens Conjugated	C		
		Intervertebral Disc Disorder					
		Respiratory Distress					
		Somnolence					
		Tinnitus					

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract	Consumer	Neurontin			
		Condition Aggravated		(Gabapentin)	PS		ORAL
ORAL		Dizziness		Magnesium	SS		
		Fatigue		Calcium	SS		
		Guillain-Barre Syndrome		Aluminum	SS		
		Nausea		Acetylsalicylic Acid	C		
				Naratriptan			
				Hydrochloride	C		
				Riboflavin	C		
				Multivitamins	C		

Age: Gender:Female I/FU:F

Outcome	PT
Required	Amenorrhoea
Intervention to	Anxiety
Prevent Permanent	Mood Swings
Impairment/Damage	Neutropenia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Red Blood Cell Count Decreased					
		Tearfulness	Health	Zoloft Tablets	PS		ORAL
DAILY:ORAL		White Blood Cell Count	Professional	Neurontin	SS		ORAL
1600.00 MG		Decreased					
TOTAL:BID:ORA							
L				Risperdal	SS		
2.00 MG							
TOTAL:DAILY							
				Depakote	C		
				Benadryl	C		
				Xanax	C		

Date:08/01/02ISR Number: 3958852-5Report Type:Expedited (15-DaCompany Report #2002051909
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Neurontin (Gabapentin)	PS		
Other		Lip Pain					
900 MG		Pain					
(DAILY),		Weight Increased		Amitriptyline Hydrochloride	C		

Date:08/02/02ISR Number: 3959867-3Report Type:Periodic Company Report #2001AP05240
 Age:54 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Amitriptyline	PS		
			Health	Gabapentin	SS		
			Professional	Zolpidem	SS		

Date:08/05/02ISR Number: 3959042-2Report Type:Expedited (15-DaCompany Report #2002-04461
 Age:68 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 300 MG QD PO	Bone Marrow Depression	Foreign	Tenofovir Df	PS		ORAL
Hospitalization - Initial or Prolonged	Confusional State	Study	Didanosine	SS		
	Dehydration	Health	Amprenavir	SS		
	Dialysis	Professional	Ritonavir	SS		
	Hiv Wasting Syndrome	Other	Bactrim	SS		
	Pancytopenia		Ibuprofen	SS		
	Pathogen Resistance		Gabapentin	SS		
	Renal Failure Acute		T20	C		
	Renal Tubular Necrosis		Fenofibrate	C		
	Staphylococcal Infection					
	Urinary Tract Infection					
	Viral Diarrhoea					

Date:08/05/02ISR Number: 3959118-XReport Type:Expedited (15-DaCompany Report #A0376571A
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Arthritis	Consumer	Sine-Off (Sine-Off)	PS		
	Bursitis		Guaiphenesin (Guaifenesin)	SS		
	Cerebrovascular Accident		Ibuprofen (Ibuprofen)	SS		
	Coronary Artery Surgery					
	Surgery					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dimetapp (Dimetapp)	SS
Alka-Seltzer Plus Cold Md (Alka-Seltzer Plus Cold Md)	SS
Tavist-D (Tavist-D)	SS
Triaminic (Triaminic)	SS
Phenylpropanolamine Hcl (Phenylpropanolamine)	SS
Vicodin (Vicodin)	SS
Clonidine (Clonidine)	SS
Coricidin (Coricidin)	SS
Tylenol Cold Medication (Tylenol Cold Medication)	SS
Medrol (Medrol)	SS
Rofecoxib (Rofecoxib)	SS
Ibuprofen (Ibuprofen)	SS
Gabapentin (Gabapentin)	SS
Vicks Dayquil (Vicks Dayquil)	SS
Nyquil (Nyquil)	SS

Date:08/05/02ISR Number: 3960125-1Report Type:Expedited (15-DaCompany Report #USA-2002-0001676
Age:40 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Oedema Peripheral Pitting Oedema	Literature Health Professional	Oxycodone Hydrochloride (Similar To Nda-20553) (Oxycodone	PS		ORAL
240 MG Q12H			Morphine Sulfate			
ORAL						

ORAL	(Similar To Nda 19-516) (Morphine Sulfate) Cr Tablet	SS	ORAL
ORAL	Morphine Sulfate (Similar To Nda 19-515) (Morphine Sulfate) Ir Tablet	SS	ORAL
ORAL	Gabapentin (Gabapentin)	SS	ORAL
	Amitriptyline (Amitriptyline)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/06/02ISR Number: 3959231-7Report Type:Expedited (15-DaCompany Report #2002052306
Age:83 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (THREE TIMES DAILY), ORAL	Arrhythmia Condition Aggravated Dyspnoea Exacerbated	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
			Amitriptyline Ginkgo Biloba Omeprazole Verapamil Spironolactone Methadone Hydrochloride Meloxicam Homeopathic Drug	C C C C C C C C C		

Date:08/06/02ISR Number: 3959232-9Report Type:Expedited (15-DaCompany Report #2002052324
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL	Haemorrhage Prostatic Operation	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:08/06/02ISR Number: 3959317-7Report Type:Expedited (15-DaCompany Report #2002051018
Age:52 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 MG, THREE	Blood Potassium Increased Drug Screen Positive	Health Professional	Neurontin (Gabapentin)	PS		ORAL

TIMES DAILY),

ORAL

20 MG

(DAILY), ORAL

Lisinopril

SS

ORAL

Amlodipine Besilate

C

Furosemide

C

Clonidine

C

Lamivudine

C

Stavudine

C

Kaletra

C

Vicodin

C

Bactrim

C

Methadone

C

Lopinavir

C

Date:08/06/02ISR Number: 3959318-9Report Type:Expedited (15-DaCompany Report #2002052184

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Surgery	Consumer	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/06/02ISR Number: 3959320-7Report Type:Expedited (15-DaCompany Report #2002051971
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nephrolithiasis Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG (600 MG, TWICE DAILY), ORAL							
				Paroxetine Hydrochloride	C		
				Amlodipine Besilate	C		
				Simvastatin	C		

Date:08/06/02ISR Number: 3959322-0Report Type:Expedited (15-DaCompany Report #2002051970
 Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aspartate Aminotransferase	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (DAILY), ORAL							
		Increased					

Date:08/06/02ISR Number: 3959345-1Report Type:Expedited (15-DaCompany Report #2002052154
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Dysgraphia	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (TID), ORAL							
				Sertraline Hydrochloride	C		
				Oxycocet	C		
				Levetiracetam	C		

Date:08/06/02ISR Number: 3959347-5Report Type:Expedited (15-DaCompany Report #2002052153
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Ejaculation Failure	Consumer	Neurontin			
Initial or Prolonged	Self Mutilation		(Gabapentin)	PS		ORAL
ORAL						

Date:08/06/02ISR Number: 3959363-3Report Type:Expedited (15-DaCompany Report #2002052155
Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Diabetes Mellitus	Consumer	Neurontin			
Initial or Prolonged	Diarrhoea		(Gabapentin)	PS		ORAL
300 MG (100						
Other	Difficulty In Walking					
MG, THREE						
	Dyspnoea					
TIMES DAILY),						
	Gastrointestinal					
ORAL						
	Infection		(Unspecified			
	Pain		Intravenous Therapy)	SS		
INTRAVENOUS	INTRAVENOUS					
	Pyrexia		Antibiotics	C		
			Thiamine	C		
			Multivitamins	C		
			Ranitidine			
			Hydrochloride	C		
			Insulin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/06/02ISR Number: 3959365-7Report Type:Expedited (15-DaCompany Report #2002051973

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Dizziness					
		Drug Ineffective		Sildenafil Citrate	SS		ORAL
(PRN), ORAL		Drug Tolerance Increased		Fexofenadine			
		Ejaculation Disorder		Hydrochloride	SS		ORAL
ORAL		Headache					
		Hypersensitivity					
		Migraine					
		Vertigo					

Date:08/06/02ISR Number: 3959400-6Report Type:Expedited (15-DaCompany Report #2002052041

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 1500 MG		Psychotic Disorder	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
(TID), ORAL			Professional				
			Company Representative	Amlodipine Besilate	C		
				Olanzapine	C		
				Valproate Sodium	C		
				Reboxetine	C		

Date:08/06/02ISR Number: 3959416-XReport Type:Expedited (15-DaCompany Report #2002052157

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Epilepsy	Foreign Health	Neurontin (Gabapentin)	PS		
1800 MG DAILY		Speech Disorder					
			Professional Company				

Representative

Date:08/06/02ISR Number: 3959417-1Report Type:Expedited (15-DaCompany Report #2002052038
 Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Epilepsy	Foreign	Neurontin			
Other		Injury Asphyxiation	Health	(Gabapentin)	PS		
300 MG DAILY			Professional Company Representative				

Date:08/06/02ISR Number: 3959418-3Report Type:Expedited (15-DaCompany Report #2002050997
 Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Effect Decreased Dysarthria	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
Other ORAL		Incoherent					
27 TABLETS		Intentional Misuse		Paracetamol	SS		ORAL
ONCE, ORAL		Suicide Attempt					
ORAL				Mirtazapine	SS		ORAL
				Alendronate Sodium Provella-14	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrocortisone C
 Multivitamins C
 B-Komplex Leciva C
 Calcium Carbonate C
 Morphine Sulfate C
 Venlafaxine C
 Lactulose C
 Ergocalciferol C

Date:08/06/02ISR Number: 3959626-1Report Type:Expedited (15-DaCompany Report #2002052323
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
1800 MG (600		Drug Ineffective		(Gabapentin)	PS		ORAL
MG, TID),		Epileptic Aura					
ORAL		Grand Mal Convulsion					
		Pain		Hydrochlorothiazide	C		
		Paraesthesia		Cetirizine			
		Pharmaceutical Product		Hydrochloride	C		
		Complaint		Tramadol			
				Hydrochloride	C		
				Metaxalone	C		
				Metoprolol	C		

Date:08/06/02ISR Number: 3959634-0Report Type:Expedited (15-DaCompany Report #A123280
 Age:71 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acute Myocardial	Health	Doxepin (Doxepin)	PS		
Hospitalization -		Infarction	Professional	Gabapentin			
Initial or Prolonged		Cerebrovascular Accident		(Gabapentin)	SS		
Required		Chest Pain		Roxanol (Morphine			
Intervention to		Coma		Sulfate)	SS		
Prevent Permanent		Drug Toxicity		Ativan (Lorazepam)	SS		
Impairment/Damage		Emphysema		Humulin (Novolin			
		Hypertension		20/80)	SS		
		Overdose		Percocet (Oxycocet)	SS		

Thermal Burn

Theodur
(Theophylline) C
Prednisone
(Prednisone) C
Imdur (Isosorbide
Mononitrate) C
Aldomet (Methyldopa) C
Lasix (Furosemide) C
Relafen (Nabumetone) C
Skelaxin
(Metaxalone) C

Date:08/06/02ISR Number: 3959652-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200696

Age: Gender:Female I/FU:F

Outcome PT
Hospitalization - Blood Thyroid Stimulating
Initial or Prolonged Hormone Increased
Other Drug Interaction
Malaise

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Medication Error
Thyroxine Free Decreased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
3000 MG	(UNKNOWN),	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL			Levothyroxine	SS		ORAL
ORAL			Clonidine Unspecified Medication	C		

Date:08/06/02ISR Number: 3959657-1Report Type:Expedited (15-DaCompany Report #A208993
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	200.00 MG	Coma	Health Professional	Zoloft Tablets	PS		ORAL
Hospitalization - TOTAL:DAILY:0		Drug Interaction					
Initial or Prolonged RAL		Pneumonia Streptococcal					
Required ORAL				Gabapentin	SS		ORAL
Intervention to Prevent Permanent Impairment/Damage				Elavil Premarin	C C		

Date:08/06/02ISR Number: 3959658-3Report Type:Expedited (15-DaCompany Report #A208992
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	100.00 MG	Coma	Health Professional	Zoloft Tablets	PS		ORAL
Hospitalization - TOTAL:DAILY:0		Drug Interaction					

Initial or Prolonged RAL	Feeling Drunk			
Required 3600.00 MG	Gait Disturbance	Gabapentin	SS	ORAL
Intervention to TOTAL:TID:ORA	Hallucination			
Prevent Permanent L Impairment/Damage	Influenza			
	Pneumonia	Anafranil	SS	
	Staphylococcal Infection	Slo-Bid Gyrocaps	C	
		Q-Tuss	C	
		Klor-Con	C	
		Albuterol	C	
		Premarin	C	
		Lotensin	C	
		Baycol	C	
		Xanax	C	
		Furosemide	C	
		Prednisone	C	
		Trazodone	C	
		Prilosec	C	
		Metoclopramide	C	
		Atrovent	C	
		Intal	C	
		Pulmicort	C	
		Miacalcin	C	
		Nasonex	C	
		Gaviscon Type		
		Unknown	C	
		Darvocet	C	
		Lonox	C	
		Prochlorperazine	C	
		Carafate	C	

Freedom Of Information (FOI) Report

Hydromet	C
Calcium With Vitamin	
D	C
Serevent	C
Singulair	C
Vitamin E	C
Ocuvite Extra	C
B-12	C
Osteo-Bi-Flex	C
Prinivil	C
Clonazepam	C
Protonix	C
Pepcid	C
Zyrtec	C
Mirapex	C

Date:08/06/02ISR Number: 3959764-3Report Type:Expedited (15-DaCompany Report #2002052182
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anal Spasm	Consumer	Neurontin			
1400 MG		Diverticulitis		(Gabapentin)	PS		
(DAILY)		Gastritis					
		Gastrointestinal		Citalopram			
		Obstruction		Hydrobromide	SS		
		Gastrointestinal Pain		Hyoscyamine	SS		
		Gastroesophageal Reflux					
		Disease					
		Headache					
		Hiatus Hernia					
		Irritable Bowel Syndrome					
		Oesophagitis					

Date:08/06/02ISR Number: 3960209-8Report Type:Expedited (15-DaCompany Report #A209649
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State	Consumer	Lithium (Lithium)	PS		ORAL
(TID), ORAL							

300 MG	Depression	Health	Gabapentin	SS	ORAL
(DAILY), ORAL	Dizziness	Professional	(Gabapentin)		
	Fatigue				
	Nausea		Prevacid		
	Osteoporosis		(Lansoprazole)	C	
	Parkinson'S Disease		Unknown Stool		
			Softener		
			(Laxative-S)	C	
			Welbutrin Sr		
			(Bupropion		
			Hydrochloride)	C	
			Propranolol	C	

Date:08/08/02ISR Number: 3960586-8Report Type:Expedited (15-DaCompany Report #2002052511
Age:61 YR Gender:Female I/FU:I

Outcome PT
Hospitalization - Aspiration
Initial or Prolonged Blood Glucose Increased
Blood Pressure Decreased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
54 GRAM	(ONCE), ORAL	Coma Drug Level Above Therapeutic Electrocardiogram Qt Corrected Interval Prolonged	Literature	Gabapentin (Gabapentin)	PS		ORAL
		Electrocardiogram St Segment Abnormal Mydriasis Overdose Pco2 Decreased Pyrexia Vomiting White Blood Cell Count Increased		Quetiapine Bupropion Clonazepam Clopidogrel Glibenclamide Losartan Metoprolol	SS C C C C C C		

Date:08/08/02ISR Number: 3960912-XReport Type:Expedited (15-DaCompany Report #2002052313
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 MG TID)		Crying Haemorrhage Ill-Defined Disorder Medication Error Nephrolithiasis Pain Procedural Complication	Consumer	Neurontin (Gabapentin) Warfarin Sodium	PS SS		

Date:08/09/02ISR Number: 3961523-2Report Type:Expedited (15-DaCompany Report #2002052323
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG (600		Convulsion Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL

MG, TID),	Epileptic Aura		
	Grand Mal Convulsion		
ORAL			
	Medication Error	Hydrochlorothiazide	C
	Pain	Cetirizine	
	Pharmaceutical Product	Hydrochloride	C
	Complaint	Tramadol	
		Hydrochloride	C
		Metaxalone	C
		Metoprolol	C

Date:08/09/02ISR Number: 3962048-0Report Type:Expedited (15-DaCompany Report #2002052421
Age: Gender:Female I/FU:I

Outcome	PT
Other	Arthritis
	Blood Glucose Decreased
	Cardiac Failure
	Congestive
	Dehydration
	Dizziness
	Feeling Abnormal
	Fluid Retention
	Intracardiac Thrombus
	Loss Of Consciousness

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Myocardial Infarction Pulmonary Embolism Weight Increased	Report Source	Product	Role	Manufacturer	Route
1600 MG (800 MG, BID), ORAL ORAL ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Troglitazone	SS		ORAL
				Atorvastatin	SS		
				Glibenclamide	SS		
				Celecoxib	SS		ORAL
				Metformin Hydrochloride	SS		
				Dopamine	C		
				("Elder" Something)	C		
				("A" Drug)	C		
				Repaglinide	C		
				Insulin	C		
				Furosemide	C		
				Hydrochlorothiazide	C		
				Potassium	C		
				Enalapril	C		
				Carvedilol	C		
				Theophylline	C		
				Paroxetine Hydrochloride	C		
				Ranitidine	C		
				Trazodone	C		
				Warfarin	C		
				Codeine	C		
				Diclofenac Sodium	C		

Date:08/09/02ISR Number: 3962703-2Report Type:Expedited (15-DaCompany Report #2002052306

Age:83 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Arrhythmia	Foreign	Gabapentin			

Initial or Prolonged Dyspnoea Consumer (Gabapentin) PS ORAL
 1200 MG
 (THREE TIMES
 DAILY), ORAL

Amitriptyline C
 Ginkgo Biloba C
 Omeprazole C
 Varapamil C
 Spironolactone C
 Methadone
 Hydrochloride C
 Meloxicam C
 Homeopathic Drug C

Date:08/12/02ISR Number: 3961579-7Report Type:Expedited (15-DaCompany Report #A100405
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 10 MG	Back Disorder	Consumer	Norvasc (Amlodipine)	PS		
Initial or Prolonged	Blood Pressure Increased	Health	Lipitor			
	Depression	Professional	(Atorvastatin)	SS		
	Hysterectomy		Trazodone			
	Weight Increased		(Trazodone)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochlorothiazide
 (Hydrochlorothiazide) SS
 Neurontin
 (Gabapentin) SS
 Atenolol (Atenolol) SS
 Vasotex (Enalapril Maleate) C

Date:08/12/02ISR Number: 3961658-4Report Type:Expedited (15-DaCompany Report #2002052663
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Pain In Extremity Visual Disturbance	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG (DAILY) , ORAL				Tamoxifen Medroxyprogesterone Acetate Aporex Gaviscon	C C C C		

Date:08/12/02ISR Number: 3961660-2Report Type:Expedited (15-DaCompany Report #2002052509
 Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (DAILY) ,		Cerebrovascular Accident Somnolence Vertigo	Foreign Consumer	Neurontin (Gabapentin)	PS		
				Other Antihypertensives	C		

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Systolic Decreased	Health Professional	Neurontin (Gabapentin)	PS		
		Coordination Abnormal		Carbamazepine	SS		
400 MG (BID)							
		Drug Interaction Drug Level Increased Somnolence		Meclozine	SS		

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Tramadol Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/12/02ISR Number: 3961985-0Report Type:Expedited (15-DaCompany Report #2002AP02500
Age:61 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening USUAL DAILY	Aspiration	Literature	Quetiapine	PS		
Hospitalization - DOSE 200 MG Initial or Prolonged 54 G ONCE PO	Coma	Health				
	Drug Level Above	Professional	Gabapentin	SS		ORAL
	Therapeutic		Bupropion	C		
	Electrocardiogram Qt Corrected Interval Prolonged		Klonopin	C		
	Electrocardiogram St Segment Abnormal		Clopidogrel	C		
	Hypotension		Glyburide	C		
	Overdose		Losartan	C		
	Pyrexia		Metoprolol	C		
	Respiratory Depression					
	Vomiting					
	White Blood Cell Count Increased					

Date:08/12/02ISR Number: 3962012-1Report Type:Expedited (15-DaCompany Report #2002052622
Age:77 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death Hospitalization - 400 MG (TID) Initial or Prolonged ORAL	Atrial Fibrillation	Health	Neurontin			
	Blood Pressure Abnormal	Professional	(Gabapentin)	PS		ORAL
Other	Cardiac Arrest					
	Cerebrovascular Accident		Rabeprazole	C		
	Cold Sweat		Absorbace	C		
	Headache		Galenic/Paracetamio/			
	Heart Sounds Abnormal		Codeine	C		
			Bisacodyl	C		
			Citalopram	C		
			Simvastatin	C		
			Metoprolol	C		
			Lisinopril	C		
			Senna	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Bilirubin Increased	Foreign Health	Keppra	PS		
2500 MG		Coagulopathy		Keppra	SS		
1500 MG		Epilepsy	Professional	Keppra	SS		
1000 MG		Haemorrhage	Other	Keppra	SS		
2000 MG		Hepatic Failure		Keppra	SS		
900 MG		Hepatic Fibrosis		Gabapentin	SS		
1500 MG		Jaundice		Gabapentin	SS		
900 MG		Platelet Count Decreased		Gabapentin	SS		
300 MG		Status Epilepticus		Valproic Acid	SS		
600 MG				Carbamazepine	SS		
900 MG				Carbamazepine	SS		
				Oxcarbamazepine	SS		

Ranitidine	C
Trazodone	C
Warfarin	C
Codeine	C
Diclofenac Sodium	C

Date:08/13/02ISR Number: 3963288-7Report Type:Expedited (15-DaCompany Report #2002051331
Age:52 YR Gender:Female I/FU:I

Outcome	PT
Other	Alanine Aminotransferase Increased Aspartate Aminotransferase Increased Blood Albumin Increased Blood Alkaline Phosphatase Increased Blood Cholesterol Increased Condition Aggravated Gamma-Glutamyltransferase Increased Haemangioma

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pain

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
ORAL		Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Professional				

Date:08/14/02ISR Number: 3962171-0Report Type:Direct Company Report #CTU 174125
 Age:92 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300MG BID		Hallucination		Neurontin 300mg	PS		ORAL
ORAL							

Duragesic Patches C
 Senna C
 Atenolol C
 Furosemide C
 Levaquin C

Date:08/14/02ISR Number: 3963535-1Report Type:Expedited (15-DaCompany Report #2002UW10802
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Pressure Increased Drug Interaction	Other	Nexium Prilosec Neurontin Bextra	PS SS SS SS		

Date:08/15/02ISR Number: 3962880-3Report Type:Direct Company Report #CTU 174280
 Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Gabapentin			

Hospitalization -	Cerebrovascular Accident	(Neurontin)	PS	ORAL
400 MG PO TID				
Initial or Prolonged	Cold Sweat	Absorbasc Oint	C	
		Acetaminophen	C	
		Bisacodyl Supp	C	
		Citalopram	C	
		Lisinopril	C	
		Metoprolol	C	
		Rabeprazole	C	
		Senna	C	
		Simvastatin	C	
		Tiazem	C	
		Warfarin	C	

Date:08/15/02ISR Number: 3962881-5Report Type:Direct Company Report #CTU 174281
Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Gabapentin (Neurontin)	PS		ORAL
600 MG PO FOR							
PAIN				Morphine	C		
				Olsalazine	C		
				Promethazine	C		
				Rabeprazole	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Senna C
Lorazepam C

Date:08/15/02ISR Number: 3963872-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200667
Age:62 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 100 MG Other (DAILY) ORAL	Duration Depression Dizziness Drug Effect Decreased Fall Gait Disturbance Head Injury Retinopathy	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Nifedipine	C		
			Captopril	C		
			Labetalol	C		
			Citalopram Hydrobromide	C		

Date:08/15/02ISR Number: 3963875-6Report Type:Expedited (15-DaCompany Report #001-0945-M0200754
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL	Duration Disturbance In Attention Dizziness Memory Impairment Somnolence Tendon Disorder Tendon Injury	Consumer Health Professional	Neurontin (Gabapentin) Di-Gesic	PS C		ORAL

Date:08/19/02ISR Number: 3964691-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200576
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG	Duration Blood Cholesterol Increased	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL

(DAILY), ORAL

Condition Aggravated

Hip Arthroplasty
Pain
Paraesthesia
Vision Blurred

Tamoxifen

C

Date:08/19/02ISR Number: 3964711-4Report Type:Expedited (15-DaCompany Report #2002050059

Age:72 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1600 MG (800 Other MG, BID)	Dehydration Diarrhoea Infectious Renal Failure Acute	Foreign Health Professional Company Representative	Neurontin (Gabapentin) (Unspecified Antihypertensive Drug) (Unspecified Antidiabetic Drug)	PS C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/19/02ISR Number: 3965005-3Report Type:Expedited (15-DaCompany Report #2002052933

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypersomnia	Consumer	Neurontin			
800 MG		Surgery		(Gabapentin)	PS		
(DAILY)		Vision Blurred					
		Weight Increased					

Date:08/20/02ISR Number: 3964279-2Report Type:Direct Company Report #CTU 174567

Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Asthenia		Gabepentin-Neurontin			
300MG 3 TIMES		Ear Discomfort		- 300mg Pfizer, Inc	PS	Pfizer	ORAL
ORAL		Facial Pain					
		Neck Pain					
		Pain					

Date:08/20/02ISR Number: 3965139-3Report Type:Expedited (15-DaCompany Report #2002053399

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Consumer	Neurontin			
Initial or Prolonged				(Gabapentin)	PS		

Date:08/20/02ISR Number: 3965141-1Report Type:Expedited (15-DaCompany Report #001-0981-M0200681

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bipolar I Disorder	Consumer	Atorvastatin			
Required		Bladder Disorder	Health	(Atorvastatin)	PS		ORAL
10 MG							

Intervention to	Cataract	Professional	
(DAILY), ORAL			
Prevent Permanent	Oedema Peripheral	Gabapentin	SS
Impairment/Damage	Pollakiuria	Lithium Carbonate	SS
	Renal Disorder		
	Sepsis		
	Weight Increased		

Date:08/20/02ISR Number: 3965373-2Report Type:Expedited (15-DaCompany Report #2002053440
Age:1 DY Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Caesarean Section
Initial or Prolonged	Central-Alveolar
Other	Hypoventilation
	Coagulation Disorder
	Neonatal
	Complications Of Maternal
	Exposure To Therapeutic
	Drugs
	Drug Withdrawal Syndrome
	Neonatal
	Hypoglycaemia Neonatal
	Maternal Drugs Affecting
	Foetus
	Neonatal Respiratory
	Distress Syndrome

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Somnolence Neonatal

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, TID)		Foreign Health Professional	Neurontin (Gabapentin)	PS		
500 MG(BID)			Valproate Sodium	SS		

Date:08/20/02ISR Number: 3965814-0Report Type:Expedited (15-DaCompany Report #2002053443
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG (100 MG, TID)	Other	Caesarean Section Complications Of Maternal Exposure To Therapeutic Drugs	Foreign Health Professional	PS		
500 MG (BID)		Grand Mal Convulsion		Valproate Sodium	SS		

Date:08/20/02ISR Number: 3965852-8Report Type:Expedited (15-DaCompany Report #2002053401
 Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	600 MG (300 MG, BID)	Coordination Abnormal	Foreign	Gabapentin	PS		
400 MG, DAILY		Drug Toxicity	Literature				
		Dysarthria	Health	Phenytoin	SS		
		Malaise Vertigo	Professional				

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG, TID, ORAL		C-Reactive Protein Increased	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Fall Interstitial Lung Disease Pyrexia Radius Fracture Spinal Compression Fracture Tuberculosis Vertigo		Tegretol (Carbamazepine)	C		

Age:91 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG (DAILY), ORAL		Akinesia Transient Ischaemic Attack	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Tamoxifen Dyazide Ibuprofen	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/21/02ISR Number: 3966261-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200747

Age:50 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG ORAL	Alopecia Chordoma	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
	Condition Aggravated Drug Ineffective	Professional	Steroids Folate Sodium Multivitamins Senna Fruit Nystatin Morphine Sulfate Diazepam Docusate Sodium	C C C C C C C C		

Date:08/21/02ISR Number: 3966263-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200569

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Blood Prolactin Increased Renal Failure Chronic	Health Professional	Neurontin (Gabapentin) Metoclopramide	PS SS		

Date:08/21/02ISR Number: 3966269-2Report Type:Expedited (15-DaCompany Report #2002053218

Age:32 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG (TID), ORAL	Hypersensitivity Panic Attack	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Clonazepam Buopropion Hydrochloride Estrogens Conjugated	C C C C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaphylactic Reaction	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Professional Company Representative				

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (100		Hirsutism					
MG, THREE		Medication Error					
TIMES DAILY),		Restless Legs Syndrome					
ORAL				Digoxin	C		
				Lisinopril	C		
				Atenolol	C		
				Pravastatin Sodium	C		

Freedom Of Information (FOI) Report

Rofecoxib C
 Hydromorphone
 Hydrochloride C

Date:08/22/02ISR Number: 3966776-2Report Type:Expedited (15-DaCompany Report #EMADSS2001007696
 Age:77 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	25 MCG/H, 1	Depressed Level Of Consciousness	Foreign Health	Durogesic (25 Mcg/Hr Patch) (Fentanyl)	PS		
TRANSDERMAL Required IN 3 DAY(S), Intervention to TRANSD		Overdose	Professional				
Prevent Permanent Impairment/Damage		Pneumonia Aspiration		Neurontin (Gabapentin)	SS		
		Somnolence		Celebrex (Celecoxib)	C		

Date:08/22/02ISR Number: 3966855-XReport Type:Expedited (15-DaCompany Report #2002053404
 Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG (300	Alanine Aminotransferase Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other MG, TID), ORAL		Aspartate	Professional				
(TID), ORAL		Aminotransferase		Oxybutynin	SS		ORAL
INTRAVENTOUS	6 GRAM (2	Blood Creatinine		Propacetamol	SS		
GRAM, TID), INTRAVENTOUS		Increased					
10 MG (5 MG,		Blood Urea Increased		Glibenclamide	SS		ORAL
		Coagulation Factor V					

BID), ORAL	Level Decreased			
10 MG	Coagulation Factor Vii	Acepromazine	SS	ORAL
(DAILY), ORAL	Level Decreased			
	Dyspnoea	Alprazolam	C	
	Echocardiogram Abnormal	Heparin-Fraction,		
	Haemodialysis	Calcium Salt	C	
	Left Ventricular Failure	Diosmin	C	
	Prothrombin Time	Mirtazapine	C	
	Prolonged	Candesartan		
	Pulmonary Embolism	Cilexetil	C	
	Somnolence	Hydrochlorothiazide	C	
		Tramadol	C	
		Morphin Chlorhydrate	C	

Date:08/23/02ISR Number: 3966220-5Report Type:Direct Company Report #CTU 174873
Age:80 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Confusional State		Gabapentin 600mg	PS		ORAL
600MG PO EVE						
Initial or Prolonged	Difficulty In Walking					
3/5 ,600MG PO						
	Disorientation					
AM 3/6-> 300						
	Renal Impairment					
MG ON 3/7						
	Somnolence		Tylenol	C		
			Allopurinol	C		
			Albuterol	C		
			Atrovent	C		
			Percocet	C		
			Fosinopril	C		
			Warfarin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Clindamycin	C
Senna	C
Colace	C
Nph	C
Reg Insulin	C
Digoxin	C
Atenolol	C
Aciphex	C
Aspirin	C
Simvastatin	C
Nitro Patch	C
Glyburide	C

Date:08/23/02ISR Number: 3966865-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200569
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Blood Prolactin Increased	Health	Neurontin			
Initial or Prolonged	Renal Failure Chronic	Professional	(Gabapentin)	PS		
			Metoclopramide	SS		

Date:08/23/02ISR Number: 3966898-6Report Type:Expedited (15-DaCompany Report #2002053412
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Convulsion	Consumer	Neurontin			
			(Gabapentin)	PS		
			Levetiracetam	C		

Date:08/23/02ISR Number: 3966923-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200754
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Dizziness	Consumer	Neurontin			
	Memory Impairment	Health	(Gabapentin)	PS		ORAL
ORAL	Somnolence	Professional	Di-Gesic	C		

Date:08/23/02ISR Number: 3967498-4Report Type:Expedited (15-DaCompany Report #2002053410

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence	Health Professional	Neurontin (Gabapentin)	PS		

Date:08/26/02ISR Number: 3966665-3Report Type:Expedited (15-DaCompany Report #2002003836

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiac Failure Congestive	Consumer	Sertraline (Sertraline)	PS		
Other		Colonic Polyp Crying		Gabapentin (Gabapentin)	SS		ORAL
900 MG (BID),		Dysuria					
ORAL		Haemoglobin Decreased Haemorrhage Thrombosis					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/26/02ISR Number: 3966674-4Report Type:Expedited (15-DaCompany Report #2002050553

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Toxic Epidermal Necrolysis	Health Professional	Neurontin (Gabapentin)	PS		

Date:08/26/02ISR Number: 3966675-6Report Type:Expedited (15-DaCompany Report #2002053982

Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Blood Creatine Phosphokinase Increased	Literature Health	Gabapentin (Gabapentin)	PS		
Hospitalization - 2700 MG							
Initial or Prolonged (TID),		Coma	Professional				
		Electroencephalogram Abnormal		Estrogens Conjugated	C		
		Status Epilepticus		Furosemide	C		
				Levothyroxine	C		
				Lisinopril	C		
				Fluoxetine	C		

Date:08/26/02ISR Number: 3966859-7Report Type:Expedited (15-DaCompany Report #2002053567

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache Hypersomnia	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (BID),							
ORAL		Nasal Septum Deviation		Topirmate	SS		ORAL
100 MG (QID),							
ORAL				Estrogens Conjugated	C		
				Bupropion Hydrochloride	C		

Date:08/26/02ISR Number: 3966885-8Report Type:Expedited (15-DaCompany Report #2002054129
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Congenital Anomaly Maternal Drugs Affecting Foetus	Literature Health Professional	Gabapentin (Gabapentin)	PS		

Date:08/26/02ISR Number: 3966886-XReport Type:Expedited (15-DaCompany Report #2002054132
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Congenital Anomaly Maternal Drugs Affecting Foetus	Literature Health Professional	Gabapentin (Gabapentin)	PS		

Date:08/26/02ISR Number: 3966887-1Report Type:Expedited (15-DaCompany Report #2002054135
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Congenital Anomaly Maternal Drugs Affecting Foetus	Literature Health Professional	Gabapentin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/26/02ISR Number: 3966892-5Report Type:Expedited (15-DaCompany Report #2002054130

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complications Of Maternal Exposure To Therapeutic Drugs Eclampsia	Literature Health Professional	Gabapentin (Gabapentin)	PS		

Date:08/26/02ISR Number: 3967463-7Report Type:Expedited (15-DaCompany Report #2002053800

Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged ORAL		Leukocytoclastic Vasculitis	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Aceprometazine	C		
				Allopurinol	C		
				Paroxetine Hydrochloride	C		
				Diacerein	C		

Date:08/26/02ISR Number: 3967742-3Report Type:Expedited (15-DaCompany Report #PHNU2001DE02643

Age:76 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Schamberg'S Disease	Foreign Health Professional	Trileptal Fmi (Oxcarbazepine) Film-Coated Tablet	PS		ORAL
300 MG, TID,			Other				
ORAL				Neurontin (Gabapentin) Capsule	SS		ORAL
1600 MG/DAY,							
ORAL				Nitrates (Nitrates)	C		

Date:08/26/02ISR Number: 3967784-8Report Type:Expedited (15-DaCompany Report #2002052184
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Surgery	Consumer	Neurontin (Gabapentin)	PS		

Date:08/26/02ISR Number: 3967850-7Report Type:Expedited (15-DaCompany Report #2002051916
Age: Gender:Female I/FU:F

Outcome	PT
Other	Abdominal Pain
	Anxiety
	Arthritis
	Back Pain
	Blood Cholesterol
	Increased
	Blood Triglycerides
	Increased
	Constipation
	Diarrhoea
	Dizziness
	Dysphagia
	Fatigue

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG	(DAILY), ORAL	Fibrocystic Breast Disease Intervertebral Disc Protrusion Mood Altered	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
10 MG,	(DAILY)	Myalgia Nervousness Red Blood Cell Count Decreased	Professional	Lipitor (Atorvastatin)	SS		
12 MG (4 MG,	THREE TIMES DAILY)	Respiratory Distress Sedation Somnolence Thinking Abnormal Thyroxine Free Decreased Tinnitus Total Cholesterol/Hdl Ratio Decreased		Valdecoxib Nortriptyline Hydrochloride Cyproheptadine Hydrochloride Applesauce Acetylsalicylic Acid Alprazolam Tramadol Hydrochloride Estrogenbs Conjugated Paracetamol	SS SS SS SS C C C C C C C C		

Date:08/26/02ISR Number: 3967946-XReport Type:Expedited (15-DaCompany Report #2002053678
Age:78 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Life-Threatening Arthralgia Dyspnoea Lymphoedema Somnolence Weight Increased	Foreign Health Professional	Neurontin (Gabapentin) Calcium Carbonate Isosorbide Paramol-118 Warfarin	PS C C C C		ORAL

Digoxin	C
Bumetanide	C
Bezafibrate	C
Dorzolamide	
Hydrochloride	C

Date:08/26/02ISR Number: 3968178-1Report Type:Expedited (15-DaCompany Report #2002054136
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Unevaluable Event	Literature Health Professional	Gabapentin (Gabapentin)	PS		

Date:08/26/02ISR Number: 3968180-XReport Type:Expedited (15-DaCompany Report #2002054137
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complications Of Maternal Exposure To Therapeutic Drugs	Literature Health Professional	Gabapentin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/26/02ISR Number: 3968182-3Report Type:Expedited (15-DaCompany Report #2002054139
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Unevaluable Event	Literature Health Professional	Gabapentin (Gabapentin)	PS		

Date:08/26/02ISR Number: 3968184-7Report Type:Expedited (15-DaCompany Report #2002054140
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complications Of Maternal Exposure To Therapeutic Drugs Convulsion Eclampsia	Literature Health Professional	Gabapentin (Gabapentin)	PS		

Date:08/27/02ISR Number: 3967614-4Report Type:Direct Company Report #CTU 175197
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening	4 DAY	Blindness		Neurontin	PS		
		Condition Aggravated Fatigue Headache Localised Infection Neuropathy Peripheral					

Date:08/27/02ISR Number: 3968248-8Report Type:Expedited (15-DaCompany Report #2002053360
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - 1800 MG,		Blood Chloride Decreased Blood Sodium Decreased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL

Initial or Prolonged Dehydration
ORAL; 900

Professional

MG, ORAL

Company

Representative

Date:08/27/02ISR Number: 3968382-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200718

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain	Consumer	Neurontin			
Other		Fluid Retention	Health	(Gabapentin)	PS		ORAL
600 MG (BID),		Hypoaesthesia	Professional				
ORAL		Nausea		Celecoxib	SS		ORAL
200 MG		Oedema Peripheral					
(DAILY), ORAL		Pharmaceutical Product					
		Complaint					
		Thrombosis					
		Vein Disorder					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/27/02ISR Number: 3968384-6Report Type:Expedited (15-DaCompany Report #001-0945-M0200633

Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		No Adverse Drug Effect		Neurontin (Gabapentin)	PS		ORAL
900 MG (300							
MG, THREE							
TIMES DAILY),							
ORAL							

Celecoxib SS

Date:08/27/02ISR Number: 3969164-8Report Type:Expedited (15-DaCompany Report #2002053814

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Retinal Haemorrhage	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:08/27/02ISR Number: 3969177-6Report Type:Expedited (15-DaCompany Report #02P-163-0188795-00

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Health	Tarka Er 4 Mg/240 Mg			
Hospitalization -		Family Stress	Professional	(Tarka Er)	PS		ORAL
MG, 2 IN 1 D,							
Initial or Prolonged		Hypotension	Other				
PER ORAL							

Intentional Misuse Hydrochlorothiazide SS

SEE IMAGE

Isoptin Sr
(Verapamil)
(Verapamil) SS

180 MG, 2 IN

ORAL

1 D, PER ORAL

Metformin	
Hydrochloride	SS
Citalopram	
Hydrobromide	SS
Gabapentin	SS
Clopidogrel Sulfate	SS
Isosorbide Dinitrate	SS
Insulin	C
Glucophage	C
Nitrate	C
Acetylsalicylic Acid	C
Citalopram	
Hydrobromide	C
Fluoxetine	C
Omeprazole	C

Date:08/27/02ISR Number: 3969309-XReport Type:Expedited (15-DaCompany Report #2002053815
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Joint Stiffness Joint Swelling	Consumer	Neurontin (Gabapentin)	PS		ORAL
2700 MG (900 MG, THREE TIMES DAILY), ORAL		Swelling					

Alprazolam	C
Bupivacaine	C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydromorphone
Hydrochloride C
Clonidine C

Date:08/28/02ISR Number: 3969510-5Report Type:Expedited (15-DaCompany Report #02P-056-0198582-00
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to Prevent Permanent Impairment/Damage	500 MG, 2 IN	Complications Of Maternal Exposure To Therapeutic Drugs	Foreign	Depakine Tablets (Sodium Valproate)	PS		ORAL
	1 D, ORAL	Epilepsy					
	2 MG, 2 IN 1	Pregnancy		Clonazepam	SS		ORAL
	D, ORAL						
	600 MG, 3 IN			Gabapentin	SS		ORAL
	1 D, ORAL						
				Folic Acid	C		

Date:08/29/02ISR Number: 3969297-6Report Type:Expedited (15-DaCompany Report #2002053802
Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	900 MG (TWICE DAILY), ORAL	Balance Disorder Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Constipation					
	ORAL	Diarrhoea		Hydrocodone	SS		ORAL
		Spinal Fracture		Risedronate Sodium	C		
		Weight Decreased		Tamoxifen	C		
				Calcium With Vitamin D	C		
				Phazyme	C		

Date:08/30/02ISR Number: 3969420-3Report Type:Expedited (15-DaCompany Report #2002051969
 Age:16 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (BID), Other ORAL		Blood Creatine Phosphokinase Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
0.5 MG (BID), ORAL				Risperidone	SS		ORAL
				Topiramate	C		
				Salbutamol	C		
				Pseudoephedrine Hydrochloride	C		
				Paracetamol	C		
				Topiramate	C		

Date:08/30/02ISR Number: 3969679-2Report Type:Direct Company Report #CTU 175360
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1 CAP TID PRN ORAL		Vertigo Vision Blurred Visual Disturbance		Neurontin 200 Mg Parke-Davis	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/30/02ISR Number: 3970277-5Report Type:Expedited (15-DaCompany Report #2002054154
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG		Amnesia Urinary Incontinence Vision Blurred	Foreign Consumer	Neurontin (Gabapentin)	PS		
(THREE TIMES A DAY)							

Date:08/30/02ISR Number: 3970545-7Report Type:Expedited (15-DaCompany Report #2002054058
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG DAILY, ORAL		Blood Oestrogen Increased Convulsion Dizziness Progesterone Decreased	Consumer	Neurontin (Gabapentin) Primidone Acetazolamide	PS C C		ORAL

Date:08/30/02ISR Number: 3970547-0Report Type:Expedited (15-DaCompany Report #2002054126
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG, ORAL		Condition Aggravated Difficulty In Walking Drug Ineffective Neuropathy Peripheral	Consumer	Neurontin (Gabapentin) Metformin Hydrochloride Glipizide Potassium Chloride Penicillin Nos Doxazosin Mesilate Fosinopril Sodium Furosemide Acetylsalicylic Acid	PS C C C C C C C C		ORAL

Timolol Maleate C
Atorvastatin C

Date:08/30/02ISR Number: 3970574-3Report Type:Expedited (15-DaCompany Report #2002050553
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis	Health	Neurontin			
		Lice Infestation	Professional	(Gabapentin)	PS		
		Skin Necrosis					

Date:09/03/02ISR Number: 3969394-5Report Type:Expedited (15-DaCompany Report #320087
Age: Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Activated Partial
Initial or Prolonged	Thromboplastin Time
	Abnormal
	Caesarean Section
	Coagulation Disorder
	Neonatal
	Drug Withdrawal Syndrome
	Neonatal
	Feeding Problem In

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Newborn Hypoglycaemia Neonatal Maternal Drugs Affecting Foetus	Consumer	Rivotril Neurontin Depakine	PS SS SS	Roche	
		Neonatal Disorder Neonatal Respiratory Distress Syndrome Prothrombin Time Prolonged Somnolence Neonatal					

Date:09/03/02ISR Number: 3971151-0Report Type:Expedited (15-DaCompany Report #02P-056-0198581-00
Age:1 DY Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Caesarean Section Coagulation Disorder Neonatal Complications Of Maternal Exposure To Therapeutic Drugs Drug Withdrawal Syndrome Neonatal Feeding Problem In Newborn Hypoglycaemia Neonatal Maternal Drugs Affecting Foetus Neonatal Respiratory Distress Syndrome Somnolence Neonatal	Foreign Health Professional	Depakine Tablets (Sodium Valproate) (Sodium Valproate) (Sodium Valproate) Clonazepam Gabapentin Folic Acid	PS SS SS C		

Date:09/04/02ISR Number: 3969979-6Report Type:Expedited (15-DaCompany Report #320345
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 8 DAY Initial or Prolonged 8 DAY		Leukopenia Overdose		Rivotril Deroxat	PS SS	Roche	

DAILY. 8 DAY Renal Failure Acute
 8 DAY Thrombocytopenia
 8 DAY
 8 DAY
 8 DAY

Methotrexate SS
 Lioresal SS
 Neurontin SS
 Topalgic SS
 Xanax C

Date:09/04/02ISR Number: 3970502-0Report Type:Direct Company Report #CTU 175621
 Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 5MG QHS		Fall		Terazosin	PS		
Initial or Prolonged 25MG QD		Hypotension		Atenolol 25mg Tab	SS		
Required 400MG BI				Gabapentin 400mg Cap	SS		
Intervention to Prevent Permanent Impairment/Damage				Rabeprazole	C		
				Citalopram	C		
				Gemfibrozil	C		
				Aspirin Enteric Coated	C		
				Multivitamin	C		
				Ipratropium Inh	C		
				Acetaminophen	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nitroglycerin C

Date:09/04/02ISR Number: 3970904-2Report Type:Expedited (15-DaCompany Report #2002053399
Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), Other ORAL		Convulsion Disturbance In Attention Movement Disorder Panic Attack Speech Disorder	Consumer	Neurontin (Gabapentin) Amitriptyline Hydrochloride	PS C		ORAL

Date:09/04/02ISR Number: 3970909-1Report Type:Expedited (15-DaCompany Report #2002054194
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1200 MG, ORAL		Breech Presentation Caesarean Section Complications Of Maternal Exposure To Therapeutic Drugs Drug Withdrawal Syndrome Maternal Drugs Affecting Foetus Petit Mal Epilepsy Postpartum Depression Unwanted Pregnancy	Consumer	Neurontin (Gabapentin) Dexamfetamine Sulfate Lorazepam Mirtazapine Thioridazine Hydrochloride Clonazepam	PS C C C C C		ORAL

Date:09/04/02ISR Number: 3970910-8Report Type:Expedited (15-DaCompany Report #2002054191
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Abortion Spontaneous Complications Of Maternal	Consumer	Neurontin (Gabapentin)	PS		ORAL

Exposure To Therapeutic
Drugs
Feeling Abnormal
Hallucination
Maternal Drugs Affecting
Foetus
Mood Swings
Postpartum Depression
Psychotic Disorder
Unwanted Pregnancy
Weight Increased

Dexamfetamine
Sulfate C
Lorazepam C
Mirtazapine C
Thioridazine
Hydrochloride C
Clonazepam C

Date:09/04/02ISR Number: 3970912-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200718

Age: Gender:Male I/FU:F

Outcome PT
Other Back Pain
Condition Aggravated
Deep Vein Thrombosis
Fluid Retention
Hypoaesthesia
Nausea
Pharmaceutical Product

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Complaint
Varicose Vein

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
600 MG (BID), ORAL		Consumer Health	Neurontin (Gabapentin)	PS		ORAL
200 MG (DAILY), ORAL		Professional	Celecoxib	SS		ORAL

Date:09/04/02ISR Number: 3971443-5Report Type:Expedited (15-DaCompany Report #2002050043
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other (TWICE DAILY), ORAL		Abdominal Distension Asthenia	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Blood Heavy Metal Increased		Lamotrigine	SS		ORAL
		Dyspnoea Feeling Abnormal Helicobacter Infection Hunger Nausea Oral Intake Reduced Weight Increased		Clonazepam	SS		

Date:09/05/02ISR Number: 3971315-6Report Type:Direct Company Report #CTU 175731
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability INTRAMUSCULAR Required A DAY	25/100MG 3X	Extensor Plantar Response Parkinson'S Disease		Sinemet 25/100	PS		

Intervention to
INTRAMUSCULAR
Prevent Permanent
Impairment/Damage
INTRAMUSCULAR 300MG 3X A

Gabapentin 300mg
Parke/Davis SS Parke/Davis

DAY

INTRAMUSCULAR

Date:09/05/02ISR Number: 3972432-7Report Type:Expedited (15-DaCompany Report #2002054310
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Bleeding Time Prolonged Haemorrhagic Diathesis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Risedronate Sodium	C		
Other				Cetirizine Hydrochloride	C		

Date:09/05/02ISR Number: 3972434-0Report Type:Expedited (15-DaCompany Report #2002054675
Age:35 YR Gender:Male I/FU:I

Outcome	PT
Other	Chills Drug Level Below Therapeutic Gait Disturbance Hemiplegia Migraine

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (TID), ORAL		Muscle Twitching Paranoia	Consumer	Neurontin(Gabapentin)	PS		ORAL
				Citalopram Hydrobromide	C		

Date:09/05/02ISR Number: 3972780-0Report Type:Expedited (15-DaCompany Report #2002050928
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other (QID),ORAL		Decreased Activity Dental Caries	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Hypothyroidism	Professional	Venlafaxine Hydrochloride	C		
		Migraine		Paroxetine Hydrochloride	C		
		Oral Intake Reduced		Tramadol	C		
		Poisoning		Clonazepam	C		
		Pyrexia		Carisoprodol	C		
		Tooth Disorder					
		Weight Increased					

Date:09/05/02ISR Number: 3972782-4Report Type:Expedited (15-DaCompany Report #2002054674
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Nausea Vomiting	Consumer	Neurontin(Gabapentin)	PS		ORAL
				Simvastatin	C		
				Amlodipine Besilate	C		
				Gemfibroxil	C		
				Estrogens Conjugated	C		
				Prinzide	C		
				Paroxetine Hydrochloride	C		

Date:09/05/02ISR Number: 3973247-6Report Type:Expedited (15-DaCompany Report #320087

Age: Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Central-Alveolar
Initial or Prolonged	Hypoventilation
	Coagulation Disorder
	Neonatal
	Complications Of Maternal
	Exposure To Therapeutic
	Drugs
	Drug Withdrawal Syndrome
	Neonatal
	Feeding Problem In
	Newborn
	Grand Mal Convulsion
	Hypoglycaemia Neonatal
	Maternal Drugs Affecting
	Foetus
	Neonatal Respiratory
	Distress Syndrome

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pregnancy Somnolence Neonatal	Report Source	Product	Role	Manufacturer	Route
2 MG 2 PER			Foreign Other	Rivotril (Clonazepam)	PS		
DAY				Neurontin (Gabapentin) 100 Mg	SS		
600 MG 3 PER				Depakine (Valproate Sodium) 500 Mg	SS		
DAY							
500 MG 2 PER							
DAY							

Date:09/06/02ISR Number: 3971404-6Report Type:Expedited (15-DaCompany Report #320345
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 8 DAY		Leukopenia	Consumer	Rivotril	PS	Roche	
Initial or Prolonged 8 DAY		Overdose		Deroxat	SS		
DAILY. 8 DAY	8 DAY	Renal Failure Acute		Methotrexate	SS		
8 DAY		Thrombocytopenia		Lioresal	SS		
8 DAY				Neurontin	SS		
8 DAY				Topalgic (Tramadol)	SS		
8 DAY				Xanax	C		

Date:09/06/02ISR Number: 3972500-XReport Type:Expedited (15-DaCompany Report #B0263843A
Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG /	Hepatitis Cholestatic Pancreatitis	Foreign	Combivir Tablet (Combivir)	PS		ORAL
	TWICE PER DAY	Rash Maculo-Papular					
	/ ORAL						
	400 MG /			Kaletra Capsule (Kaletra)	SS		ORAL
	TWICE PER DAY						
	/ ORAL						
	500 MG /			Gabapentin (Formulation Unknown) (Gabapentin)	SS		ORAL
	THREE TIMES						
	PER DAY /						
	ORAL						
				Pyrimethamine	C		
				Sulphadiazine	C		
				Calcium Folate	C		
				Clobazam	C		

Date:09/09/02ISR Number: 3973175-6Report Type:Direct
Age:23 YR Gender:Male I/FU:I

Company Report #CTU 175911

Outcome	PT
Life-Threatening Disability	Abdominal Pain Anxiety Condition Aggravated Decreased Activity Depression Suicidal

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	General Physical Health Deterioration Impaired Work Ability	Report Source	Product	Role	Manufacturer	Route
300 MG 2 DAILY		Libido Decreased Panic Attack Social Avoidant Behaviour		Neurontin 300 Mg	PS		

Date:09/09/02ISR Number: 3973236-1Report Type:Direct Company Report #USP 55036
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Medication Error		Neurontin (Gabapentin)	PS	Pfizer	
				Noroxin (Norfloxacin)	SS	Merck	

Date:09/09/02ISR Number: 3973660-7Report Type:Expedited (15-DaCompany Report #2002054696
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 800 MG DAILY, ORAL		Cellulitis Convulsion Drug Level Above Therapeutic	Consumer	Neurontin (Gabapentin)	PS		ORAL
400 MG DAILY, ORAL		Liver Function Test Abnormal		Phenytoin Sodium	SS		ORAL
25 MG DAILY		Staphylococcal Infection		Lamotrigine	SS		
				Paracetamol	SS		
				Venlafaxine Hydrochloride	C		
				Atorvastatin	C		
				Benazepril Hydrochloride	C		
				Clonazepam	C		

Obetrol	C
Quetiapine Fumarate	C
Mirapex	C
Eptacog Alfa	C
Folic Acid	C
Pyridoxine	
Hydrochloride	C
Cyanocobalamin	C

Date:09/09/02ISR Number: 3973662-0Report Type:Expedited (15-DaCompany Report #2002054897
 Age:90 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Agitation Blood Pressure Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other 1000 MG QID, ORAL	Clostridial Infection Confusional State		Metronidazole	SS		ORAL
	Drug Interaction Infection Leg Amputation Pain Paranoia Somnolence		Glibenclamide Chromagen Prednisone	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/09/02ISR Number: 3973700-5Report Type:Expedited (15-DaCompany Report #001-0719-M0100408

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 1200 MG Initial or Prolonged (BID), ORAL Other	Arthralgia	Consumer	Lopid (Gemfibrozil)	PS		ORAL
ORAL	Asthenia	Health				
5 MG (DAILY), ORAL	Back Pain Difficulty In Walking	Professional	Gabapentin (Gabapentin)	SS		ORAL
	Disorientation Dizziness		Amlodipine (Amlodipine)	SS		ORAL
	Headache					
5 MG (DAILY), ORAL	Hypertension Myalgia		Glipizide (Glipizide)	SS		ORAL
			Nifedipine (Nifedipine)	SS		
10 MG, SUBLINGUAL			(Unspecified Choleserol Lowering Drug)	SS		
			Furosemide	C		
			Acetylsalicylic Acid	C		
			Multivitamins	C		
			Terazosin	C		
			Labetalol Hydrochloride	C		
			(Sunastatin)	C		
			Clopidogrel	C		
			Meloxicam (Pericad Plus Vitamins)	C		
			(L-Carnitine)	C		
			(Glucosamine Chondroitin)	C		
			(Msm)	C		
			Celecoxib	C		
			Tamsulosin			

Hydrochloride	C
Tolterodine	
L-Tartrate	C
Lotrel	C
Metoprolol Succinate	C
Simvastatin	C
Vicodin	C
Crataegus Extract	C
Ubidecarenone	C
Calcium	C
Garlic	C
(Unspecified Pain Medication)	C
Mibefradil Hydrochloride	C

Date:09/09/02ISR Number: 3973993-4Report Type:Expedited (15-DaCompany Report #2002054934
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Health	Neurontin			
		Anger	Professional	(Gabapentin)	PS		ORAL
900 MG (TID),		Anxiety					
ORAL		Tremor		Lithium	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/09/02ISR Number: 3974006-0Report Type:Expedited (15-DaCompany Report #2002052658

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Atrial Fibrillation	Health	Neurontin			
Other		Atrial Flutter	Professional	(Gabapentin)	PS		
400 MG (BID)		Blood Pressure Decreased		Carbamazepine	SS		
		Cardiomyopathy		Meclozine	SS		
		Cardiovascular Disorder					
		Coordination Abnormal					
		Coronary Artery Disease					
		Diabetes Mellitus					
		Insulin-Dependent					
		Drug Interaction					
		Intermittent Claudication					
		Medication Error					
		Myocardial Ischaemia					
		Neuropathy Peripheral					
		Overdose					
		Renal Failure Chronic					
		Somnolence					
		Syncope					
		Tachyarrhythmia					
		Ventricular Hypertrophy					

Date:09/09/02ISR Number: 3974008-4Report Type:Expedited (15-DaCompany Report #2002054820

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Neurontin			
Disability		Blood Cholesterol		(Gabapentin)	PS		ORAL
Other		Increased					
900 MG (300		Depression					
MG, TID),		Drug Tolerance Increased					
ORAL		Eye Pain					
		Feeling Abnormal					
		Feeling Drunk					
		Hypoaesthesia					
		Ill-Defined Disorder					

Lethargy
Migraine
Palpitations
Panic Attack

Date:09/09/02ISR Number: 3974055-2Report Type:Expedited (15-DaCompany Report #2002054890
Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Bronchial Obstruction Cough	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Hypoxia	Professional	Fentanyl	SS		ORAL
		Overdose Pyrexia Somnolence		Celecoxib	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/09/02ISR Number: 3974070-9Report Type:Expedited (15-DaCompany Report #2002051362
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG		Eosinophilia Hypersensitivity	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other (TID), ORAL		Renal Failure Acute Thrombocytopenia	Professional				

Date:09/09/02ISR Number: 3974071-0Report Type:Expedited (15-DaCompany Report #2002055082
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG, ORAL		Narcolepsy Syncope	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
			Professional Company Representative	Serotonin Selective Reuptake Inhibitor (Unspecified) (Serotonin) Narcotic (Unspecified) (All Other Therapeutic Products)	C C		

Date:09/09/02ISR Number: 3974072-2Report Type:Expedited (15-DaCompany Report #2002055083
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Blood Disorder Neoplasm Malignant Skin Cancer	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:09/09/02ISR Number: 3974095-3Report Type:Expedited (15-DaCompany Report #2002054543
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness Nausea	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
400 MG, ORAL		Speech Disorder Thinking Abnormal Vomiting	Professional				

Date:09/09/02ISR Number: 3974097-7Report Type:Expedited (15-DaCompany Report #2002054786
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged ORAL		Clostridium Colitis Pyrexia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Professional	Anethole Trithione	SS		ORAL
ORAL				Ascorbic Acid	SS		ORAL
				Metronidazole Vancomycin	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/11/02ISR Number: 3974487-2Report Type:Expedited (15-DaCompany Report #2002055226
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Rash Maculo-Papular Renal Failure Acute Thrombocytopenia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:09/11/02ISR Number: 3974494-XReport Type:Expedited (15-DaCompany Report #2002055134
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Feeling Hot Headache	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:09/11/02ISR Number: 3974692-5Report Type:Expedited (15-DaCompany Report #2002053218
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG (TID), ORAL		Hypersensitivity Panic Attack Pharyngeal Oedema	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Clonazepam Bupropion Hydrochloride Estrogens Conjugated Sertraline Hydrochloride	C C C C C		

Date:09/12/02ISR Number: 3975144-9Report Type:Direct Company Report #CTU 176292
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required DAYPRO		Eye Swelling		Neurontin 800	PS		
Intervention to Prevent Permanent Impairment/Damage		Rash Erythematous Rash Maculo-Papular Rash Pruritic		Bisoprolol/Hctz 5/6.25	SS		

Date:09/12/02ISR Number: 3975242-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200328
Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Other	Alopecia Asthma Blood Potassium Decreased Condition Aggravated Haematoma Increased Appetite Muscle Spasms Nausea Petechiae Stevens-Johnson Syndrome Stupor Thrombocytopenia Trigeminal Neuralgia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL	200 MG	Vomiting Weight Increased	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
	(DAILY), ORAL		Professional	Carbamazepine	SS		ORAL
				Levothyroxine Sodium	C		

Date:09/12/02ISR Number: 3975328-XReport Type:Expedited (15-DaCompany Report #2002055494
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	Hospitalization - 300 MG	Arteriosclerosis Cardiac Failure	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Initial or Prolonged (DAILY), ORAL		Cardiomegaly	Professional				
Other		Drug Ineffective		Zopiclone	C		
		Fatigue		Ciprofloxacin	C		
		Nephrosclerosis		Fluconazole	C		
		Oedema		Paracetamol	C		
		Oedema Peripheral		Citalopram			
		Pitting Oedema		Hydrobromide	C		
		Pleural Effusion		B-Kombin Tablet	C		
		Pulmonary Oedema		Flucloxacillin			
		Scar		Sodium	C		
		Thrombosis		Oxazepam	C		
		Weight Increased		Furosemide	C		
				Insulin	C		

Date:09/12/02ISR Number: 3975413-2Report Type:Expedited (15-DaCompany Report #2002-04821
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300 MG QD PO	Pneumonia	Foreign	Viread	PS		ORAL

Proteinuria

Health
Professional
Company
Representative
Other

Abacavir
Didanosine
Kaletra
Venlafaxine
Diazepam
Domperidone
Centrum
Gabapentin
Temazepam
Mst

C
C
C
C
C
C
I
I
I

Date:09/12/02ISR Number: 3975460-0Report Type:Expedited (15-DaCompany Report #2002056050
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cold Sweat Difficulty In Walking	Consumer	Neurontin (Gabapentin)	PS		ORAL
3600 MG (1200 MG, THREE TIMES DAILY), ORAL		Insomnia Movement Disorder Pain		Rofecoxib Enalapril Fluoxetine Hydrochloride	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/12/02ISR Number: 3975463-6Report Type:Expedited (15-DaCompany Report #2002056051

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG (100 MG, BID), ORAL ORAL		Condition Aggravated Dizziness Drug Ineffective Drug Level Increased Fatigue Grand Mal Convulsion Simple Partial Seizures	Consumer	Dilantin Suspension (Phenytoin Sodium) Gabapentin Lamotrigine Levetiracetam	PS SS SS SS		ORAL ORAL

Date:09/12/02ISR Number: 3975543-5Report Type:Expedited (15-DaCompany Report #PROG00202002271

Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged DAILY PO INTRAMUSCULAR DAILY IM DAILY	30 MCG	Anomaly Ovarian Cyst Pancreatitis Acute	Health Professional	Utrogestan (Progesterone) Avonex (Interferon Beta) Gabapentin (Gabapentin) Provigil (Modafinil)	PS SS SS I		ORAL

Date:09/13/02ISR Number: 3975850-6Report Type:Expedited (15-DaCompany Report #2002055658

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hepatic Enzyme Increased Leukocytosis Pyrexia	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Company
Representative

Date:09/13/02ISR Number: 3976022-1Report Type:Expedited (15-DaCompany Report #2002055716

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Eosinophilia	Health	Neurontin			
Hospitalization - Initial or Prolonged		Rash	Professional Company Representative	(Gabapentin)	PS		

Date:09/16/02ISR Number: 3976384-5Report Type:Expedited (15-DaCompany Report #001-0719-M0100408

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Other	Arthralgia Arthropathy Asthenia Back Pain Chest Pain Coordination Abnormal Disorientation Dizziness Drug Ineffective Fatigue

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Adverse Event	Report Source	Product	Role	Manufacturer	Route
1200 MG	(BID), ORAL	Headache Infection Movement Disorder	Consumer	Lopid (Gemfibrozil)	PS		ORAL
		Myalgia	Health				
		Neck Pain	Professional	Gabapentin (Gabapentin)	SS		ORAL
		Pain Palpitations		Amlodipine (Amlodipine)	SS		ORAL
5 MG (DAILY),		Speech Disorder Tenderness					
ORAL		Tinnitus					
5 MG (DAILY),		Varicose Vein Vision Blurred		Glipizide (Glipizide)	SS		ORAL
ORAL				Nifedipine (Nifedipine)	SS		
SUBLINGUAL	10 MG,						
SUBLINGUAL				(Unspecified Cholesterol Lowering Drug)	SS		
				Furosemide	C		
				Acetylsalicylic Acid	C		
				Multivitamins	C		
				Terazosin	C		
				Labetalol Hydrochloride	C		
				(Sunastatin)	C		
				Clopidogrel	C		
				Meloxicam	C		
				(Pericad Plus Vitamins)	C		
				(L-Carnitine)	C		
				(Glucosamine Chondroitin)	C		
				(Msm)	C		
				Celecoxib	C		
				Tamsulosin Hydrochloride	C		

Tolterodine	C
L-Tartrate	C
Lotrel	C
Metoprolol Succinate	C
Simvastatin	C
Vicodin	C
Crataegus Extract	C
Ubidecarenone	C
Calcium	C
Garlic	C
(Unspecified Pain Medication)	C
Mibefradil Hydrochloride)	C

Date:09/16/02ISR Number: 3976534-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200375
Age:59 YR Gender:Female I/FU:F

Outcome	PT
Other	Blood Pressure Abnormal Diverticulitis Fall Gastrooesophageal Reflux

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Disease	Report Source	Product	Role	Manufacturer	Route
		Hypoaesthesia					
		Impaired Healing					
		Lower Limb Fracture					
		Muscle Spasms	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG, TID),			Professional				
ORAL				Cimetidine	SS		
1600 MG (BID)				Omeprazole	SS		ORAL
(DAILY), ORAL				Metformin	C		
				Losartan Potassium	C		
				Atorvastatin	C		
				Diltiazem Hydrochloride	C		
				Hydroxyprogesterone	C		
				Estrogens Conjugated	C		
				Hydrochlorothiazide	C		
				Insulin Human Injection, Isophane	C		
				Glibenclamide	C		
				Acetylsalicylic Acid	C		
				Multivitamins	C		
				Ibuprofen	C		
				Potassium Chloride	C		

Date:09/16/02ISR Number: 3976535-2Report Type:Expedited (15-DaCompany Report #2002052154

Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (TID),		Dysgraphia	Professional				
ORAL				Sertraline Hydrochloride	C		
				Oxycocet	C		
				Levetiracetam	C		

Date:09/16/02ISR Number: 3976538-8Report Type:Expedited (15-DaCompany Report #2002056416
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Prostate Cancer	Consumer	Neurontin (Gabapentin)	PS		ORAL

2400 MG (600
MG, QID),

ORAL

- Gemfibrozil C
- Atorvastatin C
- Guaifenesin C
- Allerx-D C
- Cetirizine Hydrochloride C
- Tamsulosin Hydrochloride C
- Bicalutamide C
- Morphine Sulfate C
- Tramadol Hydrochloride C
- Diazepam C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Celecoxib C
 Nefazodone
 Hydrochloride C
 Budesonide C
 Mometasone Furoate C
 Ipratropium Bromide C

Date:09/16/02ISR Number: 3976540-6Report Type:Expedited (15-DaCompany Report #2002003836
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other	Cardiac Failure Congestive Colonic Haemorrhage Colonic Polyp	Consumer Health Professional	Sertraline (Sertraline) Gabapentin (Gabapentin)	PS SS		ORAL
900 MG (BID), ORAL	Crying Haemoglobin Decreased Moaning Thrombosis Urinary Retention					

Date:09/16/02ISR Number: 3976543-1Report Type:Expedited (15-DaCompany Report #2002055686
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 20 MG Disability (DAILY), ORAL Other (TWICE DAILY) ,ORAL 2000 MG (TWICE	Condition Aggravated Difficulty In Walking Drug Ineffective Insomnia Muscular Weakness Neuropathy Peripheral Staphylococcal Sepsis	Consumer	Atorvastatin (Atorvastatin) Gabapentin Glucophage	PS SS SS		ORAL ORAL ORAL

DAILY), ORAL

Glipizide	C
Potassium Chloride	C
Penicillin	C
Doxazosin Mesilate	C
Fosinopril Sodium	C
Furosemide	C
Aspirin	C
Timolol	C

Date:09/16/02ISR Number: 3976624-2Report Type:Expedited (15-DaCompany Report #EMADSS2002005362
Age:84 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 8 MG, DAILY, ORAL	Apraxia Brain Scan Abnormal Fall	Foreign Health Professional	Reminyl (Tablet) (Galantamine)	PS		ORAL
0.5 MG, DAILY, ORAL	Gait Disturbance Hypertonia Polyneuropathy		Risperdal (1 Mg Tablet) (Risperidone)	SS		ORAL
1000 MG, DAILY, ORAL			Neurontin (Gabapentin)	SS		ORAL
1 CAP, DAILY,			Colchimax (Colchimax)	SS		ORAL

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Freedom Of Information (FOI) Report

ORAL				Seropram (Citalopram Hydrobromide)	SS		ORAL
20 MG, DAILY;							
ORAL				Bi-Tildiem (Diltiazem Hydrochloride)	SS		ORAL
180 MG,							
DAILY; ORAL				Nitriderm Tts (Glyceryl Trinitrate)	C		
				Hept-A-Myl (Heptaminol Hydrochloride)	C		
				Temesta (Lorazepam)	C		
				Imovan (Zopiclone)	C		
				Dafalgan (Paracetamol)	C		
				Kardegic (Acetylsalicylate Lysine)	C		

Date:09/16/02ISR Number: 3976654-0Report Type:Expedited (15-DaCompany Report #EMADS2002005362
 Age:84 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Apraxia Areflexia Fall	Foreign Health Professional	Risperdal (1 Mg Tablet) (Risperidone)	PS		ORAL
0.5MG, DAILY, ORAL	Gait Disturbance					
8 MG, DAILY, ORAL	Hypertonia Hyporeflexia Pain In Extremity		Reminyl (Tablet) (Galantamine)	SS		ORAL
1000 MG,	Polyneuropathy		Neurontin (Gabapentin)	SS		ORAL

DAILY, ORAL	Colchimax (Colchimax)	SS	ORAL
1 CAP, DAILY, ORAL			
20 MG, DAILY, ORAL	Seropram (Citalopram Hydrobromide)	SS	ORAL
180 MG, DAILY, ORAL	Bi-Tildiem (Diltiazem Hydrochloride)	SS	ORAL
	Nitriderm Tts (Glyceryl Trinitrate)	C	
	Hept-A-Myl (Heptaminol Hydrochloride)	C	
	Temesta (Lorazepam)	C	
	Imovane (Zopiclone)	C	
	Dafalgan (Paracetamol)	C	
	Kardegic (Acetylsalicylate Lysine)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/16/02ISR Number: 3976708-9Report Type:Expedited (15-DaCompany Report #2002054786
 Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Clostridium Colitis Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Professional	Anethole Trithione	SS		ORAL
ORAL				Ascorbic Acid	SS		ORAL
ORAL				Metronidazole	C		
				Vancomycin	C		

Date:09/17/02ISR Number: 3977268-9Report Type:Expedited (15-DaCompany Report #B0278371A
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Leukopenia Overdose Renal Failure Acute	Foreign	Paxil Tablet (Paroxetine Hydrochloride)	PS		ORAL
20 MG ORAL		Thrombocytopenia		Tramadol Hydrochloride (Formulation Unknown) (Tramadol Hydrochloride)	SS		ORAL
ORAL				Gabapentin (Formulation Unknown) (Gabapentin)	SS		ORAL
ORAL				Clonazepam Injection (Clonazepam)	SS		
INTRAVENOUS	INTRAVENOUS			Baclofen Tablet (Baclofen)	SS		ORAL
ORAL				Methotrexate Tablet (Methotrexate)	SS		ORAL
ORAL				Alprazolam	C		

Date:09/17/02ISR Number: 3978555-0Report Type:Expedited (15-DaCompany Report #2002052306

Age:83 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (THREE TIMES DAILY), ORAL		Arrhythmia Asthenia Dyspnoea	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
				Amitriptyline	C		
				Ginkgo Biloba	C		
				Omeprazole	C		
				Verapamil	C		
				Spiroinolactone	C		
				Methadone			
				Hydrochloride	C		
				Meloxicam	C		
				Homeopathic Drug	C		

Date:09/17/02ISR Number: 3979894-XReport Type:Expedited (15-DaCompany Report #2002056606

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Neurotoxicity	Consumer	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/17/02ISR Number: 3979895-1Report Type:Expedited (15-DaCompany Report #2002056605

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Medication Error	Consumer	Neurontin			
Other		Neurotoxicity		(Gabapentin)	PS		

Date:09/17/02ISR Number: 3980208-XReport Type:Expedited (15-DaCompany Report #A208992

Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	100 MG	Coma	Health	Zoloft (Sertraline)	PS		ORAL
Hospitalization - (DAILY), ORAL		Drug Ineffective	Professional				
Initial or Prolonged Required	3600 MG (1200	Drug Interaction		Neurontin			
Intervention to Prevent Permanent ORAL Impairment/Damage		Hallucination		(Gabapentin)	SS		ORAL
		Influenza					
		Pneumonia					
		Staphylococcal Infection		Anafranil (Clomipramine Hydrochloride)	SS		
				Slo-Bid Gyrocaps (Theophylline)	C		
				Q-Tuss (Ru-Tuss)	C		
				Klor-Con (Potassium Chloride)	C		
				Albuterol (Salbutamol)	C		
				Premarin (Estrogens Conjugated)	C		
				Lotensin (Benazepril Hydrochloride)	C		
				Baycol (Cerivastatin Sodium)	C		
				Xanax (Alprazolam)	C		
				Furosemide (Furosemide)	C		
				Prednisone (Prednisone)	C		

Trazodone	
(Trazodone)	C
Prilosec	
(Omeprazole)	C
Metoclopramide	
(Metoclopramide)	C
Atrovent	
(Ipratropium	
Bromide)	C
Intal (Cromoglicate	
Sodium)	C
Pulmicort	
(Budesonide)	C
Miacalcin	
(Calcitonin, Salmon)	C
Nasonex (Mometasone	
Furoate)	C
Gaviscon Type	
Unknown (Antacids)	C
Darvocet (Propacet)	C
Lonox (Lomotil)	C
Prophlorperazine	
(Prochlorperazine)	C

Freedom Of Information (FOI) Report

Carafate	
(Sucralfate)	C
Hydromet (Aldoril)	C
Calcium With Vitamin	
D (Calcium With	
Vitamin D)	C
Serevent (Salmeterol	
Xinafoate)	C
Singulair	
(Montelukast Sodium)	C
Vitamin E	
(Tocopherol)	C
Ocuvite Extra	
(Ocuvite)	C
B-12	
(Cyanocobalamin)	C
Osteo-Bi-Flex	
(General Nutrients)	C
Prinivil	
(Lisinopril)	C
Clonazepam	
(Clonazepam)	C
Protonix	
(Pantoprazole)	C
Pepcid (Famotidine)	C
Zyrtec (Cetirizine	
Hydrochloride)	C
Mirapex	
(Pramipexole)	C
Wellbutrin Sr	
(Bupropion	
Hydrochloride)	C
Alupent	
(Orciprenaline	
Sulfate)	C

Date:09/17/02ISR Number: 3980740-9Report Type:Expedited (15-DaCompany Report #A208993

Age:52 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 200 MG	Coma Drug Interaction	Health Professional	Sertraline (Sertraline)	PS		ORAL

Initial or Prolonged Pneumonia Streptococcal
(DAILY), ORAL
Required
Intervention to
ORAL
Prevent Permanent
Impairment/Damage

Gabapentin (Gabapentin)	SS	ORAL
Elavil (Amitriptyline Hydrochloride)	C	
Premarin (Estrogens Conjugated)	C	

Date:09/17/02ISR Number: 3980743-4Report Type:Expedited (15-DaCompany Report #A119156
Age:39 YR Gender:Female I/FU:F

Outcome	PT
Death	Agitation
Hospitalization -	Arrhythmia
Initial or Prolonged	Drooling
	Electrocardiogram Qt

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FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
80 MG (BID), ORAL		Prolonged Gastritis Haematemesis Respiratory Arrest	Health Professional	Ziprasidone (Ziprasidone)	PS		ORAL
1200 MG (BID), ORAL				Gabapentin (Gabapentin)	SS		ORAL
				Risperdal (Risperidone)	C		
				Trazadone	C		
				Ativan (Lorazepam)	C		
				Cogentin (Benzatropine Mesilate)	C		
				Multivitamin	C		
				Haldol (Haloperidol)	C		
				Tylenol (Paracetamol)	C		
				Protonix (Pantoprazole)	C		
				Restoril (Temazepam)	C		

Date:09/18/02ISR Number: 3978543-4Report Type:Expedited (15-DaCompany Report #02P-056-0192416-00
Age:68 YR Gender:Male I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG, 1 IN 1 D, PER ORAL		Life-Threatening Hospitalization - Initial or Prolonged Diarrhoea	Foreign Health Professional	Ritonavir Soft Gelatin Capsules (Norvir) (Ritonavir)	PS		ORAL
300 MG, 1 IN 1 D		Hiv Wasting Syndrome Hypertension Iatrogenic Injury	Other	Tenofovir	SS		

1200 MG, 1 IN	Pancytopenia	Amprenavir	SS	
1 D	Renal Failure Acute			
400 MG, 1 IN	Renal Tubular Necrosis	Didanosine	SS	
1 D	Spinal Myelogram Abnormal			
1 CAPSULE, 1	Urinary Tract Infection	Bactrim	SS	ORAL
IN 1 D, PER	Vomiting			
ORAL				
2600 MG		Ibuprofen	SS	
900 MG, 1 IN		Gabapentin	SS	
1 D				
SUBCUTANEOUS	90 MG, 2 IN 1	Ro+29-9800 (T 20-Fusion Inhibitor)	SS	
D,				
SUBCUTANEOUS				
1 DOSAGE		Loperamide	SS	ORAL
FORMS, 1 IN 1				
D, PER ORAL				
10 DROP, 1 IN		Clonazepam	SS	ORAL
1 D, PER ORAL				
1 DOSAGE		Alfuzosin	SS	ORAL
FORMS, 1 IN 1				
D, PER ORAL				
90 MG, 1 IN 1		Enfuirrtide	SS	ORAL
D, PER ORAL				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/18/02ISR Number: 3978829-3Report Type:Expedited (15-DaCompany Report #USA-2002-0000035

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Abortion	Consumer	Oxycontin Tablets 10			
Other		Anhedonia	Health	Mg(Oxycodone			
		Anxiety	Professional	Hydrochloride) Cr			
ORAL		Bronchitis Acute	Other	Tablet	PS		ORAL
		Chest Discomfort		Oxycontin Tablets			
		Chest Pain		40mg(Oxycodone			
		Complications Of Maternal		Hydochloride)Cr			
40 MG, BID,		Exposure To Therapeutic		Tablet	SS		ORAL
ORAL		Drugs					
		Depression		Nubain"Endo"(Nalbuph			
		Drug Dependence		ine Hydrochloride)			
		Drug Hypersensitivity		Injectable	C		
		Drug Ineffective		Decadron			
		Drug Withdrawal Syndrome		(Dexamethasone)	C		
		Dyspnoea		Albuterol			
		Emotional Disorder		(Salbutamol)	C		
		Headache		Plendil (Felodipine)	C		
		Maternal Drugs Affecting		K-Dur (Potassium			
		Foetus		Chloride)	C		
		Pain		Ultram	C		
		Pyrexia		Acyclovir			
		Stridor		(Aciclovir)	C		
		Wheezing		Celebrex (Celecoxib)	C		
				Hydrocortisone			
				(Hydrocortisone)	C		
				Cefadroxil			
				(Cefadroxil)	C		
				Atropine			
				W/Diphenoxylate(Diph			
				enoxylate			
				Hydrochloride,			
				Atorpine Sulfate)	C		
				Cephalexin			
				(Cefalexin)	C		
				Prednisone			
				(Prednisone)	C		
				Colchicine			
				(Colchicine)	C		
				Cytoxan			

(Cyclophosphamide)	C
Zyrtec (Cetirizine Hydrochloride)	C
Tobradex (Dexamethasone, Tobramycin)	C
Hydrocodone W/Acetaminophen	C
Acetaminophen W/Oxycodone	C
Endodan	C
Promethegan (Does Not Code)	C
Monopril (Fonsinopril Sodium)	C
Pravachol (Pravastatin Sodium)	C
Xenical (Orlistat)	C
Lipitor	

Freedom Of Information (FOI) Report

(Atorvastatin)	C
Ultracet (Does Not Code)	C
Ventolin (Salbutamol)	C
Azmacort(Triamcinolone Acetonide)	C
Proventil Inhaler (Salbutamol)	C
Tiazac	C
Aerobid (Flunisolide)	C
Maxzide (Hydrochlorothiazide , Triamterene)	C
Quinidine (Quinidine)	C
Amitriptyline (Amitriptyline)	C
Furosemide (Furosemide)	C
Levaquin(Levofloxacin)	C
Advair Diskus (Fluticasone Propionate, Salmeterol Xinafoate)	C
Bactroban (Mupirocin)	C
Clonidine (Clonidine)	C
Neurontin (Gabapentin)	I
Phenergan "Natrpharm"(Promethazine Hydrochloride)	I

Date:09/19/02ISR Number: 3977719-XReport Type:Expedited (15-DaCompany Report #B0279385A

Age:36 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide	Literature Health	Paxil (Paroxetine Hydrochloride)	PS		ORAL
ORAL							

	Overdose	Professional	Oxycodone (Oxycodone)	SS	ORAL
ORAL			Gabapentin (Gabapentin)	SS	ORAL
ORAL					

Date:09/19/02ISR Number: 3977792-9Report Type:Expedited (15-DaCompany Report #2002057053
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Life-Threatening 1200 MG (400 Other MG, TID),		Completed Suicide	Foreign Health Professional Company Representative	Gabapentin (Gabapentin)	PS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/19/02ISR Number: 3977794-2Report Type:Expedited (15-DaCompany Report #2002056561
Age:2.5 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Maternal Drugs Affecting Foetus Myasthenia Gravis	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:09/19/02ISR Number: 3978022-4Report Type:Expedited (15-DaCompany Report #2002057052
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Other 600 MG (DAILY), ORAL		Angioneurotic Oedema	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Baclofen Oxycodone Fish Oil Prednisone	PS C C C C		ORAL

Date:09/19/02ISR Number: 3978183-7Report Type:Expedited (15-DaCompany Report #NSADSS2002031884
Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged TRANSDERMAL IN 72 HOURS, TRANSD; SEE IMAGE ORAL	50 MCG/H, 1	Burning Sensation Cardiac Failure Congestive Chest Pain Convulsion Decreased Appetite Peripheral Coldness Tremor Vision Blurred	Consumer	Duragesic (Patch) (Fentanyl) Neurontin (Gabapentin) Lopressor (Metoprolol)	PS SS		ORAL

Weight Decreased

Tartrate)	C
Dilantin (Phenytoin Sodium)	C
Lipitor (Atorvastatin)	C
Lasix (Furosemide)	C
Nexium (Esomeparazole Magnesium)	C

Date:09/19/02ISR Number: 3978414-3Report Type:Expedited (15-DaCompany Report #2002057371

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Unevaluable Event	Health Professional	Neurontin (Gabapentin)	PS		

Date:09/19/02ISR Number: 3978416-7Report Type:Expedited (15-DaCompany Report #2002053410

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour Drug Dependence	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
4800 MG		Medication Error		Salbutamol	C		
(DAILY), ORAL							

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Isosorbide
 Mononitrate C
 Combivent C
 Metolazone C
 Metoprolol Succinate C
 Glyceryl Trinitrate C
 Torasemide C
 Simvastatin C
 Potassium Chloride C

Date:09/19/02ISR Number: 3978418-0Report Type:Expedited (15-DaCompany Report #2002056558

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG (TID), Other ORAL	Convulsion Erectile Dysfunction Fatigue	Consumer	Dilantin (Phenytoin Sodium)	PS		ORAL
1200 MG (TID), ORAL	Feeling Abnormal Grand Mal Convulsion		Gabapentin	SS		ORAL
	Insomnia Lethargy Lip Blister Listless Malaise Myocardial Infarction Rash Pruritic Restlessness Somnolence Urinary Incontinence		Oxcarbazepine Valproate Semisodium Pravastatin Sodium Amlodipine Besilate Tocopherol Acetylsalicylic Acid Glucosamine	SS SS C C C C		

Date:09/19/02ISR Number: 4008071-1Report Type:Periodic Company Report #S02-USA-00609-01

Age:31 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 40 MG QD PO	Convulsion	Health Professional	Celexa (Citalopram Hydrobromide)	PS		ORAL

30 MG QAM PO

Celexa (Citalopram Hydrobromide) SS ORAL

20 MG QD PO

Celexa (Citalopram Hydrobromide) SS ORAL

300 MG TID PO

Neurontin (Gabapentin) SS ORAL

Date:09/20/02ISR Number: 3977760-7Report Type:Direct
Age:60 YR Gender:Female I/FU:I

Company Report #USP 55251

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Neurontin	PS	Pfizer	
				Motrin (Ibuprofen)	SS	Pharmacia Corp	

Date:09/23/02ISR Number: 3979442-4Report Type:Expedited (15-DaCompany Report #2002057364
Age:73 YR Gender:Male I/FU:I

Outcome	PT
Other	Erectile Dysfunction Fatigue Oedema Peripheral

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Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
3600 MG, ORAL		Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
			Diuretics	C		
			Acetylsalicylic Acid	C		
			Losartan Potassium	C		
			Bisoprolol	C		

Date:09/23/02ISR Number: 3979470-9Report Type:Expedited (15-DaCompany Report #2002054154
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2100MG	Amnesia Bone Pain	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
(700MG, TID)		Conversion Disorder	Health				
ORAL		Difficulty In Walking	Professional				
		Dysphagia		Paroxetine	C		
		Nuclear Magnetic Resonance Imaging Abnormal		Cyclizine	C		
		Optic Nerve Disorder		Ranitidine	C		
		Pollakiuria		Salbutamol	C		
		Urinary Incontinence		Beclometasone			
		Vision Blurred		Dipropionate	C		
				Panadeine Co	C		
				Morphine Sulfate	C		

Date:09/23/02ISR Number: 3979497-7Report Type:Expedited (15-DaCompany Report #2002056983
Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	UNKNOWN, ORAL	Apraxia Cerebral Atrophy	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other	8 MG (4 MG,	Fall	Professional	Galantamine	SS		ORAL

TWICE DAILY),	Gait Disturbance			
ORAL	Hypertonia			
UNKNOWN	Lacunar Infarction	Colchimax	SS	ORAL
(DAILY), ORAL	Pain In Extremity			
20 MG	Polyneuropathy	Citalopram Hydrobromide	SS	ORAL
(DAILY), ORAL				
0.5 MG		Risperidone	SS	ORAL
(DAILY), ORAL				
180 MG (90		Diltiazem Hydrochloride	SS	ORAL
MG, TWICE				
DAILY), ORAL				
		Glyceryl Trinitrate	C	
		Heptaminol		
		Hydrochloride	C	
		Lorazepam	C	
		Zopiclone	C	
		Paracetamol	C	
		Acetylsalicylate		
		Lysine	C	
		Clomipramine		
		Hydrochloride	C	
		Molsidomine	C	
		Donepezil		

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Hydrochloride C
Tianeptine C

Date:09/23/02ISR Number: 3979675-7Report Type:Expedited (15-DaCompany Report #2002052155
Age:58 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Arthralgia Clostridium Colitis	Consumer Health	Neurontin (Gabapentin)	PS		
Other 300 MG (100 MG, THREE TIMES)	Condition Aggravated Diabetes Mellitus Inadequate Control Diarrhoea	Professional	(Unspecified Intravenous Therapy)	SS		
INTRAVENOUS	INTRAVENOUS					
	Dyspnoea Iatrogenic Injury Neuropathy Peripheral Pain In Extremity Pyrexia		Antibiotics Thiamine Multivitamins Ranitidine Hydrochloride Insulin	C C C C C C		

Date:09/23/02ISR Number: 3979777-5Report Type:Expedited (15-DaCompany Report #2002050494
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Cystitis Hormone Level Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other 600 MG (TID), ORAL	Lesion Of Sciatic Nerve Sciatica Sleep Disorder Tremor Weight Decreased		Estrogens Conjugated Sinemet	SS C		

Date:09/23/02ISR Number: 3979780-5Report Type:Expedited (15-DaCompany Report #2002051909
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue Pain	Consumer Health	Neurontin (Gabapentin)	PS		
3000 MG (DAILY)		Weight Increased	Professional				
				Amitriptyline Hydrochloride	C		

Date:09/23/02ISR Number: 3979819-7Report Type:Expedited (15-DaCompany Report #2002052313
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Carpal Tunnel Syndrome Nephrolithiasis	Consumer	Neurontin (Gabapentin)	PS		
1800 MG (60 Other MG, TID)				Warfarin Sodium	SS		

Date:09/23/02ISR Number: 3979831-8Report Type:Expedited (15-DaCompany Report #2002057062
Age:83 YR Gender:Female I/FU:I

Outcome
Hospitalization -
Initial or Prolonged

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Other

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Confusional State Disorientation	Consumer	Neurontin (Gabapentin)	PS		ORAL
TID, ORAL		Heart Rate Decreased		Temazepam	SS		ORAL
ORAL		Movement Disorder		Karvea Hct	C		
		Pulmonary Thrombosis		Atenolol	C		
		Somnolence		Acetylsalicylic Acid	C		
		Thrombosis		Omeprazole	C		
		White Blood Cell Count Decreased		Propacet	C		
				Pyridoxine Hydrochloride	C		
				Loperamide Hydrochloride	C		
				Potassium Os-Cal	C		

Date:09/23/02ISR Number: 3979834-3Report Type:Expedited (15-DaCompany Report #2002057071
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cough Dyspnoea	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Facial Pain					
300 MG DAILY, ORAL		Respiratory Disorder		Amlodipine Besilate	C		
		Throat Tightness		Sertraline			
		Trigeminal Neuralgia		Hydrochloride	C		

Date:09/23/02ISR Number: 3979871-9Report Type:Expedited (15-DaCompany Report #001-0945-M0100815
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Glucose Increased Blood Potassium Decreased	Health Professional	Neurontin (Gabapentin)	PS		ORAL

SEE IMAGE

200 MG	Fatigue	Celecoxib	SS	ORAL
(DAILY), ORAL	Hypersomnia			
500 MG	International Normalised	Levofloxacin	SS	ORAL
(DAILY), ORAL	Ratio Decreased			
	Joint Stiffness	Clonazepam	C	
	Macular Degeneration	Estrogens Conjugated	C	
	Maculopathy	Furosemide	C	
	Pain In Extremity	Digoxin	C	
	Pneumonia	Warfarin Sodium	C	
	Retinal Disorder	Cyanocobalamin	C	
	Somnolence			
	Tarsal Tunnel Syndrome			
	Vision Blurred			
	Visual Acuity Reduced			
	Vitreous Detachment			

Date:09/23/02ISR Number: 3980040-7Report Type:Expedited (15-DaCompany Report #NSADSS2002032637
Age:20 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health	Fentanyl (Patch) (Fentanyl)	PS		
INGESTION -			Professional				
INTENTIONAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

SUICIDE

Propoxyphene
(Dextropropoxyphene) SS

INTENTIONAL

SUICIDE

Gabapentin
(Gabapentin) SS

INTENTIONAL

SUICIDE

Date:09/24/02ISR Number: 3979550-8Report Type:Expedited (15-DaCompany Report #2002056560
Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Arrhythmia Intracardiac Thrombus	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Rash Maculo-Papular	Professional	Valsartan Hydrochlorothiazide Fluindione	C C		

Date:09/24/02ISR Number: 3980912-3Report Type:Expedited (15-DaCompany Report #S02-FRA-01975-01
Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 20 MG QD PO		Apraxia Cauda Equina Syndrome	Foreign Health	Seropram (Citalopram Hydrobromide)	PS		ORAL
		Fall Gait Disturbance Hypertonia Pain In Extremity Polyneuropathy	Professional Other	Neurontin (Gabapentin) Galantamine Risperdal (Risperidone) Bi-Tildiem (Diltiazem Hydrochloride) Colchimax Aricept (Donepezil Hydrochloride)	SS SS SS SS C		

Anafranil	
(Clomipramine	
Hydrochloride)	C
Corvasal	
(Molsidomine)	C
Nitriderm Tts	
(Glyceryl	
Trinitrate)	C
Hept-A-Mil	
(Heptaminol	
Hydrochloride)	C
Temesta (Lorazepam)	C
Imovane (Zopiclone)	C
Dafalgan	
(Paracetamol)	C
Kardegic	
(Acetylsalicylate	
Lysine)	C
Stablon (Tianeptine)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/24/02ISR Number: 3980985-8Report Type:Expedited (15-DaCompany Report #K200201632

Age:19 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour Disinhibition Euphoric Mood	Foreign Health Professional	Altace Capsules (Ramipril) Capsule, 2.5 Mg	PS		ORAL
2.5 MG, QD, ORAL		Excitability	Other				
				Prednisone (Prednisone) 8 Mg	SS		ORAL
8 MG, QD, ORAL							
				Benzodiazepine Derivatives()	SS		
UNKNOWN							
				Gabapentin (Gabapentin) Tablet	SS		ORAL
ORAL							

Date:09/24/02ISR Number: 3981271-2Report Type:Expedited (15-DaCompany Report #NSADSS2002032637

Age:20 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health	Fentanyl (Patch) (Fentanyl)	PS		
INGESTION - INTENTIONAL			Professional				
				Propoxyphene (Dextropropoxyphene)	SS		
INTENTIONAL SUICIDE							
				Cabapentin (Gabapentin)	SS		
INTENTIONAL SUICIDE							

Date:09/24/02ISR Number: 3981494-2Report Type:Expedited (15-DaCompany Report #2002057832
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Agranulocytosis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID),			Professional				
ORAL				Cefotaxime Sodium	C		

Date:09/24/02ISR Number: 3981496-6Report Type:Expedited (15-DaCompany Report #2002056669
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hepatitis Leukopenia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
DAILY, ORAL			Professional				
Other		Neutropenia					
1800 MG		Tuberculosis					
DAILY ORAL				Ganciclovir	SS		ORAL
ORAL				Nevirapine	SS		ORAL
BID, ORAL				Kaletra	SS		ORAL
BID, ORAL				(Tenofir)	SS		ORAL
DAILY, ORAL				Valganciclovir	SS		ORAL
BID, ORAL				Bactrim	C		
				Calcium Folate	C		
				Paracetamol	C		
				Loperamide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/24/02ISR Number: 3981537-6Report Type:Expedited (15-DaCompany Report #2002057367

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Neoplasm	Consumer	Neurontin			
(DAILY), ORAL		Drug Tolerance Decreased		(Gabapentin)	PS		ORAL
				Estrogen Product	SS		ORAL
ORAL				Atenolol	C		
				Amitriptyline			
				Hydrochloride	C		
				Vitamins	C		
				Calcium	C		
				Minerals Nos	C		
				Tamoxifen	C		

Date:09/25/02ISR Number: 3981330-4Report Type:Direct Company Report #CTU 177116

Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -							
Initial or Prolonged		Neuroleptic Malignant		Risperdal 3mg			
3MG 00/11 HS		Syndrome		Janssen	PS	Janssen	ORAL
PO		Rhabdomyolysis					
				Neurontin 600mg			
600MG 1/1AM				Parke-Davis	SS	Parke-Davis	
00/11HS							

Date:09/25/02ISR Number: 3982050-2Report Type:Expedited (15-DaCompany Report #2002057840

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -							
Initial or Prolonged		Abdominal Pain Upper	Foreign	Neurontin			
900 MG TID		Meningitis Viral	Health	(Gabapentin)	PS		
			Professional				

Company
Representative

Date:09/25/02ISR Number: 3982413-5Report Type:Expedited (15-DaCompany Report #2002055716

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Eosinophilia Rash	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:09/25/02ISR Number: 3982542-6Report Type:Expedited (15-DaCompany Report #NSADSS2002032666

Age:50 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Ultram (50 Mg Tablet) (Tramadol Hydrochloride)	PS		ORAL
ORAL				Gabapentin (Gabapentin)	SS		ORAL
ORAL				Rofecoxib	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/27/02ISR Number: 3982397-XReport Type:Expedited (15-DaCompany Report #FLUV00302002330

Age:91 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Coordination Abnormal Dizziness Somnolence	Foreign Other	Floxyfral (Fluvoxamine Maleate)	PS		ORAL
100 MG DAILY							
PO				Neurontin (Gabapentin)	SS		ORAL
500 MG DAILY							
PO, 300 MG							
TID PO				Plendil (Felodipine) Comilorid (Amiloride Hcl W/Hydrochlorothiazid e) Seresta (Oxazepam)	C C C		

Date:09/27/02ISR Number: 3982761-9Report Type:Expedited (15-DaCompany Report #2002057829

Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Back Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG							
(TID), ORAL				Levothyroxine Sodium Paracetamol	C C		

Date:09/27/02ISR Number: 3982778-4Report Type:Expedited (15-DaCompany Report #2002058031

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Coma	Consumer	Neurontin (Gabapentin)	PS	ORAL
ORAL	Convulsion				
	Myocardial Infarction				
	Nausea				
	Vomiting				

Date:09/27/02ISR Number: 3982834-0Report Type:Expedited (15-DaCompany Report #2002057843

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		
Other		Bradycardia Extrasystoles					
700 MG		Hypotension					
(DAILY),		Neutrophilia Platelet Count Decreased		Gabapril Lamotrigine	C C		

Date:09/27/02ISR Number: 3983430-1Report Type:Expedited (15-DaCompany Report #2002057870

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		Coordination Abnormal Dizziness					
500 MG, ORAL		Somnolence	Professional	Fluvoxamine Maleate	SS		ORAL
100 MG, ORAL		Vertigo		Felodipine Moduretic "Msd"	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Oxazepam C

Date:09/27/02ISR Number: 3983437-4Report Type:Expedited (15-DaCompany Report #2002055562
Age:94 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 400 MG		Coordination Abnormal Fall	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Initial or Prolonged (DAILY), ORAL		Haematoma	Professional				
Other		Tremor Wound		Glyceryl Trinitrate	C		
				Furosemide	C		
				Ramipril	C		
				Isosorbide Dinitrate	C		
				Digoxin	C		
				Aporex	C		
				Zopiclone	C		

Date:09/27/02ISR Number: 3984730-1Report Type:Expedited (15-DaCompany Report #2002057367
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Brain Neoplasm	Consumer	Neurontin (Gabapentin)	PS		ORAL
DAILY, ORAL				Unknown Estrogen Product	SS		ORAL
ORAL				Atenolol	C		
				Amitriptyline Hydrochloride	C		
				Vitamins	C		
				Calcium	C		
				Minerals Nos	C		
				Tamoxifen	C		

Date:10/01/02ISR Number: 3983145-XReport Type:Expedited (15-DaCompany Report #2002056560
Age:80 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Arrhythmia Atrial Thrombosis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Drug Toxicity	Professional	Clobazam	SS		ORAL
		Encephalopathy Exanthem Rash Maculo-Papular Somnolence		Valsartan Hydrochlorothiazide Fluindione	C C		

Date:10/01/02ISR Number: 3983453-2Report Type:Expedited (15-DaCompany Report #2002058135
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 400 MG, ORAL		Laryngeal Oedema	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
			Professional Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/01/02ISR Number: 3983716-0Report Type:Direct
Age:49 YR Gender:Female I/FU:I

Company Report #CTU 177629

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
36 G QD DAILY [APPROX 1 MONTH]	1	Leukopenia		Gabapentin	PS		

Date:10/01/02ISR Number: 3984856-2Report Type:Expedited (15-DaCompany Report #2002054820
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (600 Disability MG, BID), Other ORAL		Anxiety Blood Cholesterol Increased Disturbance In Attention Drug Tolerance Increased Eye Pain Feeling Abnormal Feeling Drunk Headache Hypoaesthesia Ill-Defined Disorder Major Depression Palpitations Panic Attack	Consumer Health Professional	Neurontin (Gabapentin) Tizanidine Hydrochloride Lorazepam Morphine Sulfate Alprazolam Metoprolol Succinate Rofecoxib	PS C C C C C C		ORAL

Date:10/01/02ISR Number: 3984895-1Report Type:Expedited (15-DaCompany Report #001-0945-990998
Age:49 YR Gender:Female I/FU:I

Outcome	PT
Disability	Abdominal Pain Alopecia Anorexia Arthralgia

Back Pain
Bone Pain
Chest Pain
Chills
Choking Sensation
Contusion
Coordination Abnormal
Cystitis
Diarrhoea
Disturbance In Attention
Dizziness
Emotional Disorder
Fall
Fatigue
Feeling Abnormal
Female Orgasmic Disorder
Flatulence
Gingival Bleeding
Hypertonia
Insomnia
Joint Stiffness
Madarosis
Muscle Tightness
Nausea
Pyrexia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100 MG (HS), ORAL, SEE IMAGE		Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Oxycodone Flexeril (Cyclobenzaprine Hydrochloride) Vicodin Valium (Diazepam) Baclofen	C C C C C		

Date:10/02/02ISR Number: 3985517-6Report Type:Direct Company Report #CTU 177733
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 100MG 3 TIMES Intervention to ORAL Prevent Permanent Impairment/Damage		Heart Rate Irregular		Neurontin 100mg	PS		ORAL
				Lipitor Paxil Fosomax Calcium Centrum Silver Vit D	C C C C C C		

Date:10/02/02ISR Number: 4022225-XReport Type:Periodic Company Report #2000590
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1000 MG 2/D		Convulsion	Consumer	Keppra	PS		ORAL

PO

1400 MG PO

1700 MG PO

Neurontin	SS	ORAL
Neurontin	SS	ORAL
Tegretol	C	

Date:10/03/02ISR Number: 3982624-9Report Type:Expedited (15-DaCompany Report #WAES 0209USA02503

Age:50 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional	Vioxx	PS	Merck & Co., Inc	ORAL
Other				Gabapentin	SS		
				Tramadol	SS		

Date:10/03/02ISR Number: 3985226-3Report Type:Expedited (15-DaCompany Report #2002052931

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Dependence Medication Error	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Tramadol Hydrochloride	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/03/02ISR Number: 3985227-5Report Type:Expedited (15-DaCompany Report #2002057071
 Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
300 MG		Cough		(Gabapentin)	PS		ORAL
(DAILY), ORAL		Dyspnoea					
		Lung Disorder		Amloldipine Besilate	C		
		Throat Tightness		Sertraline			
		Trigeminal Neuralgia		Hydrochloride	C		

Date:10/03/02ISR Number: 3985249-4Report Type:Expedited (15-DaCompany Report #2002050043
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Distension	Consumer	Neurontin			
(TWICE		Blood Arsenic Increased		(Gabapentin)	PS		ORAL
DAILY), ORAL		Blood Heavy Metal					
ORAL		Increased		Lamotrigine	SS		ORAL
		Hypertension		Clonazepam	SS		
		Laboratory Test Abnormal					
		Nausea					
		Respiratory Disorder					
		Weight Increased					

Date:10/03/02ISR Number: 3986919-4Report Type:Expedited (15-DaCompany Report #2002058571
 Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Epilepsy	Foreign	Neurontin			
Initial or Prolonged		Overdose	Health	(Gabapentin)	PS		ORAL
900 MG (300							
MG,TID), ORAL			Professional				
			Company	Vitamins	C		

Representative Carbamazepine C
Rinace C

Date:10/03/02ISR Number: 3987784-1Report Type:Expedited (15-DaCompany Report #2002059032
Age:19 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Epilepsy	Foreign	Neurontin			
Initial or Prolonged	Hepatitis C	Health	(Gabapentin)	PS		ORAL
300 MG (TID),						
Other	Memory Impairment	Professional				
ORAL						
	Post Procedural		Amityptilin	C		
	Complication		Metamizole Sodium	C		
			Milgamma	C		
			Keltican/ Old Form	C		
			Capsaicin	C		
			Phytodolor	C		
			Emla Plaster	C		

Date:10/03/02ISR Number: 3987818-4Report Type:Expedited (15-DaCompany Report #2002053360
Age:48 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

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2.4 GRAM	(DAILY), ORAL	Blood Chloride Decreased Blood Creatinine Increased	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Blood Sodium Decreased Dehydration	Company Representative	Durogesic Plaster	C		

Date:10/03/02ISR Number: 3987825-1Report Type:Expedited (15-DaCompany Report #2002058564
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Neutropenia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/03/02ISR Number: 3987921-9Report Type:Expedited (15-DaCompany Report #2002058715
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	1200 MG, QID ORAL	Blister Foot Fracture Gastritis	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Impaired Healing Infection Irritable Bowel Syndrome		Sertraline Hydrochloride Bupropion Hydrochloride Librax Metformin Hydrochloride Pioglitazone Valsartan Unspecified Pain	C C C C C C C		

Medication

C

Date:10/03/02ISR Number: 3987922-0Report Type:Expedited (15-DaCompany Report #2002058714

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Sarcoidosis	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG QID							
ORAL							

Sertraline	
Hydrochloride	C
Bupropion	
Hydrochloride	C
Librax	C
Metformin	
Hydrochloride	C
Pioglitazone	C
Valsartan	C
Unspecified Pain	
Medication	C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/03/02ISR Number: 3987931-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200613

Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600							
MG TID) ORAL				Lansoprazole	C		

Date:10/03/02ISR Number: 3987949-9Report Type:Expedited (15-DaCompany Report #2002058579

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aplastic Anaemia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:10/03/02ISR Number: 3987990-6Report Type:Expedited (15-DaCompany Report #044-0945-M0100204

Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Cholestasis Hepatic Steatosis	Foreign Literature	Neurontin (Gabapentin)	PS		ORAL
900 MG (300							
Other		Hepatitis	Health				
MG, TID),		Jaundice Cholestatic	Professional				
ORAL				Human Mixtard	C		
				Metformin	C		
				Amitriptyline	C		
				Dihydrocodeine	C		
				Ramipril	C		

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coma	Health	Neurontin			
		Grand Mal Convulsion	Professional	(Gabapentin)	PS		ORAL
900 MG (TID),							
ORAL		Hyperventilation					
		Memory Impairment		Amitriptyline	SS		
		Somnolence		Levothyroxine Sodium	C		
				Glipizide	C		
				Metformin			
				Hydrochloride	C		

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Accidental Overdose	Literature	Oxycontin Tablets			
		Head Injury	Health	(Oxycodone			
		Insomnia	Professional	Hydrochloride) Cr			
SEE IMAGE		Medication Error		Tablet	PS		
				Neurontin			
1200 MG				(Gabapentin)	SS		
				Methadone			
SEE TEXT				(Methadone)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/04/02ISR Number: 3989083-0Report Type:Expedited (15-DaCompany Report #2002055566

Age:65 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Difficulty In Walking	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL				All Other Therapeutic Products	C		

Date:10/07/02ISR Number: 3987115-7Report Type:Expedited (15-DaCompany Report #2002054675

Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (TID),		Drug Level Decreased	Health				
ORAL		Hemiplegia	Professional				
		Migraine		Bupropion Hydrochloride	SS		
		Muscle Twitching		Citalopram Hydrobromide	C		
		Paranoia					

Date:10/07/02ISR Number: 3987121-2Report Type:Expedited (15-DaCompany Report #001-0945-950546

Age:12 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Coma	Health	Neurontin (Gabapentin)	PS		ORAL
600 (TID),		Convulsion	Professional				
Other		Dehydration					
ORAL		Haemodialysis		Clavulin	SS		ORAL
ORAL		Loss Of Consciousness		Carnitine	C		
		Mental Status Changes		Phenytoin Sodium	C		
		Nephropathy Toxic		Valproate Sodium	C		
		Pneumonia		Primidone	C		
		Renal Failure					

Respiratory Failure
Somnolence
Uraemic Encephalopathy

Date:10/07/02ISR Number: 3987123-6Report Type:Expedited (15-DaCompany Report #2002059319

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Operation	Consumer	Neurontin			
		Drug Ineffective	Health	(Gabapentin)	PS		ORAL
ORAL		Pharmaceutical Product	Professional				
		Complaint					

Date:10/07/02ISR Number: 3987349-1Report Type:Expedited (15-DaCompany Report #2002059746

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Decreased	Consumer	Dilantin (Phenytoin			
		Convulsion		Sodium)	PS		ORAL
ORAL		Hypersomnia		Neurontin			
		Tremor		(Gabapentin)	SS		ORAL
QD, ORAL		Vomiting		Famotidine	C		
		Weight Increased		Metoclopramide	C		
				Omeprazole	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ipratropium Bromide C
 Seravent C
 Asthmacort C
 Diazepam C
 Oxybutynin
 Hydrochloride C
 Furosemide C
 Pirbuterol Acetate C
 Metoprolol C
 Promethazine
 Hydrochloride C
 Potassium Citrate C

Date:10/07/02ISR Number: 3987353-3Report Type:Expedited (15-DaCompany Report #2002059747
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Nephrolithiasis	Consumer	Dilantin (Phenytoin Sodium)	PS		ORAL
1200 MG, ORAL			Neurontin (Gabapentin)	SS		ORAL
			Famotidine	C		
			Metoclopramide	C		
			Omeprazole	C		
			Ipratropium Bromide	C		
			Seravent	C		
			Asthmacort	C		
			Diazepam	C		
			Oxybutynin			
			Hydrochloride	C		
			Furosemide	C		
			Pirbuterol Acetate	C		
			Metoprolol	C		
			Promethazine			
			Hydrochloride	C		
			Potassium Citrate	C		

Date:10/07/02ISR Number: 3987511-8Report Type:Expedited (15-DaCompany Report #2002059601
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue Gastrointestinal Disorder	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/07/02ISR Number: 3989265-8Report Type:Expedited (15-DaCompany Report #2002050348
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Hypermetropia Myopia	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
600 MG			Professional				
(DAILY), ORAL		Presbyopia Visual Disturbance					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/08/02ISR Number: 3989841-2Report Type:Expedited (15-DaCompany Report #2002060106

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
4800 MG (1600 MG TID) ORAL		Spinal Fracture					

Date:10/08/02ISR Number: 3989856-4Report Type:Expedited (15-DaCompany Report #02P-163-0201355-00

Age:19 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Literature Health Professional	Valproic Acid (Depakene) (Valproic Acid) Olanzapine Gabapentin	PS SS SS		

Date:10/08/02ISR Number: 3989999-5Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged ORAL		Back Pain Brain Neoplasm	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Disability Other			Professional Company Representative				

Date:10/08/02ISR Number: 3990001-XReport Type:Expedited (15-DaCompany Report #2002051678

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams	Consumer	Neurontin			

(TID), ORAL

Anorexia

Health

(Gabapentin)

PS

ORAL

Blood Pressure Increased
Depression
Drug Withdrawal Syndrome
Fear
Medication Error
Suicidal Ideation
Weight Increased

Professional

Date:10/08/02ISR Number: 3990004-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200295
Age:74 YR Gender:Male I/FU:F

Outcome
Death
Hospitalization -
Initial or Prolonged
Other

PT
Arrhythmia
Cardiac Arrest
Coordination Abnormal
Decreased Activity
Deep Vein Thrombosis
Electromechanical
Dissociation
Fall
Post Procedural
Complication
Pulmonary Embolism

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pyrexia Subdural Haematoma Vomiting	Report Source	Product	Role	Manufacturer	Route
2400 MG (600 MG, QID)			Health Professional Company	Neurontin (Gabapentin)	PS		
400 MG (100 MG, Q6H)			Representative	Cerebyx (Fosphenytoin Sodium)	SS		
2000 MG (500 MG, QID)				Metronidazole (Metronidazole)	SS		
800 MG (200 MG, QID)				Carbamazepine	SS		
				Clavulin	SS		

Date:10/08/02ISR Number: 3990009-4Report Type:Expedited (15-DaCompany Report #2002053412
Age: Gender:Female I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1500 MG (DAILY)		Blood Pressure Increased Convulsion	Consumer	Neurontin (Gabapentin)	PS		
2000 MG (DAILY), ORAL		Dyskinesia Dyspnoea Feeling Hot		Levetiracetam	SS		ORAL
		Grand Mal Convulsion Petit Mal Epilepsy					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Disability	Back Pain	Foreign	Neurontin			
Other	Chromaturia	Health	(Gabepentin)	PS		
1200 MG (TID)						
	Depression	Professional	Clonazepam	C		
	Dysuria					
	Feeling Abnormal					
	Headache					
	Insomnia					
	Irritability					
	Mental Disorder					
	Nausea					
	Pruritus					
	Somnolence					
	Urinary Retention					
	Vertigo					

Outcome	PT
Disability	Abdominal Distension
	Abnormal Behaviour
	Aggression
	Asthenia
	Back Pain
	Bone Neoplasm

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Condition	Report Source	Product	Role	Manufacturer	Route
1200 MG		Chest Pain Condition Aggravated Constipation	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
(TID), OAL		Dyspnoea					
		Ganglion		Amitriptyline			
		Nasal Congestion		Hydrochloride	C		
		Neck Pain		Diazepam	C		
		Neuropathy Peripheral		Sertraline	C		
		Oedema Peripheral					
		Pain					
		Pain In Extremity					
		Peripheral Coldness					
		Sciatica					
		Tremor					

Date:10/09/02 ISR Number: 3990847-8 Report Type:Expedited (15-DaCompany Report #2002051548
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Pressure Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Difficulty In Walking					
300 MG DAILY,		Dizziness					
ORAL		Retinal Haemorrhage		Unspecified High Blood Pressure Medication	C		
		Vision Blurred					

Date:10/10/02 ISR Number: 3991889-9 Report Type:Expedited (15-DaCompany Report #2002060109
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Difficulty In Walking	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Fall					
		Muscular Weakness		Phenelzine Sulfate	C		

Paresis
Phantom Pain
Somnolence

Anovlar C
Omeprazole C
Clonazepam C
Trazodone C

Date:10/10/02ISR Number: 3991891-7Report Type:Expedited (15-DaCompany Report #2002060118

Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence	Health Professional	Neurontin(Gabapentin)	PS		ORAL
4000 MG							
(BID), ORAL				Ibuprofen	C		

Date:10/10/02ISR Number: 3992124-8Report Type:Expedited (15-DaCompany Report #2002060110

Age: Gender:Female I/FU:I

Outcome	PT
Disability	Dizziness
Other	Exanthem
	Fall
	Gastroenteritis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Syncope

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
600 MG	(DAILY), ORAL	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Candesartan Cilexetil	C		

Date:10/11/02ISR Number: 3991308-2Report Type:Expedited (15-DaCompany Report #2002052323
 Age: Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1800 MG (600 MG TID)	ORAL	Convulsion Drug Ineffective	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Pain		Hydrochlorothiazide	C		
			Paraesthesia		Cetirizine	C		
			Pharmaceutical Product Complaint		Hydrochloride	C		
					Tramadol	C		
					Hydrochloride	C		
					Metaxalone	C		
					Metoprolol	C		

Date:10/11/02ISR Number: 3992628-8Report Type:Expedited (15-DaCompany Report #2002060656
 Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		
					Ethanol	SS		
					Unspecified Antidepressants	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Chronic Obstructive Pulmonary Disease	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Convulsion		Prednisone	SS		ORAL
ORAL		Medication Error		Theophylline	SS		ORAL
ORAL		Weight Increased		Paroxetine Hydrochloride Donepezil Hydrochloride	C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Condition Aggravated Dizziness	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
300 MG (DAILY,ORAL		Dysphonia	Professional				
		Herpes Zoster		Bendroflumethiazide Estradiol	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/02ISR Number: 3993148-7Report Type:Expedited (15-DaCompany Report #061-0945-M0200096
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pain Of Skin	Foreign	Neurontin			
2400 MG		Tremor	Health	(Gabapentin)	PS		ORAL
(DAILY), ORAL			Professional				
			Company	Nifedipine	C		
			Representative	Atorvastatin	C		
				Sertraline			
				Hydrochloride	C		

Date:10/15/02ISR Number: 3993150-5Report Type:Expedited (15-DaCompany Report #2002060940
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Acute Febrile	Foreign	Neurontin			
900 MG (300		Neutrophilic Dermatitis	Health	(Gabapentin)	PS		ORAL
MG, TID),			Professional				
ORAL							

Date:10/15/02ISR Number: 3993173-6Report Type:Expedited (15-DaCompany Report #2002-BP-04878BP(0)
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Circulatory Collapse	Health	Clonidine			
SEE IMAGE	18 DAY	Colitis Ischaemic	Professional	Hydrochloride			
6MG (2MG, 2		Gastrointestinal	Other	Tablets (Clonidine)			
MG TID)	44 DAY	Dysplasia		(Ta)(Clonidine-Hcl)	PS		ORAL
		Loss Of Consciousness		Cilansetron (Nr)	SS		ORAL
		Protein S Increased					
		Rectal Adenoma		Gabapentin			

600 MG(2002	Rhinitis	(Gabapentin)(Nr)	SS	ORAL
MG, 200MG	Shock			
TID)	Suprapubic Pain			
34	DAY			
	Upper Respiratory Tract Infection	Paracetamol (Paracetamol) (Nr)	C	
		Omeprazole (Omeprazole)(Nr)	C	
		Doxepin (Doxepin)(Nr)	C	
		Aspirin (Nr)	C	

Date:10/15/02ISR Number: 3993489-3Report Type:Expedited (15-DaCompany Report #001-0945-990998
Age:49 YR Gender:Female I/FU:F

Outcome	PT
Disability	Alopecia
	Anorexia
	Anorgasmia
	Arthralgia
	Back Pain
	Bone Pain
	Chest Pain
	Contusion
	Coordination Abnormal
	Cystitis
	Diarrhoea
	Dissociation

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Disturbance In Attention Dizziness Emotional Disorder	Report Source	Product	Role	Manufacturer	Route
100 MG (HS), ORAL		Fall Fatigue Flatulence Gastrointestinal Pain Gingival Bleeding Hypertonia Insomnia Joint Stiffness Madarosis Medication Error Nausea Pyrexia Reaction To Medical Agent Preservatives Retching Thirst Urinary Retention	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Oxycodone Cyclobenzaprine Hydrochloride Vicodin Diazepam Baclofen	C C C C C		

Date:10/15/02ISR Number: 3993525-4Report Type:Expedited (15-DaCompany Report #2002059319

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Drug Ineffective Pharmaceutical Product Complaint	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:10/15/02ISR Number: 3993527-8Report Type:Expedited (15-DaCompany Report #2002060230

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged DAILY		Cholesterosis Chronic Fatigue Syndrome	Health Professional	Neurontin (Gabapentin)	PS		

Other		Colitis		Progesterone	SS	ORAL
DAILY, ORAL						
		Colonic Polyp		Interferon Beta	SS	
INTRAMUSCULAR	30 MCG					
		Food Allergy				
WEEKLY,						
		Gastrooesophageal Reflux				
INTRAMUSCULAR						
		Disease		Modafinil	C	
		Irritable Bowel Syndrome				
		Nephrolithiasis				
		Ovarian Cyst				
		Pancreatitis Acute				
		Teratoma				

Date:10/15/02ISR Number: 3993540-0Report Type:Expedited (15-DaCompany Report #2002060225
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2800 MG, ORAL		Confusional State Drug Toxicity	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
		Somnolence		Amitriptyline Hydrochloride	SS		ORAL
75 MG, (DAILY), ORAL				Fentanyl	SS		
TRANSDERMAL	TRANSD			Metoclopramide Hydromorphone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride C
 Rofecoxib C
 Pantoprazole C

Date:10/15/02ISR Number: 3993544-8Report Type:Expedited (15-DaCompany Report #2002060616
 Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 2400 Initial or Prolonged MG(DAILY), ORAL		Cardiac Failure Hypertensive Crisis	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:10/16/02ISR Number: 3994599-7Report Type:Expedited (15-DaCompany Report #US010179
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2 MG QID ORAL		Agranulocytosis	Health	Gabitril	PS		ORAL
525 MG QD		Bradycardia	Professional	Neurontin	SS		ORAL
ORAL		Hypotension					
1500 MG QD		Thrombocytopenia		Neurontin	SS		ORAL
ORAL		White Blood Cell Count Decreased		Lamictal	SS		ORAL
700 MG QD				Lamictal	SS		ORAL
ORAL				Lithium	C		
400 MG QD				Sonata	C		
ORAL				Claritin	C		

Multivitamins C
Colace C

Date:10/16/02ISR Number: 3994742-XReport Type:Expedited (15-DaCompany Report #2002057367
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Intolerance	Consumer Health	Nuerontin (Gabapentin)	PS		ORAL
400 MG (200 MG, TWICE DAILY), ORAL			Professional				
ORAL				Estrogen Nos	SS		ORAL
				Atenolol	C		
				Amitriptyline Hydrochloride	C		
				Vitamins	C		
				Calcium	C		
				Mineral Nos	C		
				Tamoxifen	C		
				Nortriptyline	C		

Date:10/16/02ISR Number: 3994793-5Report Type:Expedited (15-DaCompany Report #DSA_22041_2002
Age:55 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Literature Health Professional	Diltiazem (Long-Acting)	PS		
				Atenolol	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Gabapentin SS

Date:10/16/02ISR Number: 4010525-9Report Type:Periodic Company Report #001-0945-M0200515
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG (QID), ORAL		Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Cilest (Nestabs) (Nestabs) Unspecified Antibiotics	C C C		

Date:10/16/02ISR Number: 4010526-0Report Type:Periodic Company Report #001-0945-M0200421
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (TID), ORAL		Depression Dizziness Dyspnoea	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Emotional Disorder Infection Intentional Misuse Oedema Peripheral Pain Rash Visual Disturbance		Lisinopril Amlodipine Propranolol Hydrochloride Estropipate	C C C C		

Date:10/16/02ISR Number: 4010527-2Report Type:Periodic Company Report #2002051142
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Consumer	Neurontin			

3600 MG (1200	Intentional Misuse	(Gabapentin)	PS	ORAL
MG, TID),	Medication Error			
ORAL	Pain			
		Methadone	C	

Date:10/16/02ISR Number: 4010528-4Report Type:Periodic Company Report #001-0945-M0101300
 Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract	Consumer	Neurontin			
200 MG (100		Diplopia	Health	(Gabapentin)	PS		ORAL
MG, BID),		Dizziness	Professional				
ORAL		Electrocardiogram					
		Abnormal		Nabumetone	C		
		Hypertension					
		Liver Function Test					
		Abnormal					
		Nausea					
		Pain					
		Photophobia					
		Retinal Disorder					
		Speech Disorder					
		Visual Disturbance					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4010529-6Report Type:Periodic
Age:54 YR Gender:Male I/FU:I

Company Report #2002050434

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Urea Increased	Health	Neurontin (
ORAL		Headache	Professional	Gabapentin)	PS		ORAL
		Hypertonia		Glyceryl Trinitrate	C		
		Rash		Isosorbide			
		Testicular Disorder		Mononitrate	C		
				Metronidazole	C		
				Selium Shampoo	C		
				Simvastatin	C		
				Atenolol	C		
				Hydrochlorothiazide	C		
				Glibenclamide	C		
				Rabeprazole	C		
				Lisinopril	C		
				Furosemide	C		
				Testosterone	C		
				Trazodone	C		
				Fluoxetine	C		

Date:10/16/02ISR Number: 4010530-2Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-M0200016

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coordination Abnormal	Consumer	Neurontin (
1200 MG		Gait Disturbance		Gabapentin)	PS		ORAL
(400MG TID),		Intentional Misuse					
ORAL		Neuropathy					

Date:10/16/02ISR Number: 4010531-4Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-M0101413

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Arthralgia	Health	Neurontin (Tablets)			

Initial or Prolonged Pyrexia Professional (Gabapentin) PS ORAL
300 MG

(DAILY), ORAL Urticaria

Paroxetine
Hydrochloride C
Rabeprazole C
Estrogens Conjugated C
Levothyroxine Sodium C
Rofecoxib C
Alprazolam C
Ranitidine C
Nadolol C

Date:10/16/02 ISR Number: 4012445-2 Report Type:Periodic Company Report #2001085961US
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Intentional Misuse	Health Professional	Xanax (Alprazolam) Tablet	PS		ORAL
SUBCUTANEOUS	20 MG, QD,			Copaxone (Glatiramer Acetate)	SS		
SUBCUTANEOUS				Neurontin (Gabapentin)	SS		ORAL
1200 MG, QD,							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Tylenol With Codeine
(Codeine Phosphate,
Paracetamol) SS
Roxicet(Oxycodone
Hydrochloride) SS

ORAL

0.25 MG, QD,

ORAL

Flonase (Fluticasone
Propionate) C
Detrol (Tolterodine
L-Tartrate) C

Date:10/16/02ISR Number: 4012450-6Report Type:Periodic
Age:33 YR Gender:Female I/FU:I

Company Report #001-0945-M0200173

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1600 MG (BID), ORAL		Amenorrhoea Anxiety Emotional Disorder Leukopenia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
200 MG (DAILY), ORAL				Sertraline Hydrochloride(Sertra line Hydrochloride)	SS		ORAL
2 MG (DAILY)				Risperidone	SS		
				Alprazolam Valproate Semisodium Diphenhydramine Hydrochloride	C C C C		

Date:10/16/02ISR Number: 4012452-XReport Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-M0200052

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion	Health	Neurontin(Gabapentin			

900 MG (TID),	Depression	Professional)	PS	ORAL
ORAL	Intentional Misuse				
	Medication Error		Morphine Pump	C	
	Muscle Twitching		Bupivacaine	C	
	Pain		Lorazepam	C	
	Tremor		Hydromorphone		
			Hydrochloride	C	
			Estrogens	C	
			Trazodone	C	
			Amitriptyline		
			Hydrochloride	C	
			Orphenadrine Citrate	C	
			Unspecified		
			Narcotics	C	

Date:10/16/02ISR Number: 4012454-3Report Type:Periodic Company Report #001-0945-M0200091
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression	Health	Neurontin(Gabapentin			
300 MG		Pain	Professional)	PS		ORAL
(DAILY), ORAL		Suicidal Ideation					
				Estradiol	C		
				(Non Steriodal			
				Anti-Inflammatory			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Drugs) C
 Vicodin C
 Donnatal C
 Colestipol
 Hydrochloride C

Date:10/16/02ISR Number: 4012455-5Report Type:Periodic Company Report #001-0945-M0200111
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin(Gabapentin)	PS		

Date:10/16/02ISR Number: 4012456-7Report Type:Periodic Company Report #001-0945-M0200112
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin(Gabapentin)	PS		

Date:10/16/02ISR Number: 4012458-0Report Type:Periodic Company Report #001-0945-M0200113
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin(Gabapentin)	PS		

Date:10/16/02ISR Number: 4012460-9Report Type:Periodic Company Report #001-0945-M0200114
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin(Gabapentin)	PS		

Date:10/16/02ISR Number: 4012461-0Report Type:Periodic Company Report #001-0945-M0200115

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:10/16/02ISR Number: 4012467-1Report Type:Periodic Company Report #001-0945-M0200116

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin(Gabapentin)	PS		

Date:10/16/02ISR Number: 4012469-5Report Type:Periodic Company Report #001-0945-M0200117

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4012472-5Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-M0200262

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin			
Other		Convulsion	Health Professional	(Gabapentin)	PS		ORAL
ORAL				Sumatriptan Succinate	SS		
				Axotal (Old Form)	SS		ORAL
ORAL				Fluoxetine Hydrochloride	SS		ORAL
ORAL				Buspirone Hydrochloride	C		
				Metronidazole	C		
				Estrogens Conjugated	C		
				Zolpidem Tartrate	C		
				Alprazolam	C		
				Levothyroxine Sodium	C		
				Estradiol	C		
				Cetirizine Hydrochloride	C		
				Estrogens Conjugated	C		

Date:10/16/02ISR Number: 4012478-6Report Type:Periodic
 Age:65 YR Gender:Female I/FU:I

Company Report #001-0945-M0200270

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin			
Other		Lymphadenopathy	Consumer	(Gabapentin)	PS		ORAL
DAILY, ORAL				Nortriptyline	C		
				Pantoprazole	C		
				Losartan Potassium	C		
				Hydrochlorothiazide	C		
				Hyoscyamine Sulfate	C		

Date:10/16/02ISR Number: 4012485-3Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-M0200272

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Agitation Bipolar I Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
20 MG, DAILY, ORAL		Diarrhoea		Ziprasidone	SS		ORAL
DAILY, ORAL		Hypotension					
		Infection		Azithromycin	SS		ORAL
		Rash		Methylphenidate Hydrochloride	C		
		Somnolence		Alprazolam	C		
		Syncope		Venlafaxine			
		Tachycardia		Hydrochloride	C		
		Tremor		Citalopram			
		Weight Increased		Hydrobromide	C		
				Topiramate	C		

Date:10/16/02ISR Number: 4012501-9Report Type:Periodic Company Report #001-0945-M0200288
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other		Cerebrovascular Disorder Myocardial Infarction	Consumer	Neurontin (Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4012502-0Report Type:Periodic
Age:64 YR Gender:Male I/FU:I

Company Report #001-0945-M0200319

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Neurontin (Gabapentin)	PS		ORAL
DAILY, ORAL			Company Representative	Losartan Potassium	C		
				Unk	C		
				Glipizide	C		
				Acetylsalicylic Acid	C		

Date:10/16/02ISR Number: 4012503-2Report Type:Periodic
Age:80 YR Gender:Female I/FU:I

Company Report #001-0945-M0200335

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia Hypertonia	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 Other MG, TID),		Hypotension					
ORAL		Somnolence					
		Syncope		Risperidone	C		
		Unevaluable Event		Sinemet	C		
				Clopidogrel	C		
				Fosinopril Sodium	C		

Date:10/16/02ISR Number: 4012504-4Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-M0200352

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dyspnoea Hypercholesterolaemia	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (200 Other MG, TID) ORAL		Hyperlipidaemia					
		Neuralgia		Ibuprofen	C		
		Pneumonia		Oxycocet	C		
				Bulbital W/Asprin,			

Caffeine C
Baclofen C
Paroxetine C
Hydrochloride C
Lorazepam C

Date:10/16/02ISR Number: 4012506-8Report Type:Periodic Company Report #001-0945-M0200360
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/16/02ISR Number: 4012508-1Report Type:Periodic Company Report #001-0945-M0200388
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Hypoventilation	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 MG (400 Other MG, TID),							
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4012509-3Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-M0200439

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Agitation Anxiety Arthralgia Arthropathy Coordination Abnormal Gait Disturbance Headache Insomnia Intentional Misuse Medication Error Migraine	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:10/16/02ISR Number: 4012511-1Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-M0200455

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death	Death Intentional Misuse	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/16/02ISR Number: 4012512-3Report Type:Periodic
 Age:31 YR Gender:Male I/FU:I

Company Report #001-0945-M0200457

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (300 Other MG, THREE TIMES DFAILY) ,ORAL 40 MG	Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Citalopram Hidrobromide	SS		ORAL

(DAILY), ORAL

Date:10/16/02ISR Number: 4012514-7Report Type:Periodic Company Report #001-0945-M0200478
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Consumer	Neurontin			
		Insomnia		(Gabapentin	PS		
		Intentional Misuse		Carisoprodol	C		
		Pain		Glibenclamide	C		
		Syncope					

Date:10/16/02ISR Number: 4012516-0Report Type:Periodic Company Report #001-0945-M0200505
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bradycardia	Health	Neurontin			
			Professional	(Gabapentin)	PS		
				Quetiapine	SS		
				Paroxetine			
				Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4012517-2Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-M0200519

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (TID), ORAL	Nephrolithiasis	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:10/16/02ISR Number: 4012519-6Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #001-0945-M0200587

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Stevens-Johnson Syndrome	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/16/02ISR Number: 4012520-2Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-M0200597

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 400 MG , ORAL	Blood Creatine Phosphokinase Increased Death	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Pravastatin (Pravastatin)	SS		
			Ciclosporin	C		
			Azathioprine	C		
			Hydrochlorothiazide	C		
			Diltiazem	C		
			Furosemide	C		
			Captopril	C		
			Isodril	C		
			Tocopherol	C		
			Ascorbic Acid	C		

Date:10/16/02ISR Number: 4012521-4Report Type:Periodic
Age:76 YR Gender:Female I/FU:I

Company Report #001-0945-M0200644

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Constipation	Consumer Health	Neurontin (Gabapentin	PS		ORAL
900 MG (300 MG, TID), ORAL		Dizziness Hyperglycaemia Hypertension Insomnia Pain	Professional	Soft Stool Methylcellulose Senakot-S Alendroante Sodium Warfarrin Sodium Triamcinolone Ointment Captopril Vicodin	C C C C C C C C		

Date:10/16/02ISR Number: 4012522-6Report Type:Periodic Company Report #001-0945-M0200658
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Weight Decreased	Health Professional	Neurontin (Gabapentin	PS		ORAL
ORAL							

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Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4012523-8Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-M0200661

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin			
Other		Amnesia		(Gabapentin	PS		ORAL
200 MG		Hypoaesthesia					
(DAILY), ORAL		Infection					
		Intentional Misuse		Lansoprazole	C		
		Medication Error		Cyanocobalamin	C		
		Oedema Peripheral		Alendronate Sodium	C		
		Osteoporosis		Ergocalciferol	C		
		Thyroid Disorder		Calcium	C		
		Weight Increased					

Date:10/16/02ISR Number: 4012524-XReport Type:Periodic
Age: Gender:I/FU:I

Company Report #001-0945-M0200118

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin			
Death		Death		(Gabapentin)	PS		

Date:10/16/02ISR Number: 4012526-3Report Type:Periodic
Age: Gender:I/FU:I

Company Report #001-0945-M0200119

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin			
Death		Death		(Gabapentin)	PS		

Date:10/16/02ISR Number: 4012527-5Report Type:Periodic
Age: Gender:I/FU:I

Company Report #001-0945-M0200120

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin			
Death		Death		(Gabapentin)	PS		

Date:10/16/02ISR Number: 4012528-7Report Type:Periodic Company Report #001-0945-M0200127
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Dehydration	Health	Neurontin			
Initial or Prolonged	Diarrhoea	Professional	(Gabapentin)	PS		
Other	Leukopenia					
	Oedema Peripheral					

Date:10/16/02ISR Number: 4012529-9Report Type:Periodic Company Report #001-0945-M0200164
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Death	Health	Neurontin			
		Professional	(Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4012530-5Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #001-0945-M0200188

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 4800 MG (1600 MG, TID), ORAL	Cardiac Failure Intentional Misuse Medication Error	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Bumetanide	C
Captopril	C
Meformin	
Hydrochloride	C
Digoxin	C
Levothyroxine Sodium	C
Mirtazapine	C
Morphine Sulfate	C
Welchol (Colesevelam Hydrochloride)	C
Unspecified Medications As Needed	C

Date:10/16/02ISR Number: 4012532-9Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-M0200235

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other ORAL	Hyperhidrosis Renal Impairment Weight Decreased	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:10/16/02ISR Number: 4012534-2Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #001-0945-M0200236

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Convulsion Muscle Twitching	Consumer	Neurontin (Gabapentin)	PS		

Tongue Disorder

Sertraline
Hydrochloride
(Sertraline
Hydrochloride) SS

Date:10/16/02ISR Number: 4012535-4Report Type:Periodic
Age:75 YR Gender:Female I/FU:I

Company Report #001-0945-M0200246

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pain	Health	Neurontin			
		Retinal Disorder	Professional	(Gabapentin)	PS		ORAL
ORAL							

Amitriptyline
Hydrochloride
(Amitriptyline) SS
Estrogens Conjugated C
Nitrofurantoin C
Polysaccharide-Iron
Complex C
Calcium Carbonate C
Calcitonin, Salmon C
Oxybutynin C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4012537-8Report Type:Periodic
Age:54 YR Gender:Female I/FU:I

Company Report #001-0945-M020253

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Insomnia	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Oedema					
		Pharyngitis					
900 MG (300 MG, TID PRN), ORAL							
				Sertraline Hydrochloride	C		
				Estradiol	C		

Date:10/16/02ISR Number: 4012931-5Report Type:Periodic
Age:52 YR Gender:Male I/FU:I

Company Report #001-0945-M0200672

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Bradycardia					
		Chest Pain					
600 MG (BID), ORAL							
		Diplopia		Tylenol Pm	C		
		Dizziness					
		Hypotension					

Date:10/16/02ISR Number: 4012935-2Report Type:Periodic
Age:56 YR Gender:Female I/FU:I

Company Report #001-0945-M0200682

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/16/02ISR Number: 4012938-8Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-M0200706

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health	Neurontin			
Other		Pancreatitis	Professional	(Gabapentin)	PS		

Date:10/16/02ISR Number: 4012940-6Report Type:Periodic Company Report #001-0945-M0200719
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Asthenia Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL		Emotional Disorder Gait Disturbance Headache Increased Appetite Intentional Misuse Neuropathy Peripheral Oedema Peripheral Pain Somnolence Speech Disorder Visual Disturbance Weight Increased		Didanosine Nevirapine Stavudine Clonazepam Warfarin Sodium Metoprolol Trandolapril Omeprazole	C C C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4012943-1Report Type:Periodic
Age: Gender: I/FU:I

Company Report #001-0945-M0200732

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	3200 MG (800 Other MG, FOUR TIMES DAILY)	Confusional State Encephalopathy	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Intentional Misuse					
		Medication Error					
							ORAL

Date:10/16/02ISR Number: 4012946-7Report Type:Periodic
Age:39 YR Gender:Female I/FU:I

Company Report #001-0945-M0200732

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	100 MG (DAILY), ORAL	Amnesia Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Paranoia					
		Speech Disorder		Atenolol Atorvastatin	C C		

Date:10/16/02ISR Number: 4012950-9Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #001-0945-M0200738

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Level Increased Medication Error Renal Failure Acute	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/16/02ISR Number: 4012954-6Report Type:Periodic
Age:64 YR Gender:Female I/FU:I

Company Report #001-0945-M0200741

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Hypertension Paraesthesia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Speech Disorder	Professional	Nicotinic Acid	C		
					Potassium	C		
					Calcium	C		
					Clonidine	C		
					Hydrochlorothiazide	C		
					Lotrel	C		
					Acetylsalicylic Acid	C		
					Insulin	C		
					Meclozine	C		

Date:10/16/02ISR Number: 4012958-3Report Type:Periodic Company Report #001-0945-M0200751
Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Convulsion	Consumer	Neurontin (Gabapentin)	PS		
Other			Renal Impairment		Topiramate	SS		
			Unevaluable Event		Valproate Semisodium	C		
			Urinary Tract Infection					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4012960-1Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #001-0945-M0200756

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea	Consumer	Neurontin (Gabapentin)	PS		
1800 MG,		Neuralgia					
(TID),		Rectal Haemorrhage					
ORAL		Renal Failure Acute		Spirolactone	SS		ORAL
				Acetylsalicylic Acid	SS		
				Furosemide	C		
				Trazodone	C		
				Carvedilol	C		
				Lisinopril	C		
				Simvastatin	C		
				Insulin	C		

Date:10/16/02ISR Number: 4012966-2Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2002050042

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intentional Misuse	Health	Neurontin			
2700 MG		Medication Error	Professional	(Gabapentin)	PS		
		Muscle Twitching	Company				
		Renal Failure Acute	Representative				
		Tremor					

Date:10/16/02ISR Number: 4012968-6Report Type:Periodic
Age:74 YR Gender:Female I/FU:I

Company Report #2002050045

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cataract	Consumer	Neurontin			
Initial or Prolonged		Hypercholesterolaemia		(Gabapentin)	PS		ORAL
400 MG (200		Pain					
MG, TWICE							

DAILY), ORAL

10 MG

(DAILY), ORAL

Atorvastatin SS ORAL

Latanoprost	C
Repaglinide	C
Metformin	
Hydrochloride	C
Loratadine	C
Pirbuterol Acetate	C
Budesonide	C
Naproxen Sodium	C

Date:10/16/02ISR Number: 4012970-4Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2002050088

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cataract	Health	Neurontin			
Other		Eye Disorder	Professional	(Gabapentin)	PS		ORAL
ORAL				(Darvocet)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4012973-XReport Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2002050429

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
		Intentional Misuse		(Gabapentin)	PS		ORAL
		Neoplasm					
900 MG (TID),							
ORAL							
				Phenytoin Sodium	C		
				Mirtazapine	C		
				Pentoxifylline	C		
				Acetylsalicylic Acid	C		
				Tocopherol	C		
				Potassium	C		

Date:10/16/02ISR Number: 4012980-7Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2002050495

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Disorder	Consumer	Neurontin			
		Depression		(Gabapentin)	PS		ORAL
100 MG							
(DAILY), ORAL							
		Hypoglycaemia					
		Hypokinesia		Rosiglitazone			
		Insomnia		Maleate	SS		
		Pain		Glibenclamide	C		
		Weight Increased		Losartan Potassium	C		
				Oxycocet	C		

Date:10/16/02ISR Number: 4012985-6Report Type:Periodic
Age:29 YR Gender:Female I/FU:I

Company Report #2002051085

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Disorder	Consumer	Neurontin			
		Hypoaesthesia		(Gabapentin)	PS		ORAL
ORAL							
		Myalgia		Birth Control Pills	SS		
		Pain		Vicodin	C		

Unintended Pregnancy

Date:10/16/02ISR Number: 4012987-XReport Type:Periodic
 Age:41 YR Gender:Female I/FU:I

Company Report #2002051128

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (200 MG, THREE TIMES DAILY), ORAL		Ketosis Nausea Pancreatitis Somnolence	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
INTRAVENOUS MG, NINE TIMES DAILY), INTRAVENOUS	13.5 MG (1.5			Epoprostenol Sodium	SS		
				Furosemide	C		
				Warfarin Sodium	C		
				Spirolactone	C		
				Esomeprazole	C		
				Amitriptyline Hydrochloride	C		
				Sertraline Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4012990-XReport Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #2002051146

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (TID), ORAL		Intentional Misuse Pain Syncope	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Salbutamol Ipratropium Bromide	C C		

Date:10/16/02ISR Number: 4012991-1Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2002051282

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (THREE TIMES Other DAILY)		Convulsion	Consumer	Neurontin (Gabapentin)	PS		
				Tocopherol Calcium	SS SS		

Date:10/16/02ISR Number: 4012993-5Report Type:Periodic
 Age:77 YR Gender:Female I/FU:I

Company Report #2002051328

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), ORAL		Cerebrovascular Disorder Depersonalisation	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Rabeprazole Celecoxib Clopidogrel Potassium Chloride Furosemide Losartan Potassium Alendronate Sodium	C C C C C C C		

Famotidine C
Benzonatate C

Date:10/16/02ISR Number: 4012997-2Report Type:Periodic Company Report #2002051611
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Neurontin (Gabapentin) Unspecified Other Drugs	PS C		

Date:10/16/02ISR Number: 4013000-0Report Type:Periodic Company Report #2002052152
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 300 MG (DAILY), ORAL		Asthenia Labile Blood Pressure	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Alendronate Sodium	C		
				Atenolol	C		
				Clopidogrel	C		
				Alprazolam	C		
				Atorvastatin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Olopatadine
Hydrochloride C
Raloxifene
Hydrochloride C

Date:10/16/02ISR Number: 4013002-4Report Type:Periodic Company Report #2002052179
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fracture	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Nausea					
ORAL							

Date:10/16/02ISR Number: 4013005-XReport Type:Periodic Company Report #2002052304
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Disorder	Health	Neurontin (Gabapentin)	PS		ORAL
		Coordination Abnormal	Professional				
1800 MG (600							
MG, THREE							
TIMES DAILY),							
ORAL							
				Irbesartan	C		
				Atenolol	C		
Hyperlipidaemia							
Osteoporosis							
Visual Disturbance							

Date:10/16/02ISR Number: 4013007-3Report Type:Periodic Company Report #2002052623
Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Hypertension	Health	Neurontin (Gabapentin)	PS		ORAL
Other			Professional				
600 MG (TID),							
ORAL							

Morphine	C
Olsalazine	C
Promethazine	C
Rabeprazole	C
Senna	C
Lorazepam	C

Date:10/16/02ISR Number: 4013009-7Report Type:Periodic Company Report #2002052813
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pneumonia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1800 MG							
(TID), ORAL							
				Seretide Mite	C		
				Gatifloxacin	C		
				Lansoprazole	C		
				Metoclopramide	C		
				Amitriptyline Hydrochloride	C		
				Celecoxib	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4013012-7Report Type:Periodic
Age:59 YR Gender:Male I/FU:I

Company Report #2002053406

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Consumer	Neurontin			
		Asthma		(Gabapentin)	PS		ORAL
		Back Pain					
		Dysgeusia		Oxycodone			
		Headache		Hydrochloride	SS		
		Parosmia		Nifedipine	C		
		Somnolence		Valdecocixib	C		

Date:10/16/02ISR Number: 4013014-0Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2002053813

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tremor	Consumer	Neurontin			
				(Gabapentin)	PS		ORAL
				Oxybutynin	C		
				Quetiapine Fumarate	C		
				Nefazodone			
				Hydrochloride	C		
				Trihexyphenidyl	C		

Date:10/16/02ISR Number: 4013016-4Report Type:Periodic
Age:26 YR Gender:Female I/FU:I

Company Report #2002053863

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Breast Neoplasm	Consumer	Neurontin			
		Galactorrhoea		(Gabapentin)	PS		ORAL
		Headache					
		Visual Disturbance		Bupropion			
				Hydrochloride	C		

Oxycodone
Hydrochloride C

Date:10/16/02ISR Number: 4013017-6Report Type:Periodic Company Report #2002053912
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ear Disorder	Health	Neurontin			
		Intentional Misuse	Professional	(Gabapentin)	PS		
800 MG (THREE							
TO FOUR TIMES		Medication Error					
DAILY)		Retinal Haemorrhage					
				Lithium Carbonate	C		
				Fluvoxamine Maleate	C		
				Trazodone	C		

Date:10/16/02ISR Number: 4013018-8Report Type:Periodic Company Report #001-0945-M0100462
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin			
Other				(Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4013020-6Report Type:Periodic
Age: Gender: I/FU:F

Company Report #001-0945-M0100463

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							

Date:10/16/02ISR Number: 4013025-5Report Type:Periodic
Age: Gender: I/FU:F

Company Report #001-0945-M0100464

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							

Date:10/16/02ISR Number: 4013027-9Report Type:Periodic
Age: Gender: I/FU:F

Company Report #001-0945-M0100465

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							

Date:10/16/02ISR Number: 4013029-2Report Type:Periodic
Age: Gender: I/FU:F

Company Report #001-0945-M0100466

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							

Date:10/16/02ISR Number: 4013031-0Report Type:Periodic
Age: Gender: I/FU:F

Company Report #001-0945-M0100467

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							

Date:10/16/02ISR Number: 4013033-4Report Type:Periodic Company Report #001-0945-M0100468
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin	PS		
Other				(Gabapentin)			

Date:10/16/02ISR Number: 4013040-1Report Type:Periodic Company Report #001-0945-M0100469
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin	PS		
Other				(Gabapentin)			

Date:10/16/02ISR Number: 4013041-3Report Type:Periodic Company Report #001-0945-M0100470
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin	PS		
Other				(Gabapentin)			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/16/02ISR Number: 4013042-5Report Type:Periodic Company Report #001-0945-M0100471

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							

Date:10/17/02ISR Number: 3995803-1Report Type:Expedited (15-DaCompany Report #NSADSS2002032637

Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide	Literature Health	Fentanyl (Patch) (Fentanyl)	PS		
INGESTION - INTENTIONAL SUICIDE		Pallor	Professional				
INTENTIONAL SUICIDE		Somnolence		Propoxyphene (Dextropropoxyphene)	SS		
INTENTIONAL SUICIDE				Gabapentin (Gabapentin)	SS		

Date:10/17/02ISR Number: 3995828-6Report Type:Expedited (15-DaCompany Report #NSADSS2002032666

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Enzymes Increased Completed Suicide Coordination Abnormal	Literature Health Professional	Ultram (50 Mg Tablet) (Tramadol Hydrochloride)	PS		ORAL
ORAL		Cyanosis Dehydration		Gabapentin (Gabapentin)	SS		ORAL
ORAL							

ORAL	Electromechanical	Rofecoxib	SS	ORAL
	Dissociation			
	Heart Rate Decreased			
	Hypotension			
	Mental Status Changes			
	Myoglobin Urine Present			
	Renal Failure			
	Somnolence			
	White Blood Cell Count			
	Increased			

Date:10/17/02ISR Number: 3996436-3Report Type:Expedited (15-DaCompany Report #2002056050

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Cold Sweat	Consumer	Neurontin			
Other		Difficulty In Walking		(Gabapentin)	PS		ORAL
3600 MG (1200		Insomnia					
MG , THREE		Medication Error					
TIMES), ORAL		Movement Disorder		Rofecoxib	C		
		Pain		Enalapril	C		
				Fluoxetine			
				Hydrochloride	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/17/02ISR Number: 3997309-2Report Type:Expedited (15-DaCompany Report #2002060915

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Disorder	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Disability		Convulsion					
1200 MG, ORAL							
Other		Hemiparesis	Professional	Glimepiride	C		
				Carvedilol	C		
				Furosemide	C		
				Trimipramine	C		

Date:10/17/02ISR Number: 3997310-9Report Type:Expedited (15-DaCompany Report #2002061334

Age:91 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dermatitis Bullous	Foreign Health	Neurontin (Gabapentin)	PS		
900 MG (300							
MG, TID)			Professional				
			Company Representative	Acetylsalicylate Lysine	C		

Date:10/17/02ISR Number: 3997315-8Report Type:Expedited (15-DaCompany Report #002#2#2002-00109

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia	Consumer	Reglan (Metoclopramide Hcl)	PS		ORAL
10 MG, 4 IN 1							
Disability		Blood Pressure Increased					
D, ORAL							
		Bruxism		Prochlorperazine -			
		Conversion Disorder		Edisylate	SS		
		Dizziness		Haloperidol	SS		
		Dystonia		Gabapentin	SS		
		Eye Disorder		Olanzapine	SS		
		Feeling Abnormal		Lisinopril	C		
		Grimacing		Diazepam	C		

Haemangioma
Halo Vision
Movement Disorder
Muscle Spasms
Nausea
Nervousness
Nightmare
Pain
Paraesthesia
Photopsia
Photosensitivity Reaction
Restlessness
Strabismus
Tardive Dyskinesia

Mylanta C
Metoprolol C
Heparin C
Clopidogrel C

Date:10/17/02ISR Number: 3997322-5Report Type:Expedited (15-DaCompany Report #2002061107
Age: Gender:Female I/FU:I

Outcome PT
Other Alopecia
Arthralgia
Arthritis
Blood Cholesterol
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					
		Carpal Tunnel Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
1600 MG (FOUR TIMES DAILY), ORAL		Constipation Fibromyalgia					
		Gingival Abscess					
		Gingival Recession					
10 MG (DAILY), ORAL		Hypothyroidism Insomnia		Lipitor (Atorvastatin)	SS		ORAL
		Muscle Disorder					
		Muscle Tightness		Lorazepam	C		
		Myalgia		Bactrim	C		
		Pain In Extremity		Paracetamol	C		
		Stress					

Date:10/18/02ISR Number: 3996341-2Report Type:Expedited (15-DaCompany Report #2002053866
Age:70 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Diplopia	Foreign Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG (300 MG, TID), ORAL			Health Professional				
				Torasemide	C		
				Co-Diovan	C		
				Simvastatin	C		
				Doxepin			
				Hydrochloride	C		
				Phenprocoumon	C		
				Digoxin	C		

Date:10/18/02ISR Number: 3996351-5Report Type:Expedited (15-DaCompany Report #2002056561
Age:2.5 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Complications Of Maternal Exposure To Therapeutic Drugs Constipation Developmental Coordination Disorder Hyperhidrosis Maternal Drugs Affecting Foetus Muscular Weakness Myasthenic Syndrome Temperature Intolerance	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:10/18/02ISR Number: 3996353-9Report Type:Expedited (15-DaCompany Report #2002057364
Age:73 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Erectile Dysfunction Fatigue Tremor	Foreign Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
3600 MG (DAILY), ORAL				Diuretics Acetylsalicylic Acid Losartan Potassium	C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Bisoprolol C

Date:10/18/02ISR Number: 3997600-XReport Type:Expedited (15-DaCompany Report #2002061738
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Diabetes Mellitus Inadequate Control	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG, TID), ORAL		Diarrhoea	Professional				

Insulin Human SS

Date:10/18/02ISR Number: 3998183-0Report Type:Expedited (15-DaCompany Report #2002053218
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hypersensitivity Panic Attack	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG (TID), ORAL			Professional				

Clonazepam C
Bupropion C
Hydrochloride C
Estrogens Conjugated C
Sertraline C
Hydrochloride C

Date:10/18/02ISR Number: 3998184-2Report Type:Expedited (15-DaCompany Report #001-0945-990998
Age:49 YR Gender:Female I/FU:F

Outcome	PT
Disability	Abdominal Pain
Other	Abdominal Rigidity
	Abnormal Faeces
	Alopecia

Anorexia
Anorgasmia
Arthralgia
Back Pain
Bone Pain
Chest Pain
Chills
Contusion
Coordination Abnormal
Cystitis
Dissociation
Disturbance In Attention
Dizziness
Drug Toxicity
Emotional Disorder
Escherichia Infection
Fall
Fatigue
Flatulence
Gastrointestinal Pain
Gingival Bleeding
Hypertonia
Insomnia
Joint Stiffness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nasopharyngitis Nausea Pain	Health Professional	Neurontin (Gabapentin)	PS		ORAL
SEE IMAGE		Pyrexia Reaction To Medical Agent					
		Preservatives Retching Sexual Assault Victim Suffocation Feeling Thirst Urinary Retention Urinary Tract Infection Vomiting		Oxycodone Cyclobenzaprine Hydrochloride Vicodin Diazepam Baclofen	C C C C C		

Date:10/21/02ISR Number: 3997799-5Report Type:Expedited (15-DaCompany Report #2002061162
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Potassium Decreased Fall	Consumer	Neurontin (Gabapentin)	PS		ORAL
DOSE STRENGHT 1200MG (QID), ORAL		Frequent Bowel Movements Gait Disturbance					
		Pollakiuria Swelling Weight Decreased		Indapamide Diuretics Verapamil Hydrochloride Vicodin Nabumetone Amitriptyline Hydrochloride	C C C C C C		

Date:10/21/02ISR Number: 3998511-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Acute Sinusitis Anxiety

Disability
Other

Back Pain
Bladder Disorder
Blood Alkaline
Phosphatase Increased
Bronchitis Acute
Carotid Artery Disease
Cholelithiasis
Cognitive Disorder
Colitis
Computerised Tomogram
Abnormal
Convulsion
Depression
Diverticulum Intestinal
Ear Pain
Ependymoma
Fatigue
Feeling Abnormal
Gastrointestinal Disorder
Hepatic Steatosis
Hyperaemia
Hyperhidrosis

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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		Memory Impairment Monocyte Count Increased Obstructive Airways Disorder Oedema Peripheral	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Plantar Fasciitis Sensory Disturbance Somnolence Tremor Ultrasound Doppler Abnormal Weight Decreased	Professional Company Representative				

Date:10/21/02ISR Number: 3998548-7Report Type:Expedited (15-DaCompany Report #2002052313

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	600 MG (300	Asthenia Carpal Tunnel Syndrome	Consumer Health	Neurontin (Gabapentin)	PS		
Other (MG, BID)		Cerebrovascular Accident	Professional				
		Crying Depression Diabetic Amyotrophy Dizziness Haemorrhage Hemiparesis Hypoaesthesia Insomnia Nephrolithiasis Pain Paraesthesia Procedural Complication		Warfarin Sodium Acetylsalicylic Acid Clonidine Potassium Chloride Furosemide Digoxin Metoprolol Succinate Captopril Isosorbide Mononitrate Capsaicin	SS C C C C C C C C C		

Date:10/21/02ISR Number: 3998551-7Report Type:Expedited (15-DaCompany Report #2002052184

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Surgery	Consumer	Neurontin	
		Health	(Gabapentin)	PS
		Professional		

Date:10/21/02ISR Number: 3998553-0Report Type:Expedited (15-DaCompany Report #2002061355
Age:52 YR Gender:Male I/FU:I

Outcome	PT
Other	Abdominal Pain Lower
	Adenoma Benign
	Anal Fissure
	Circulatory Collapse
	Cold Sweat
	Colitis Ischaemic
	Diarrhoea
	Drug Tolerance Decreased
	Excoriation
	Faeces Hard
	Gastrointestinal
	Infection
	Haematochezia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Influenza Like Illness Intestinal Ischaemia Large Intestinal Ulcer					
600 MG (TID), ORAL		Loss Of Consciousness Malaise Nausea	Health Professional	Gabapentin (Gabapentin)	PS		ORAL
6 MG (TID), ORAL		Rectal Haemorrhage Renal Cyst		Cilansetron	SS		ORAL
50 MCG (BID), ORAL		Skin Discolouration Syncope		Clonidine	SS		ORAL
1000 MG, ORAL		Upper Respiratory Tract		Lemsip	SS		ORAL
2400 MG, ORAL		Infection		Acetylsalicylic Acid	SS		ORAL
				Paracetamol	C		
				Omeprazole	C		
				Doxepin	C		

Date:10/21/02ISR Number: 3999906-7Report Type:Expedited (15-DaCompany Report #2002129159FR
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour Drug Toxicity Gastrointestinal Perforation Visual Disturbance	Foreign Health Professional Other	Celebrex (Celecoxib) Capsule Neurontin (Gabapentin) Anti Ulcerous Treatment	PS SS C		

Date:10/21/02ISR Number: 4005134-1Report Type:Expedited (15-DaCompany Report #2002061334
Age:90 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Dermatitis Bullous	Foreign	Neurontin			

Initial or Prolonged	Epilepsy	Health	(Gabapentin)	PS
900 MG (300				
Other		Professional		
MG, TID)				
		Company	Acetylsalicylate	
		Representative	Lysine	C
			Pentoxifylline	C
			Sinemet	C

Date:10/23/02ISR Number: 3999397-6Report Type:Expedited (15-DaCompany Report #2002062065
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Cholestasis	Foreign	Neurontin			
Initial or Prolonged		Health	(Gabapentin)	PS		
		Professional	Phenobarbital	C		
		Company				
		Representative				

Date:10/23/02ISR Number: 3999411-8Report Type:Expedited (15-DaCompany Report #2002061740
 Age: Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Arthritis Infective
Initial or Prolonged	Drug Ineffective
Other	Haematoma
	Muscle Spasms

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Post Procedural Complication Rash Papular	Report Source	Product	Role	Manufacturer	Route
(TID), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Carisoprodol	SS		
				Omeprazole	C		
				Mirtazapine	C		
				Alprazolam	C		
				Eye Drops	C		

Date:10/23/02ISR Number: 3999521-5Report Type:Expedited (15-DaCompany Report #2002062009
Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (TID), ORAL		Angioneurotic Oedema Conjunctivitis Drug Hypersensitivity Dysphagia Dysphonia	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:10/23/02ISR Number: 3999579-3Report Type:Expedited (15-DaCompany Report #2002058031
Age:82 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG Other (TID), ORAL		Convulsion Dysphagia Fatigue Feeling Cold Myocardial Infarction Pneumonia Stress Vomiting	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Simvastatin	C		
				Atenolol	C		
				Estrogens Conjugated	C		
				All Other Therapeutic Products	C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Depression Drug Dependence Drug Withdrawal Syndrome	Consumer Other	Oxycontin Tablets(Oxycodone Hydrochloride) Cr Tablet			ORAL
ORAL		Pain Pneumonia Aspiration Respiratory Failure		Oxycontin Tablets 20 Mg(Oxycodone Hydrochloride) Cr Tablet	PS		ORAL
20 MG, ORAL;							
40 MG, ORAL							
ORAL				Cyclobenzaprine(Cycl obenzaprine) Tablet	SS		ORAL
ORAL				Hydroxyzine Pamoate(Hydroxyzine Embonate)	SS		ORAL
ORAL				Valium(Diazepam) Tablet	SS		ORAL
ORAL	7 DAY			Neurontin(Gabapentin) Tablet	SS		ORAL
ORAL	7 DAY			Tegretal(Carbamazepi ne) Tablet	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL	7	DAY	Celexa (Citalopram Hydrobromide) Tablet	SS	ORAL
ORAL	7	DAY	Sonata(Zaleplon) Tablet	SS	ORAL
ORAL	7	DAY	Trazodone(Trazodone) Tablet	SS	ORAL
7	DAY		Lorcet (Paracetamol Hydrocodone Bitartrate)	SS	

Date:10/25/02ISR Number: 4000521-XReport Type:Expedited (15-DaCompany Report #2002062012
 Age:56 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Adrenal Insufficiency	Consumer	Neurontin			
Initial or Prolonged	Blood Chloride Decreased		(Gabapentin)	PS		ORAL
900 MG (300	Blood Potassium Decreased					
MG, TID),	Blood Sodium Decreased					
ORAL	Oedema Peripheral		Metolazone	SS		
	Pain In Extremity		Citalopram			
	Pneumonia		Hydrobromide	C		
			Estrogens Conjugated	C		
			Diazepam	C		
			Furosemide	C		
			Panadeine Co	C		
			Hydrocortisone	C		
			Fexofenadine			
			Hydrochloride	C		
			Multivitamins	C		
			Calcium	C		
			Potassium	C		

Date:10/28/02ISR Number: 4000827-4Report Type:Expedited (15-DaCompany Report #2002054897
 Age:90 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Agitation Blood Pressure Systolic	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other 1000 MG		Increased Clostridium Colitis	Professional	Metronidazole	SS		ORAL
(QID), ORAL		Confusional State Drug Interaction Infection Leg Amputation Pain Paranoia Somnolence		Glibenclamide Chromagen Prednisone	C C C		

Date:10/28/02ISR Number: 4000828-6Report Type:Expedited (15-DaCompany Report #2002061973
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG		Aphasia Feeling Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
(ONCE), ORAL		Syncope Tongue Oedema		Paroxetine Hydrochloride Levothyroxine Sodium	C C		

Freedom Of Information (FOI) Report

Metoprolol Succinate C
 Propacet C

Date:10/28/02ISR Number: 4001193-0Report Type:Expedited (15-DaCompany Report #2002054696
 Age:39 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 800 MG (DAILY), ORAL	Cellulitis Convulsion Drug Level Increased	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
400 MG (DAILY) ORAL	Liver Function Test Abnormal Staphylococcal Infection		Dilantin (Phenytoin Sodium)	SS		ORAL
25 MG (DAILY)			Lamotrigine	SS		
			Paracetamol	SS		
			Venlafaxine Hydrochloride	C		
			Atorvastatin	C		
			Benazepril Hydrochloride	C		
			Clonazepam	C		
			Obetrol	C		
			Quetiapine Fumarate	C		
			Mirapex (Pramipexole	C		
			Eptacog Alfa	C		
			Folic Acid	C		
			Pyridoxine Hydrochloride	C		
			Cyanocobalamin	C		

Date:10/28/02ISR Number: 4002601-1Report Type:Expedited (15-DaCompany Report #2002062558
 Age:35 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization -	Coma	Foreign	Neurontin			

Initial or Prolonged 2700 MG (TID)	Confusional State	Consumer	(Gabapentin)	PS	ORAL
ORAL	Hypoaesthesia				
ORAL	Paraesthesia		Morphine	SS	ORAL

Date:10/29/02ISR Number: 4001561-7Report Type:Expedited (15-DaCompany Report #2002053982
Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Areflexia	Literature	Gabapentin			
Hospitalization - 2700 MG (TID)		Blood Creatine	Health	(Gabapentin)	PS		
Initial or Prolonged		Phosphokinase Abnormal	Professional	Estrogens Conjugated	C		
		Coma		Furosemide	C		
		Corneal Reflex Decreased		Levothyroxine	C		
		Drug Level Changed		Lisinopril	C		
		Electroencephalogram Abnormal		Fluoxetine	C		
		Pupil Fixed					
		Status Epilepticus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/29/02ISR Number: 4002072-5Report Type:Expedited (15-DaCompany Report #2002062643

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erectile Dysfunction	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID),		Memory Impairment					
ORAL		Mental Impairment					
		Weight Increased		Carbamazepine Vicodin	SS C		

Date:10/29/02ISR Number: 4002076-2Report Type:Expedited (15-DaCompany Report #2002056416

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		No Adverse Drug Effect	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
2400 MG (600			Professional				
MG, QID),							
ORAL							
				Gemfibrozil	C		
				Atorvastatin	C		
				Guaifenesin	C		
				Allerx-D	C		
				Cetirizine			
				Hydrochloride	C		
				Tamsulosin			
				Hydrochloride	C		
				Bicalutamide	C		
				Morphine Sulfate	C		
				Tramadol			
				Hdyrochloride	C		
				Diazepam	C		
				Celecoxib	C		
				Nefazodone			
				Hydrochloride	C		
				Budesonide	C		
				Mometasone Furoate	C		
				Ipratropium Bromide	C		

Outcome PT
Other Abdominal Distension
Asthenia
Blood Arsenic Increased
Dizziness
Dyspnoea
Feeling Abnormal
Headache
Helicobacter Infection
Hunger
Hypertension
Laboratory Test Abnormal
Muscle Spasms
Nausea
Nervousness
Oesophageal Spasm
Pharmaceutical Product
Complaint
Respiratory Disorder

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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
(TWICE DAILY), ORAL		Consumer Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Professional	Lamotrigine	SS		ORAL
			Clonazepam	SS		

Date:10/29/02ISR Number: 4002151-2Report Type:Expedited (15-DaCompany Report #001-0945-M010093
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams Anxiety	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
SEE IMAGE		Diarrhoea	Professional	Carbamazepine	SS		ORAL
25 MG		Drug Ineffective					
(DAILY), ORAL		Dysgeusia		Levothyroxine Sodium	SS		ORAL
25 MCG		Eye Disorder					
(DAILY), ORAL		Face Oedema Fatigue		Gabapentin Combination	SS		
TOPICAL	TOPICAL	Feeling Abnormal Heart Rate Increased Insomnia Medication Error Nausea Road Traffic Accident Somnolence Thyroid Disorder Tongue Oedema Vision Blurred Vomiting		Cromoglicate Sodium	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	20 MG DAILY	Alanine Aminotransferase	Foreign	Losec	PS		ORAL
PO		Increased	Health				
4 MG DAILY PO	11 MON	Aspartate	Professional	Coumadin "Boots"	SS		ORAL
0.125 MG QD		Aminotransferase	Other	Lanoxin	SS		ORAL
PO		Increased					
100 MG TID PO		Asthenia		Neurontin	SS		ORAL
		Blood Lactate		Cardizem Cd	C		
		Dehydrogenase Increased		Cosopt	C		
		Coagulation Time		Effexor-Xr	C		
		Prolonged		Entrophen	C		
		Confusional State		Flomax "Boehringer			
		Decreased Appetite		Ingelheim"	C		
		Dehydration		Iron	C		
		Drug Level Increased		Lasix	C		
		Dyspnoea		Multivitamins Plus			
		Influenza Like Illness		Iron	C		
		Nephritis Interstitial		Novolin	C		
		Renal Failure Acute		Oscal	C		
				Serax	C		
				Xalatan	C		
				Novolin 70/30	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/29/02ISR Number: 4003698-5Report Type:Expedited (15-DaCompany Report #2002062399
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour	Foreign	Neurontin			
		Gastric Ulcer Perforation	Health	(Gabapentin)	PS		
		Medication Error	Professional	Celecoxib	SS		
		Overdose	Company	Antiulcerous			
		Therapeutic Agent	Representative	Treatment	C		
		Toxicity					
		Visual Disturbance					

Date:10/29/02ISR Number: 4003711-5Report Type:Expedited (15-DaCompany Report #2002062644
Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident	Foreign	Gabapentin			
Hospitalization - 800 MG (BID), Initial or Prolonged ORAL			Consumer	(Gabapentin)	PS		ORAL
				Digoxin	C		
				Pentoxifyline (Pentoxifylline)	C		
				Methadone	C		

Date:10/29/02ISR Number: 4003854-6Report Type:Expedited (15-DaCompany Report #2002062179
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Fall	Consumer	Neurontin			
		Ill-Defined Disorder		(Gabapentin)	PS		

Date:10/29/02ISR Number: 4003881-9Report Type:Expedited (15-DaCompany Report #2002057840
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -
Initial or Prolonged
600 MG (TID),

Abdominal Pain Upper
Meningitis Viral

Foreign
Health

Neurontin
(Gabapentin)

PS

ORAL

ORAL

Professional

Company
Representative

Date:10/29/02ISR Number: 4003885-6Report Type:Expedited (15-DaCompany Report #2002059032

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

Outcome
Hospitalization -
Initial or Prolonged
Other

PT
Apraxia
Balance Disorder
Disturbance In Attention
Epilepsy
Fatigue
Gait Disturbance
Grand Mal Convulsion
Hepatitis C
Hyponatraemia
Memory Impairment
Muscle Spasms
Nausea
Pain

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Post Procedural Complication Vomiting	Report Source	Product	Role	Manufacturer	Route
300 MG TID			Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Oxcarbazepine	SS		
				Amityptilin	C		
				Metamizole Sodium	C		
				Milgamma	C		
				Keltican / Old Form/	C		
				Capsaicin	C		
				Phytodolor	C		
				Emla Plaster	C		
				Lidocaine			
				Hydrochloride	C		

Date:10/29/02ISR Number: 4003886-8Report Type:Expedited (15-DaCompany Report #2002062757
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG BID		Accident Asthenia	Foreign Literature	Gabapentin (Gabapentin)	PS		
		Cholecystitis					
		Face Oedema					
		Medication Error					
		Oedema Peripheral					
		Pulmonary Oedema					
		Somnolence					

Date:10/30/02ISR Number: 4002378-XReport Type:Direct Company Report #CTU 179937
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 600 MG QHS		Anorexia		Gabapentin	PS		

Initial or Prolonged 40 MG QD	Confusional State	Fluoxetine	SS
	Depressed Level Of Consciousness	Fluoxetine	C
	Hallucination	Donepezil	C
		Albuterol	C
		Rabeprazole	C
		Flunisolide	C
		Nitropatch	C
		Perdnisone	C
		Kcl	C
		Mov	C
		Folate	C
		Ipratropium	C
		Calcium	C

Date:10/30/02ISR Number: 4003581-5Report Type:Expedited (15-DaCompany Report #A0376571A
Age: Gender:Female I/FU:F

Outcome PT
Other Arthritis
Bursitis
Cerebrovascular Accident
Double Vessel Bypass
Graft

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Surgery

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Consumer	Sine-Off (Sine-Off)	PS		
			Guaiphenesin (Guaifenesin)	SS		
			Ibuprofen (Ibuprofen)	SS		
			Dimetapp (Dimetapp)	SS		
			Alka-Seltzer Plus Cold Md (Alka-Seltzer Plus Cold Md)	SS		
			Tavist-D (Tavist-D)	SS		
			Triaminic (Triaminic)	SS		
			Phenylpropanolamine Hcl (Phenylpropanolamine)	SS		
			Vicodin (Vicodin)	SS		
			Clonidine (Clonidine)	SS		
			Coricidin (Coricidin)	SS		
			Tylenol Cold Medication (Tylenol Cold Medication)	SS		
			Medrol (Medrol)	SS		
			Rofecoxib (Rofecoxib)	SS		
			Ibuprofen (Ibuprofen)	SS		
			Gabapentin (Gabapentin)	SS		
			Vicks Dayquil (Vicks Dayquil)	SS		
			Nyquil (Nyquil)	SS		

Date:10/31/02ISR Number: 4002141-XReport Type:Direct
Age:44 YR Gender:Female I/FU:I

Company Report #CTU 180052

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization - Generalised Oedema
300 BID ORAL
Initial or Prolonged

Gabapentin Pfizer PS Pfizer ORAL

Date:10/31/02ISR Number: 4003051-4Report Type:Direct Company Report #CTU 180084
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache		Neurontin	PS		ORAL
600 MG BID PO		Medication Error					
		Tremor					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/31/02ISR Number: 4004414-3Report Type:Expedited (15-DaCompany Report #2002062760

Age:87 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anorexia	Foreign	Gabapentin			
Other		Depression	Health	(Gabapentin)	PS		ORAL
(300 MG) ORAL		Headache	Professional	Amiodarone	C		
		Malaise		Acetylsalicylic Acid	C		
		Somnolence		Dipyridamole	C		
		Tremor		Furosemide	C		

Date:10/31/02ISR Number: 4004419-2Report Type:Expedited (15-DaCompany Report #2002062761

Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Alanine Aminotransferase	Foreign	Neurontin			
Initial or Prolonged		Increased	Health	(Gabapentin)	PS		
1200 MG (600		Aspartate	Professional				
MG BID)		Aminotransferase	Company				
		Increased	Representative				
		Blood Alkaline					
		Phosphatase Increased					
		Necrosis					

Date:10/31/02ISR Number: 4004716-0Report Type:Expedited (15-DaCompany Report #2002062742

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Decreased Appetite	Consumer	Norvasc (Amlodipine)	PS		ORAL
5 MG (DAILY),		Fluid Retention					
Initial or Prolonged		Myocardial Infarction		Neurontin			
ORAL		Nerve Compression		(Gabapentin)	SS		ORAL
Other		Osteoporosis					
300 MG							
(DAILY), ORAL							

Date:10/31/02ISR Number: 4004942-0Report Type:Expedited (15-DaCompany Report #2002063526

Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 150 MG DAILY, Initial or Prolonged ORAL		Haematocrit Decreased	Health	Zoloft (Sertraline)	PS		ORAL
Other ORAL		Haemoglobin Decreased	Professional				
		Pain Surgery		Neurontin (Gabapentin)	SS		ORAL

Date:10/31/02ISR Number: 4005192-4Report Type:Expedited (15-DaCompany Report #2002055134

Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (DAILY), ORAL		Feeling Hot Headache	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Company Representative	Fentanyl Methadone Hydrochloride	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/31/02ISR Number: 4005196-1Report Type:Expedited (15-DaCompany Report #2002062771
Age:68 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (300 Other MG, QID) , ORAL	Alcohol Interaction Dysphagia Transient Ischaemic Attack	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL			Ethanol	SS		ORAL
			Valaciclovir Hydrochloride Diclofenac Sodium	C C		

Date:11/01/02ISR Number: 4003240-9Report Type:Direct Company Report #180206
Age:48 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - QWK 2 MON Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage	Chest Discomfort Diarrhoea Dyspnoea Hyperhidrosis Posturing Vision Blurred Vomiting		Cisplatin Premarin Tylenol #3 Vicodin Neurontin	PS C C C I		

Date:11/01/02ISR Number: 4003364-6Report Type:Direct Company Report #CTU 180185
Age:41 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 400 MG, QID, Initial or Prolonged BY MOUTH	Dizziness Dyspnoea Nausea		Gabapentin Acetaminophen Tab Nicotine Transdermal	PS C		ORAL

Patch C
 Lorazepam Tab C
 Flunisolide
 (Aerobid) Oral
 Inhalant I
 Ipratropium Soln,
 Inhl I

Date:11/01/02ISR Number: 4005345-5Report Type:Expedited (15-DaCompany Report #GB9327229OCT2002
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Drug Interaction Somnolence	Health Professional Other	Efexor (Venlafaxine Hydrochloride, Tablet,0)	PS		ORAL
75 MG DAILY							
ORAL				Gabapentin (Gabapentin, 0)	SS		ORAL
300 MG OD -							
TD OVER 3D	7	DAY		Morphine Sulfate (Morphine Sulfate)	C		
				Levothyroxine (Levothyroxine)	C		
				Diclofenac (Diclofenac)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/01/02ISR Number: 4005570-3Report Type:Expedited (15-DaCompany Report #2002AP03531
Age:55 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse Suicide Attempt	Literature Health Professional	Atenolol Diltiazem Gabapentin	PS SS SS		

Date:11/01/02ISR Number: 4005794-5Report Type:Expedited (15-DaCompany Report #2002063465
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3200 MG, ORAL Other		Balance Disorder Influenza Like Illness	Consumer	Neurontin (Gabapentin) Topiramate Rizatriptan Benzoate	PS C C		ORAL

Date:11/01/02ISR Number: 4005795-7Report Type:Expedited (15-DaCompany Report #2002063521
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 3600 MG (THREE TIMES A DAY), ORAL		Hypertension Increased Appetite	Consumer	Neurontin (Gabapentin) Unspecified Psychiatric Medications	PS C		ORAL

Date:11/05/02ISR Number: 4006926-5Report Type:Expedited (15-DaCompany Report #HQ4987731OCT2002
Age:20 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Completed Suicide Intentional Misuse	Literature	Fentanyl Citrate (Fentanyl Citrate, Injection)	PS	ORAL
FETAL					
OVERDOSE,					
AMOUNT					
UNKNOWN, ORAL			Gabapentin (Gabapentin,)	SS	
			Propoxyphene (Dextropropoxyphene,)	SS	

Date:11/06/02ISR Number: 4007371-9Report Type:Expedited (15-DaCompany Report #2002063516
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthropathy Back Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (THREE		Fibromyalgia					
TIMES A DAY),		Hyperhidrosis					
ORAL		Pain In Extremity		Arthrotec Tramadol Hydrochloride	C		
				Hormone Replacement Colesevelam	C		

Freedom Of Information (FOI) Report

Hydrochloride C
 Glibomet C
 Levothyroxine Sodium C

Date:11/06/02ISR Number: 4007383-5Report Type:Expedited (15-DaCompany Report #2002064142
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Convulsion	Literature	Gabapentin	PS		
Other			Health Professional	(Gabapentin)			

Date:11/06/02ISR Number: 4007385-9Report Type:Expedited (15-DaCompany Report #2002064840
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Decreased Activity Lethargy	Literature	Gabapentin	PS		
			Health Professional	(Gabapentin)			

Date:11/06/02ISR Number: 4007650-5Report Type:Expedited (15-DaCompany Report #2002064709
 Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea Exertional Medication Error	Consumer	Neurontin	PS		ORAL
				(Gabapentin)			

600 MG (300

MG. BID),

ORAL

Theophylline C
 Cetirizine
 Hydrochloride C
 Citalopram
 Hydrobromide C
 Prednisone C
 Zafirlukast C

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased	Consumer	Neurontin			
400 MG, ORAL		Drug Ineffective	Health	(Gabapentin)	PS		ORAL
75 MG		Exophthalmos	Professional	Venlafaxine			
(DAILY), ORAL		Eye Pain		Hydrochloride	SS		ORAL
		Fatigue					
		Heart Rate Increased		Atorvastatin	C		
		Hyperhidrosis		Eugynon	C		
		Medication Error		Ranitidine			
		Multiple Sclerosis		Hydrochloride	C		
		Muscle Twitching					
		Pharmaceutical Product					
		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/06/02ISR Number: 4007673-6Report Type:Expedited (15-DaCompany Report #2002064047
Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG Other (DAILY), ORAL	Duration Cartilage Injury Face Oedema Fall Joint Swelling Oedema Peripheral Paraesthesia Somnolence Weight Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Oxycocet	C		
			Diazepam	C		
			Ranitidine	C		
			All Other			
			Therapeutic Products	C		
			Sibutramine			
			Hydrochloride	C		

Date:11/06/02ISR Number: 4008290-4Report Type:Expedited (15-DaCompany Report #2002057364
Age:73 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 3600 MG (DAILY), ORAL	Duration Erectile Dysfunction Fatigue Myalgia Pain In Extremity Tremor	Foreign Consumer Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Diuretics	C		
			Acetylsalicylic Acid	C		
			Losartan Potassium	C		
			Bisoprolol	C		

Date:11/06/02ISR Number: 4008489-7Report Type:Expedited (15-DaCompany Report #2002063981
Age:41 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 400 MG (DAILY), ORAL	Duration Asthenia Paraesthesia	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL

Date:11/06/02ISR Number: 4008561-1Report Type:Expedited (15-DaCompany Report #055-0945-M0100042
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	1200 MG, TID,	Aggression	Foreign	Gabapentin	PS		ORAL
ORAL		Asthenia	Consumer				
		Back Pain		Amitrptyline			
		Condition Aggravated		Hydrochloride	C		
		Constipation		Diazepam	C		
		Depression		Sertraline	C		
		Difficulty In Walking					
		Dyspnoea					
		Inflammation					
		Lesion Of Sciatic Nerve					
		Nasal Congestion					
		Neck Pain					
		Neuralgia					
		Neuropathy Peripheral					
		Pain In Extremity					
		Psychotic Disorder					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/06/02ISR Number: 4008603-3Report Type:Expedited (15-DaCompany Report #2002063979
Age:95 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening 900 MG (THREE TIMES A DAY), ORAL	Blood Pressure Decreased Dizziness Nervousness Oxygen Saturation Decreased Somnolence	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Acetylsalicylic Acid Furosemide Nitrazepam Isosorbide Panadeine Co	C C C C C		

Date:11/07/02ISR Number: 4008991-8Report Type:Expedited (15-DaCompany Report #2002062065
Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (TWICE DAILY), ORAL 100 MG (DAILY), ORAL	Cholestasis Drug Interaction Hepatocellular Damage	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Phenobarbital	PS SS		ORAL ORAL
			Fluindione Phloroglucinol Dipyridamole Clonazepam Baclofen Naftidrofuryl Oxalate	C C C C C C		

Date:11/07/02ISR Number: 4009406-6Report Type:Expedited (15-DaCompany Report #2002064435

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Agitation	Consumer	Neurontin			
Initial or Prolonged	Bipolar Disorder		(Gabapentin)	PS		
	Condition Aggravated		Lamotrigine	C		
	Incontinence		Quetiapine Fumarate	C		
			Metoprolol	C		

Date:11/08/02ISR Number: 4009078-0Report Type:Expedited (15-DaCompany Report #2002064869

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Completed Suicide	Consumer	Neurontin			
Other			(Gabapentin)	PS		

Date:11/12/02ISR Number: 4008028-0Report Type:Direct Company Report #CTU 180735

Age:32 YR Gender:Female I/FU:I

Outcome	PT
Disability	Confusional State
	Coordination Abnormal
	Decreased Activity
	Dizziness
	Drug Level Decreased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG BID		Incoherent Nausea Nervous System Disorder		Neurontin 300mg Parke Davis	PS	Parke Davis	ORAL
ORAL		Pain Somnolence Thinking Abnormal		Neurontin 600mg Parke Davis	SS	Parke Davis	ORAL
600MG TID				Nortriptyline Accupuncture Manipulative Osteopathic Medical Treatment	C C C C		

Date:11/12/02ISR Number: 4008431-9Report Type:Direct
Age:40 YR Gender:Male I/FU:I

Company Report #CTU 180837

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 300MG PO BID		Anxiety		Gabapentin 300mg	PS		ORAL
Hospitalization - ~6 WEEKS	6 WK	Ventricular Tachycardia		Azimilide Or Placebo-Shield Study	SS		ORAL
Initial or Prolonged Required PO ~ 2 WEEKS	2 WK			Toprol Xl Kcl Zocor Norvasc Prilosec Mg Oxidie Multivit Asa	C C C C C C C		
Intervention to Prevent Permanent Impairment/Damage							

Date:11/12/02ISR Number: 4011471-7Report Type:Expedited (15-DaCompany Report #B0273503A
Age:64 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Acute Myocardial Infarction	Foreign Study Health Professional	Gabapentin (Gabapentin) Combivir Tablet (Combivir)	PS		ORAL
1 TABLET					SS		
TWICE PER DAY/ ORAL							
600 MG/ PER DAY/ ORAL				Efavirenz Tablet (Efavirenz)	C		ORAL
				Serevent (Salmeterol Xinafoate)	C		
				Amiodarone	C		
				Carvedilol	C		
				Ramipril	C		
				Frusemide	C		

Date:11/13/02ISR Number: 4010212-7Report Type:Expedited (15-DaCompany Report #2002064772
Age:41 YR Gender:Male I/FU:I

Outcome PT
Other Convulsion
Drug Interaction

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Somnolence

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, THREE TIMES DAILY), ORAL		Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
75 MG (DAILY), ORAL			Venlafaxine	SS		ORAL
			Morphine Sulfate	C		
			Levothyroxine Sodium	C		
			Diclofenac	C		

Date:11/13/02ISR Number: 4010781-7Report Type:Expedited (15-DaCompany Report #2002064849
Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 1200 MG Initial or Prolonged (TID), ORAL		Cardiovascular Disorder Haemorrhage	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
		Respiratory Failure					

Date:11/13/02ISR Number: 4010845-8Report Type:Expedited (15-DaCompany Report #2002065231
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG (600 MG, TID),		Blood Urine Present Nephrolithiasis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Weight Decreased					

ORAL

Date:11/13/02ISR Number: 4011343-8Report Type:Expedited (15-DaCompany Report #2002063022

Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coordination Abnormal Epilepsy	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
3000 MG (600 MG, DAILY)		Medication Error	Professional				

ORAL

Trihexyphenidyl	C
Propranolol	C
Aporex	C

Date:11/13/02ISR Number: 4012036-3Report Type:Expedited (15-DaCompany Report #2002056558

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Convulsion
Initial or Prolonged	Drug Ineffective
Other	Enuresis
	Erectile Dysfunction
	Fatigue
	Feeling Abnormal
	Grand Mal Convulsion
	Insomnia
	Lethargy
	Lip Blister

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Lip Ulceration Listless Myocardial Infarction					
300 MG (TID), ORAL		Rash Pruritic Somnolence	Consumer Health	Dilantin (Phenytoin Sodium)	PS		ORAL
		Stent Placement	Professional				
1200 MG (TID), ORAL				Neurontin (Gabapentin)	SS		ORAL
				Oxcarbazepine	SS		
				Valproate Semisodium	SS		
				Pravastatin Sodium	C		
				Amlodipine Besilate	C		
				Tocopherol	C		
				Acetylsalicylic Acid	C		
				Glucosamine	C		

Date:11/13/02ISR Number: 4012044-2Report Type:Expedited (15-DaCompany Report #2002065066
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (DAILY), ORAL		Carotid Artery Occlusion Cerebrovascular Accident	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Grand Mal Convulsion		Antihypertensives Antidepressants	C C		

Date:11/13/02ISR Number: 4012388-4Report Type:Expedited (15-DaCompany Report #HQ5059005NOV2002
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Drug Interaction Extrapyramidal Disorder Syncope	Health Professional	Efexor Xr (Venlafaxine Hydrochloride, Capsule, Extended			

75 MG 1X PER

Release)

PS

ORAL

1 DAY

Neurontin
(Gabapentin,)

SS

ORAL

400 MG 2X PER

1 DAY

Date:11/14/02ISR Number: 4009021-4Report Type:Expedited (15-DaCompany Report #A0373437A

Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - INTRAVENOUS Initial or Prolonged times per day 5	1.5MG Nine YR	Decreased Activity Hyperosmolar State		Flolan	PS	Glaxo Wellcome	
		International Normalised Ratio Abnormal Loss Of Consciousness Pain In Extremity		Neurontin Furosemide Spironolactone Coumadin Digoxin Oxycontin Nexium	SS C C C C C C	Glaxo Wellcome Glaxo Wellcome Glaxo Wellcome	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/14/02ISR Number: 4012306-9Report Type:Expedited (15-DaCompany Report #EMADSS2002006751
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Somnolence	Foreign Health Professional	Tramal (Unspecified) (Tramadol Hydrochloride)	PS		ORAL
100 MG,							
DAILY, ORAL							
1800 MG,				Neurontin (Gabapentin)	SS		
DAILY							

Date:11/14/02ISR Number: 4012956-XReport Type:Expedited (15-DaCompany Report #2002061973
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dissociation Dysphasia	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG		Feeling Abnormal					
(ONCE), ORAL		Syncope Tongue Oedema		Paroxetine Hydrochloride	C		
				Levothyroxine Sodium	C		
				Metoprolol Succinate	C		
				Propacet	C		

Date:11/14/02ISR Number: 4013285-0Report Type:Expedited (15-DaCompany Report #2002-BP-04878BP (2)
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abdominal Pain Abnormal Faeces Adenoma Benign Circulatory Collapse	Foreign Health Professional Other	Clonidine Hydrochloride Tablets(Clonidine)(T a) (Clonidine-Hcl)	PS		ORAL
SEE IMAGE	18 DAY						

6 MG (2 MG, 2			Cold Sweat		Cilansetron(Nr)	SS	ORAL
MG TID)	44	DAY	Colitis Ischaemic				
			Diarrhoea		Gabapentin(Gabapenti		
600 MG (200			Dizziness		n)(Nr)	SS	ORAL
MG, 200 MG			Haematochezia				
TID)	34	DAY	Loss Of Consciousness				
			Malaise		Paracetamol(Paraceta		
			Protein S Increased		mol)(Nr)	C	
			Shock		Omeprazole(Omeprazol		
			Ulcer		e)	C	
			Upper Respiratory Tract		Doxepin(Doxepin)(Nr)	C	
			Infection		Aspirin(Nr)	C	

Date:11/14/02ISR Number: 4013287-4Report Type:Expedited (15-DaCompany Report #2002065211
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Transient Ischaemic	Foreign	Neurontin			
Other		Attack	Health	(Gabapentin)	PS		ORAL
900 MG , ORAL			Professional	Ethanol	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/18/02ISR Number: 4013267-9Report Type:Expedited (15-DaCompany Report #2002062009
Age:80 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Conjunctivitis	Health	Neurontin			
		Drug Hypersensitivity	Professional	(Gabapentin)	PS		ORAL
		Dysphagia					
		Dysphonia					
		Laryngeal Oedema					

Date:11/18/02ISR Number: 4014018-4Report Type:Expedited (15-DaCompany Report #2002052509
Age:67 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cerebrovascular Accident	Foreign	Neurontin (Tablets)			
Initial or Prolonged		Cerebrovascular Disorder	Consumer	(Gabapentin)	PS		ORAL
400 MG (TID),							
Disability		Nuclear Magnetic	Health				
ORAL							
		Resonance Imaging	Professional	Furosemide	C		
		Abnormal		Verapamil	C		
		Somnolence		Irbesartan	C		
		Vertigo		Moxonidine	C		
				Omeprazole	C		
				Gaviscon	C		
				Asasantin	C		

Date:11/18/02ISR Number: 4014934-3Report Type:Expedited (15-DaCompany Report #033-0945-M0200035
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Epilepsy	Foreign	Neurontin			
			Consumer	(Gabapentin)	PS		
2200 MG							
(DAILY)							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Foreign	Gabapentin			
UNKNOWN	900 MG	Drug Interaction	Literature	(Gabapentin)	PS		
(DAILY),		Drug Level Increased					
UNKNOWN		Electroencephalogram					
UNKNOWN	40 MG	Abnormal		Atorvastatin			
(UNKNOWN),		Eye Movement Disorder		(Atorvastatin)	SS		
UNKNOWN		Fatigue					
UNKNOWN				Carbamazepine	SS		
UNKNOWN				Roxithromycin	SS		
UNKNOWN							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Hernia	Foreign	Neurontin			
ORAL		Fatigue	Consumer	(Gabapentin)	PS		ORAL
		Pain					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/19/02ISR Number: 4015133-1Report Type:Expedited (15-DaCompany Report #2002065565

Age:55 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Gabapentin			
Other		Overdose		(Gabapentin)	PS		ORAL
ORAL				Diltiazem			
				(Diltiazem)	SS		ORAL
ORAL				Atenolol	SS		ORAL

Date:11/19/02ISR Number: 4015137-9Report Type:Expedited (15-DaCompany Report #2002066162

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Rhinorrhoea	Health Professional	Neurontin			
ORAL				(Gabapentin)	PS		ORAL
				Terazosin			
				Hydrochloride	C		
				Diltiazem	C		
				Finasteride	C		
				Oxycocet	C		

Date:11/20/02ISR Number: 4015166-5Report Type:Expedited (15-DaCompany Report #2002066174

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cataract	Health Professional	Neurontin			
		Keratoconjunctivitis		(Gabapentin)	PS		
		Sicca					
		Macular Degeneration					

Date:11/20/02ISR Number: 4015215-4Report Type:Expedited (15-DaCompany Report #2002066580

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (TID) Other		Delirium Hallucination	Consumer	Neurontin (Gabapentin) Famotidine	PS C		

Date:11/20/02ISR Number: 4015674-7Report Type:Expedited (15-DaCompany Report #2002062757
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (BID),		Abdominal Pain Upper Asthenia Chest X-Ray Abnormal Cholecystitis Dyspnoea Eosinophilia Face Oedema Hepatic Enzyme Increased Oedema Peripheral	Foreign Literature	Gabapentin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/20/02ISR Number: 4015752-2Report Type:Expedited (15-DaCompany Report #2002066569

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diplopia	Foreign	Neurontin			
ORAL		Vitreous Detachment	Health	(Gabapentin)	PS		ORAL
			Professional				

Date:11/20/02ISR Number: 4015773-XReport Type:Expedited (15-DaCompany Report #2002066182

Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation	Foreign	Gabapentin			
Hospitalization -		Bronchial Obstruction	Literature	(Gabapentin)	PS		
Initial or Prolonged		Delirium	Health	Amitriptyline	SS		
Other		Drug Interaction	Professional	Paroxetine	SS		
		Dyspnoea		Carbamazepine	SS		
		Eyelid Ptosis		Methadone	SS		
75 MG (25 MG,							
EVERY 8		Stridor					
HOURS)							
				Dexamethasone	SS		
SUBCUTANEOUS	20 MG (10 MG,						
TWICE A DAY),							
SUBCUTANEOUS							
				Midazolam	SS		
SUBCUTANEOUS	240 MG (10 MG						
PER HOUR),							
SUBCUTANEOUS							
				Hydromorphone	SS		
SUBCUTANEOUS	300 MG						
(DAILY),							
SUBCUTANEOUS							

75 MG (25 MG,
 EVERY 8
 HOURS)

Levomepromazine SS
 Phenobarbital C

Date:11/21/02ISR Number: 4016372-6Report Type:Expedited (15-DaCompany Report #2002066596
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (100 MG, TID), ORAL		Anorexia Aptyalism Confusional State Dizziness	Foreign Literature Health Professional	Phenytoin Suspension (Phenytoin Sodium)	PS		ORAL
ORAL		Drug Hypersensitivity Insomnia		Gabapentin (Gabapentin)	SS		ORAL
ORAL		Memory Impairment Pain Rash Somnolence		Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		ORAL
35 MG (BID), ORAL				Fentanyl	SS		
TRANSDERMAL TRANSDERMAL	25 UG/H,			Morphine	SS		ORAL
PRN, ORAL				Oxybutynin Docusate Sodium Lorazepam Sennoside A Glycerol	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/21/02ISR Number: 4016713-XReport Type:Expedited (15-DaCompany Report #2002061738
 Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Diabetes Mellitus	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		Inadequate Control					
ORAL		Diarrhoea	Professional	Insulin Human	C		

Date:11/21/02ISR Number: 4017216-9Report Type:Expedited (15-DaCompany Report #2002066579
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Creatinine Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID),		Cytomegalovirus Infection					
ORAL		Haemodialysis Kidney Transplant Rejection Treatment Noncompliance		All Other Therapeutic Products (All Other Therapeutic Products)	SS		
INTRAVENOUS	INTRAVENOUS			All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:11/22/02ISR Number: 4013668-9Report Type:Direct Company Report #CTU 181528
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Abdominal Pain		Neurontin 500mg	PS		
		Alopecia		Neurontin 900mg	SS		
		Arthralgia					
		Bone Pain					
		Eye Pain					
		Fatigue					

Gastrooesophageal Reflux
Disease
Hypoaesthesia
Lethargy
Mood Swings
Muscle Twitching
Myalgia
Oedema Peripheral
Pain
Paraesthesia
Peripheral Sensory
Neuropathy
Rash
Skin Disorder
Tremor

Date:11/25/02ISR Number: 4014379-6Report Type:Expedited (15-DaCompany Report #A0386384A
Age:42 YR Gender:Male I/FU:I

Outcome PT
Other Agranulocytosis
Bradycardia
Condition Aggravated

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hypotension Thrombocytopenia	2MG	Four times per day		Health Professional	Lamictal Gabitril	PS SS	Glaxo Wellcome	ORAL ORAL
					Neurontin Lithium Sonata Claritin Multivitamins Colace	SS C C C C C	Glaxo Wellcome	ORAL

Date:11/25/02ISR Number: 4014438-8Report Type:Expedited (15-DaCompany Report #WAES 0211USA01406
Age:30 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Drug Interaction Heat Stroke		Cogentin Wellbutrin Neurontin	PS SS SS	Merck & Co., Inc	

Date:11/26/02ISR Number: 4015890-4Report Type:Direct Company Report #CTU 181755
Age:73 YR Gender:Female I/FU:I

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

Date:11/26/02ISR Number: 4017530-7Report Type:Expedited (15-DaCompany Report #2002067329
Age:51 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG			Dizziness Nystagmus	Foreign Health	Neurontin (Gapabentin)	PS		ORAL

(TID), ORAL

Professional

Carbamazepine	C
Atenolol	C
Bendroflumethiazide	C
Amlodipine	C

Date:11/26/02ISR Number: 4017558-7Report Type:Expedited (15-DaCompany Report #2002066599

Age:52 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 (BID), ORAL	Glomerulonephritis	Foreign Health Professional	Gabapentin(Gabapentin) Paracetamol	PS C		ORAL

Date:11/26/02ISR Number: 4018007-5Report Type:Expedited (15-DaCompany Report #2002061973

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

Outcome Other	PT Dizziness Feeling Abnormal Speech 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
300 MG	(ONCE), ORAL	Syncope Tongue Oedema	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Paroxetine Hydrochloride	C		
				Levothyroxine Sodium	C		
				Metoprolol Succinate	C		
				Propacet	C		

Date:11/26/02ISR Number: 4018009-9Report Type:Expedited (15-DaCompany Report #2002066621
Age:62 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 (300 MG, Other TID), ORAL		Abdominal Pain Upper Blood Pressure Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Constipation					
ORAL		Cough		Celecoxib	SS		ORAL
ORAL		Disease Recurrence		Hydromorphone	SS		ORAL
1 MG (DAILY), ORAL		Dizziness		Lorazepam	SS		ORAL
		Dry Mouth					
		Fatigue		Metoprolol Succinate	SS		
		Medication Error		Docusate Sodium	SS		
		Myocardial Ischaemia		Thyroid	C		
		Pain In Extremity					
		Tongue Disorder					
		Visual Disturbance					
		Wheezing					

Date:11/26/02ISR Number: 4018010-5Report Type:Expedited (15-DaCompany Report #2002066595
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Neurontin			
		Dizziness		(Gabapentin)	PS		ORAL
600 MG (BID),		Dyskinesia					
ORAL		Exophthalmos		All Other			
		Eye Movement Disorder		Therapeutic Products	C		
		Gait Disturbance		Propacet	C		
		Hallucination		Clonazepam	C		
		Migraine		Imipramine			
		Paralysis		Hydrochloride	C		
		Trismus		Levothyroxine Sodium	C		
		Visual Disturbance		Hydromorphone			
				Hydrochloride	C		
				Morphine Sulfate	C		
				Midrid	C		
				Cafergot	C		

Date:11/26/02ISR Number: 4018063-4Report Type:Expedited (15-DaCompany Report #2002053431
Age: Gender:Female I/FU:I

Outcome	PT
Disability	Agitation
	Anger
	Anxiety

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG,BID)		Balance Disorder Dysarthria Memory Impairment Mood Swings Nervous System Disorder Neuropathy Peripheral Paranoia Psychotic Disorder Vision Blurred	Health Professional	Neurontin (Gabapentin)	PS		

Date:11/26/02ISR Number: 4018090-7Report Type:Expedited (15-DaCompany Report #2002066912
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (QD) TWICE DAILY IN MORNING		Bronchospasm Disease Recurrence Drug Dependence Dry Mouth Dysarthria Nausea Weight Increased	Consumer	Neurontin (Gabapentin) Lorazepam Paracetamol	PS SS C		

Date:11/27/02ISR Number: 4015823-0Report Type:Expedited (15-DaCompany Report #WAES 0211USA01405
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Convulsion Drug Interaction		Fosamax Gabitril Neurontin	PS SS SS	Merck & Co., Inc	ORAL

Date:11/27/02ISR Number: 4019033-2Report Type:Expedited (15-DaCompany Report #2002067636
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Gait Disturbance Tremor	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL Other				Lotrel	C		
				Metoprolol Succinate	C		
				Fenofibrate	C		
				Clopidogrel Sulfate	C		
				Tramadol,			
				Acetaminophen	C		

Date:11/27/02ISR Number: 4019175-1Report Type:Expedited (15-DaCompany Report #2002067099
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (TID), ORAL		Epilepsy Thrombocytopenia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/27/02ISR Number: 4019176-3Report Type:Expedited (15-DaCompany Report #2002067330

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG Other (DAILY), ORAL	Hypercapnia Somnolence	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
1.5 MG, ORAL			Bromazepam	SS		ORAL

Date:11/27/02ISR Number: 4019319-1Report Type:Expedited (15-DaCompany Report #2002067264

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Intentional Misuse Toxicologic Test Abnormal	Health Professional	Neurontin (Gabapentin)	PS		
			Nefazodone Hydrochloride	SS		
			All Other Therapeutic Products	SS		

Date:11/27/02ISR Number: 4019588-8Report Type:Expedited (15-DaCompany Report #2002067435

Age:56 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG, ORAL	Supraventricular Tachycardia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:11/29/02ISR Number: 4018158-5Report Type:Expedited (15-DaCompany Report #2002060106

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Spinal Fracture	Health Professional	Neurontin (Gabapentin)	PS		ORAL
4800 MG (1600 MG TID) ORAL							

Date:11/29/02ISR Number: 4018176-7Report Type:Expedited (15-DaCompany Report #2002067845
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:11/29/02ISR Number: 4018292-XReport Type:Expedited (15-DaCompany Report #2002067847
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence Hip Fracture	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL Pelvic Fracture Road Traffic Accident							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/29/02ISR Number: 4018295-5Report Type:Expedited (15-DaCompany Report #2002067737

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence	Health Professional	Neurontin (Gabapentin)	PS		
				Benadryl (Diphenhydramine)	SS		

Date:11/29/02ISR Number: 4018350-XReport Type:Expedited (15-DaCompany Report #2002068180

Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Clonic Convulsion	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
2400 MG							
(DAILY), ORAL							
				Codeine	C		
				Paracetamol	C		
				Misoprostol	C		
				Other Anti-Asthmatics, Inhalants	C		

Date:11/29/02ISR Number: 4018351-1Report Type:Expedited (15-DaCompany Report #2002068300

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression Headache	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
1800 MG (ORAL							
		Insomnia		Carbamazepine	C		
				Clonazepam	C		
				Valproate Sodium	C		

Date:11/29/02ISR Number: 4018352-3Report Type:Expedited (15-DaCompany Report #2002057052

Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other		Angioneurotic Oedema Stridor	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG, THREE TIMES DAILY) ORAL			Professional Company Representative				

Baclofen	C
Oxycodone Hydrochloride	C
Fish Oil	C
Prednisolone	C
Ketoprofen	C
Bisacodyl	C
Calcipotriol	C

Date:11/29/02ISR Number: 4018419-XReport Type:Expedited (15-DaCompany Report #2002063521
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Hypertension	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG (TWICE							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

DAILY), ORAL

All Other
 Therapeutic Products C
 Dextropropoxyphene C
 Loratadine C
 Esomeprazole C

Date:11/29/02ISR Number: 4023015-4Report Type:Expedited (15-DaCompany Report #USA-2002-007068
 Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Balance Disorder Confusional State Contusion	Consumer Health Professional	Betaseron (Interferon Beta-1b) Injection			PS
SUBCUTANEOUS	8MIU, EVERY 2D,	Dehydration					
SUBCUTANEOUS		Fall					
		Hallucination Laceration Mental Status Changes Mobility Decreased Multiple Sclerosis Nervous System Disorder Pain Sepsis		Neurontin (Gabapentin) Topamax (Topiramate) Oxycontin (Oxycodone Hydrochloride) Vicodin (Hydrocodone Bitartrate)			SS C C C

Date:12/03/02ISR Number: 4018206-2Report Type:Expedited (15-DaCompany Report #B0286643A
 Age:9 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation Anger Anxiety Depression Drug Ineffective Insomnia Suicidal Ideation	Consumer	Paxil Lithium Risperidone Gabapentin	PS SS SS SS	Glaxo Wellcome Glaxo Wellcome	

Date:12/03/02ISR Number: 4018240-2Report Type:Expedited (15-DaCompany Report #B0285837A
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Confusional State Encephalopathy Hypokalaemia	Health Professional	Abacavir Sulfate + Lamivudine + Zidovudine	PS	Glaxo Wellcome	ORAL
1UNIT Twice per day UNKNOWN UNKNOWN	Hypothermia Respiratory Alkalosis Somnolence		Gabapentin Nicardipine	SS SS		

Date:12/03/02ISR Number: 4018266-9Report Type:Expedited (15-DaCompany Report #B0286643A
Age:9 YR Gender:Male I/FU:I

Outcome	PT
Other	Agitation Anger Anxiety Bipolar Disorder Drug Ineffective Insomnia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Malaise Suicidal Ideation	Report Source	Product	Role	Manufacturer	Route
				Paxil	PS	Glaxo Wellcome	
				Lithium	SS	Glaxo Wellcome	
				Risperidone	SS		
				Gabapentin	SS		

Date:12/03/02ISR Number: 4020718-2Report Type:Expedited (15-DaCompany Report #2002068084
Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Depression	Consumer	Neurontin (Gabapentin)	PS		

Date:12/03/02ISR Number: 4021016-3Report Type:Expedited (15-DaCompany Report #2002068179
Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other			Cardio-Respiratory Arrest Delirium		Neurontin (Gabapentin)	PS		
120 MG BID					Geodon (Ziprasidone)	SS		
					Clozappine	SS		
					Lithium	SS		

Date:12/03/02ISR Number: 4021042-4Report Type:Expedited (15-DaCompany Report #PERCOCET2002-00411
Age:64 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Coma	Literature	Percocet Endo	PS	Endo	ORAL
PO			Completed Suicide	Health	Vicodin Knoll	SS	Knoll	ORAL
PO			Hepatic Failure	Professional	Oxycodone	SS		ORAL

	Respiratory Depression	Risperidone	SS	ORAL
PO		Valproic Acid	SS	ORAL
PO		Gabapentin	SS	ORAL
PO		Meloxicam	SS	ORAL

Date:12/03/02ISR Number: 4021367-2Report Type:Expedited (15-DaCompany Report #2002067815
Age:83 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Clonic Convulsion Confusional State	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL	Incoherent	Professional	Topiramate	SS		ORAL
	Pyrexia		Amiodarone Ramipril;	C C		

Date:12/03/02ISR Number: 4021958-9Report Type:Expedited (15-DaCompany Report #055-0945-M0100042
Age: Gender:Female I/FU:F

Outcome	PT
Disability	Aggression Asthenia Back Pain Cardiovascular Disorder Chest Pain

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Condition Aggravated	Report Source	Product	Role	Manufacturer	Route
1200 MG (TID)		Constipation Depression Difficulty In Walking Dyspnoea	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
ORAL		Hair Growth Abnormal					
		Irritability		Amitriptyline			
		Lymphadenopathy		Hydrochloride	C		
		Mass		Diazepam	C		
		Nasal Congestion		Sertraline	C		
		Neck Pain		Venlafaxine	C		
		Neuropathy Peripheral		Alprazolam	C		
		Oedema		Acetylsalicylic Acid	C		
		Oedema Peripheral					
		Pain In Extremity					
		Paraesthesia					
		Peripheral Coldness					
		Psychotic Disorder					
		Sciatic Nerve Neuropathy					
		Shock					
		Spinal Disorder					
		Tremor					

Date:12/04/02ISR Number: 4022246-7Report Type:Expedited (15-DaCompany Report #2002068595

Age:91 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation	Foreign	Gabapentin			
300 MG		Condition Aggravated	Consumer	(Gabapentin)	PS		ORAL
(DAILY), ORAL		Insomnia					
		Meningioma Benign		Phenobarbital	C		
		Somnolence		Cinnarizine	C		
		Transient Ischaemic		Ginkgo Biloba	C		
		Attack		Dimenhydrinate	C		

Date:12/04/02ISR Number: 4022429-6Report Type:Expedited (15-DaCompany Report #2002064047

Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG		Balance Disorder Cartilage Injury	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other (DAILY), ORAL		Face Oedema	Professional				
		Fall Oedema Peripheral Pain Paraesthesia Somnolence Weight Increased		Sibutramine Hydrochloride Venlafaxine Hydrochloride Oxycocet Diazepam Ranitidine ..	SS SS C C C		

Date:12/04/02ISR Number: 4022479-XReport Type:Expedited (15-DaCompany Report #2002068312
Age: Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Chest Pain Cholelithiasis Gastrooesophageal Reflux

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Disease

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
			Gabapentin	SS		
			Metaxalone	SS		

Date:12/05/02ISR Number: 4021979-6Report Type:Expedited (15-DaCompany Report #2002068411
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 5600 MG (800 Other MG, 7 TIMES A DAY)		Cancer Pain Neurofibromatosis	Consumer	Neurontin (Gabapentin)	PS		
200 MG (100 MG, BID)				Zoloft (Sertraline)	SS		
200 MG (100 MG, BID)				Trazodone Hydrochloride	SS		
1 MG (0.25 MG, QID)				Alprazolam	SS		
15/1500 MG (DOSE STRENGTH)				Vicodin	SS		
5/500 1000 MG (200				Ibuprofen	SS		

MG, FIVE

TIMES A DAY)

162 MG (81

MG, BID)

Acetylsalicylic Acid SS

Date:12/05/02ISR Number: 4021994-2Report Type:Expedited (15-DaCompany Report #2002068601

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Oedema Peripheral	Consumer	Neurontin (Gabapentin)	PS		
Initial or Prolonged						

Date:12/05/02ISR Number: 4022552-6Report Type:Expedited (15-DaCompany Report #2002068668

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Dyspnoea	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		
Initial or Prolonged	Foreign Body Aspiration					
800 MG	Rhonchi	Professional				
(DAILY)						

Date:12/05/02ISR Number: 4022554-XReport Type:Expedited (15-DaCompany Report #2002051749

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	No Adverse Drug Effect	Foreign Health	Neurontin (Gabapentin)	PS		
Initial or Prolonged						
UNKNOWN	600 MG TWICE	Professional				
DAILY UNKNOWN		Company Representative				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/05/02ISR Number: 4022563-0Report Type:Expedited (15-DaCompany Report #2002062761
Age:72 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 800 MG (400 MG, BID), UNKNOWN	Alanine Aminotransferase Increased Aspartate Aminotransferase Increased Blood Alkaline Phosphatase Increased Liver Disorder Skin Necrosis	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:12/05/02ISR Number: 4022565-4Report Type:Expedited (15-DaCompany Report #061-0945-M0200098
Age:66 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 3600 MG (1200 MG, TID), ORAL SUBCUTANEOUS	Alanine Aminotransferase Increased Aspartate Aminotransferase Increased Blood Alkaline Phosphatase Increased Confusional State Disorientation Drug Ineffective Gamma-Glutamyltransferase Increased Pain Urinary Tract Infection	Foreign Health Professional Company Representative	Neurontin (Tablets) (Gabapentin) Ketamine Morphine Amitriptyline Capecitabine	PS SS C C C		ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG (BID), UNKNOWN	Asthenia Chest X-Ray Abnormal Cholecystitis Dyspnoea Echography Abnormal Eosinophilia Face Oedema Hepatic Enzyme Increased Lung Disorder Medication Error Oedema Oedema Peripheral	Foreign Literature Health Professional	Gabapentin (Gabapentin)	PS		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 300 MG QD TITRATED UP TO THREE TID	Confusional State Dysarthria Muscle Rigidity Parkinsonian Gait Tremor		Neurontin	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/06/02ISR Number: 4020335-4Report Type:Direct
Age:77 YR Gender:Female I/FU:I

Company Report #CTU 182290

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300 TID	Anorexia		Neurontin 300 Tid	PS		
Initial or Prolonged	Jaundice Cholestatic		Verapamil	C		
	Nausea		Celebrex	C		
	Neuropathy Peripheral		Lisinopril	C		
	Pruritus		Asa	C		
			Lipitor	C		
			Maxzide	I		

Date:12/09/02ISR Number: 4023154-8Report Type:Expedited (15-DaCompany Report #2002057832
Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 900 MG (TID), Initial or Prolonged ORAL	Agranulocytosis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
	Chest Pain					
	Erythema Infectiosum	Professional				
	Leukopenia		Cefotaxime Sodium	C		
	Myocarditis					
	Pericardial Effusion					
	Splenomegaly					
	Staphylococcal Infection					

Date:12/09/02ISR Number: 4023156-1Report Type:Expedited (15-DaCompany Report #2002055658
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (400 Other MG, TID)	Abscess	Foreign Health	Neurontin (Gabapentin)	PS		
	Hepatocellular Damage					
	Inflammation	Professional				
1200 MG (400	Leukocytosis	Company	Carbamazepine	SS		

MG, TID)

Lung Disorder
Overdose
Pyrexia

Representative

Date:12/09/02ISR Number: 4023273-6Report Type:Expedited (15-DaCompany Report #2002068428
Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dehydration	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		Haematochezia					
900 MG (TID),		Shock	Professional				
ORAL				All Other Therapeutic Products	SS		
				Valaciclovir Hydrochloride	C		
				Ciprofloxacin	C		

Date:12/09/02ISR Number: 4023464-4Report Type:Expedited (15-DaCompany Report #2002066679
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Coordination Abnormal	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
300 MG (THREE		Dyspepsia					
TIMES DAILY),		Paraesthesia	Professional				
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Diclofenac Sodium C

Date:12/09/02ISR Number: 4023512-1Report Type:Expedited (15-DaCompany Report #2002069532
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Drug Ineffective Euphoric Mood	Foreign Literature	Gabapentin (Gabapentin)	PS		ORAL
(DAILY), ORAL			Health	Reboxetine	SS		ORAL
ORAL			Professional	Venlafaxine	SS		ORAL
				All Other Therapeutic Products	C		
				Tranylcypromine	C		
				Fluoxetine	C		
				Lithium	C		
				Valproate Semisodium	C		
				Lamotrigine	C		

Date:12/09/02ISR Number: 4023514-5Report Type:Expedited (15-DaCompany Report #2002068856
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100 MG, ORAL		Anaphylactic Reaction	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
			Professional Company Representative	Ibuprofen Dextropropoxyphene Hydrochloride Amitriptyline Codeine Tramadol Panadeine Co	C C C C C C		

Date:12/09/02ISR Number: 4023526-1Report Type:Expedited (15-DaCompany Report #2002069039
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Anxiety Clonic Convulsion	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
1800 MG		Condition Aggravated	Professional				
(DAILY), ORAL		Loss Of Consciousness		Clozapine	SS		ORAL
750 MG		Urinary Incontinence					
(DAILY), ORAL				Venlafaxine	C		

Date:12/11/02ISR Number: 4021683-4Report Type:Direct Company Report #CTU 182559
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG BID		Condition Aggravated		Neurontin 300mg	PS		ORAL
Initial or Prolonged ORAL		Renal Impairment					
Required 600MG TID				Neurontin 600mg	SS		ORAL
Intervention to ORAL							
Prevent Permanent Impairment/Damage							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/11/02ISR Number: 4022274-1Report Type:Direct
Age:42 YR Gender:Male I/FU:I

Company Report #CTU 182517

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Alanine Aminotransferase		Gabapentin	PS		
200 MG QID							
Intervention to		Abnormal		Niferex	C		
Prevent Permanent		Aspartate		Vancomycin	C		
Impairment/Damage		Aminotransferase Abnormal		Oxycodone Sr	C		
		Eosinophilia		Metazapine	C		
		Thrombocytopenia		Mgox	C		
				Ducosate	C		

Date:12/11/02ISR Number: 4024243-4Report Type:Expedited (15-DaCompany Report #200206908
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Hyperhidrosis	Foreign	Neurontin			
Initial or Prolonged		Lactose Intolerance	Consumer	(Gabapentin)	PS		ORAL
1800 MG							
(TID), ORAL		Vomiting					
				Methadone	C		

Date:12/12/02ISR Number: 4022813-0Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 182590

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Arthralgia		Neurontin 300mg 1			
Intervention to		C-Reactive Protein		Qam , 3 Q Hs	PS		ORAL
1200 MG/D PO							
Prevent Permanent		Increased		Paxil	C		
Impairment/Damage		Chest Discomfort		Percocet	C		
		Face Oedema					
		Oedema					
		Red Blood Cell					
		Sedimentation Rate					
		Increased					
		Weight Increased					

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia	Consumer	Neurontin			
1600 MG (400		Arthritis	Health	(Gabapentin)	PS		ORAL
MG, FOUR		Blood Cholesterol	Professional				
TIMES DAILY),		Increased					
ORAL		Blood Triglycerides					
10 MG		Increased		Lipitor			
(DAILY), ORAL		Carpal Tunnel Syndrome		(Atorvastatin)	SS		ORAL
		Constipation					
		Fibromyalgia		Lorazepam	C		
		Gingival Abscess		Bactrim	C		
		Gingival Disorder		Paracetamol	C		
		Hypothyroidism		Amitriptyline			
		Iatrogenic Injury		Hydrochloride	C		
		Insomnia		Zipradisone			
		Stress		Hydrochloride	C		
				Quetiapine Fumarate	C		
				Risperidone	C		
				Carbamazepine	C		
				Estrogens Conjugated	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Levothyroxine Sodium C

Date:12/12/02ISR Number: 4025720-2Report Type:Expedited (15-DaCompany Report #2002062558
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2700 MG		Coma Hypoaesthesia	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
(TID), ORAL		Medication Error	Health				
ORAL		Paraesthesia	Professional	Morphine	SS		ORAL
TRANSDERMAL	100 MCG,			Fentanyl	SS		
TRANSDERMAL				Trandolapril	C		
				Diazepam	C		
				Etodolac	C		
				Beclometasone			
				Dipropionate	C		
				Salbutamol	C		
				Tramadol			
				Hydrochloride	C		
				Pethidine	C		

Date:12/13/02ISR Number: 4026355-8Report Type:Expedited (15-DaCompany Report #1005340
 Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Glucose Fluctuation	Foreign	Keppra	PS		
2500 MG		Closed Head Injury	Health	Keppra	SS		
1500 MG		Coagulopathy	Professional	Keppra	SS		
1000 MG		Convulsion		Keppra	SS		
2000 MG		Haemorrhage		Keppra	SS		

900 MG	Hepatocellular Damage	Gabapentin	SS
	Hypoglycaemia	Gabapentin	SS
1500 MG			
	Hypotension	Gabapentin	SS
900 MG			
	Hypothermia	Valproic Acid	SS
300 MG			
	Prothrombin Time Shortened	Carbamazepine	SS
600 MG		Carbamazepine	SS
	Status Epilepticus	Oxcarbazepine	SS
900 MG			
	Stupor	Pantozol	C
	Thrombocytopenia	Lasix	C
		Bactrim	C
		Elobact	C

Date:12/16/02ISR Number: 4026803-3Report Type:Expedited (15-DaCompany Report #2002070919
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Electrocardiogram Qt Prolonged	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Amiodarone	C		

Date:12/16/02ISR Number: 4026859-8Report Type:Expedited (15-DaCompany Report #2002070374
Age: Gender:Male I/FU:I

Outcome
Hospitalization -
Initial or Prolonged

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Freedom Of Information (FOI) Report

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Neutropenia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:12/16/02ISR Number: 4026860-4Report Type:Expedited (15-DaCompany Report #2002070294
Age:57 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 800 MG (BID), Initial or Prolonged ORAL			Confusional State Muscular Weakness Pneumonia	Foreign Health Professional Company Representative	Neurontin (Tablets) (Gabapentin) Phenytoin Sodium Valproate Sodium	PS C C		ORAL

Date:12/16/02ISR Number: 4026862-8Report Type:Expedited (15-DaCompany Report #2002052677
Age:48 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG,TID,), ORAL			Abnormal Behaviour Back Pain Blood Glucose Increased Disinhibition Disturbance In Attention Dizziness Drug Ineffective Haematocrit Decreased High Density Lipoprotein Decreased Medication Error Memory Impairment	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Oxycodone Hydrochloride Paracetamol Atorvastatin Prochlorperazine Oxycodone Hydrochloride Tramadol	PS C C C C C		ORAL

Nausea
Neck Pain
Personality Change
Therapeutic Response
Unexpected
Vertigo
Vomiting

Hydrochloride

C

Date:12/17/02ISR Number: 4026123-7Report Type:Direct
Age:45 YR Gender:Male I/FU:I

Company Report #CTU 182829

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Discomfort Cholelithiasis		Neutronin 600mg Tablets Pfizer	PS	Pfizer	ORAL
ONE DAY ORAL		Dysgeusia Gastrooesophageal Reflux Disease		Darvocet	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/17/02ISR Number: 4027966-6Report Type:Expedited (15-DaCompany Report #2002070421
 Age:87 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		General Physical Health	Consumer	Neurontin			
Life-Threatening		Deterioration		(Gabapentin)	PS		
Other		Medication Error		Lamotrigine	SS		
		Respiratory Failure		Risperidone	SS		
				All Other Therapeutic Products	SS		

Date:12/17/02ISR Number: 4028479-8Report Type:Expedited (15-DaCompany Report #055-0945-M0100042
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Abdominal Distension	Foreign	Gabapentin			
800 MG (400 MG, BID), ORAL		Abnormal Behaviour	Consumer	(Gabapentin)	PS		ORAL
		Aggression					
		Back Pain					
		Blood Cholesterol Increased		Amitriptyline Hydrochloride	C		
		Blood Triglycerides Increased		Diazepam	C		
		Cardiovascular Disorder Condition Aggravated		Sertraline	C		
		Constipation		Venlafaxine	C		
		Depression		Alprazolam	C		
		Difficulty In Walking		Acetylsalicylic Acid	C		
		Dizziness					
		Dyspnoea					
		Electric Shock					
		Ganglion					
		Nasal Congestion					
		Neck Pain					
		Neuropathy Peripheral					
		Oedema					
		Oedema Peripheral					
		Pain					
		Pain In Extremity					
		Peripheral Coldness					

Psychotic Disorder
Sciatica
Tremor
Weight Increased

Date:12/18/02ISR Number: 4025151-5Report Type:Expedited (15-DaCompany Report #WAES 0205USA01611
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Blood Pressure Decreased	Health	Prinivil	PS	Merck & Co., Inc	ORAL
Initial or Prolonged		Dizziness	Professional	Neurontin	SS		
		Drug Interaction		Prinivil	SS		ORAL

Date:12/18/02ISR Number: 4029491-5Report Type:Expedited (15-DaCompany Report #2002068085
Age:63 YR Gender:Female I/FU:I

Outcome	PT
Other	Asthenia Crying

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Depression Suicidal Ideation	Report Source	Product	Role	Manufacturer	Route
300 MG (QD), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Docetaxel	C		
				Zoledronic Acid	C		
				Terazosin Hydrochloride	C		
				Hyzaar	C		
				Potassium Chloride	C		
				Ranitidine Hydrochloride	C		
				Naproxen	C		
				Acetylsalicylic Acid	C		
				Vitamins	C		
				Amino Acids	C		

Date:12/18/02ISR Number: 4029493-9Report Type:Expedited (15-DaCompany Report #2002068084

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Depression	Consumer	Neurontin (Gabapentin)	PS		

Date:12/18/02ISR Number: 4030402-7Report Type:Expedited (15-DaCompany Report #2002071293

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Fatigue	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
			Professional Company Representative	Mirtazapine	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (100 MG, TID), ORAL		Anorexia Aptyalism Asthenia Confusional State	Foreign Literature Health Professional	Phenytoin Suspension (Phenytoin Sodium)	PS		ORAL
ORAL		Crying Depressed Mood		Gabapentin(Gabapenti n)	SS		ORAL
ORAL		Dizziness Drug Hypersensitivity Fatigue Insomnia		Amitriptyline Hydrochloride(Amitri ptyline Hydrochloride)	SS		ORAL
45 MG (BID) ORAL		Memory Impairment					
TRANSDERMAL	25 UG/H,	Nervous System Disorder		Fentanyl	SS		
TRANSDERMAL		Pain					
PRN, ORAL, SUBCUTANEOUS		Paraesthesia Rash Somnolence		Morphine Oxybutynin Docusate Sodium Lorazepam	SS C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sennoside A C
Glycerol C

Date:12/20/02ISR Number: 4030868-2Report Type:Expedited (15-DaCompany Report #2002070769
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence	Health Professional	Neurontin (Gabapentin)	PS		

Date:12/20/02ISR Number: 4031017-7Report Type:Expedited (15-DaCompany Report #001-0945-M0200668
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Drug Abuser	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Drug Screen False					
Other		Positive		Dilantin Suspension (Phenytoin Sodium)	SS		ORAL
ORAL							

Date:12/20/02ISR Number: 4031277-2Report Type:Expedited (15-DaCompany Report #2002050043
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Distension	Consumer	Neurontin (Gabapentin)	PS		ORAL
(TWICE		Adrenal Adenoma	Health Professional				
DAILY), ORAL		Asthenia					
ORAL		Blood Arsenic Increased		Lamotrigine	SS		ORAL
		Blood Heavy Metal Increased		Clonazepam	SS		
		Dizziness					
		Dyspnoea					
		Headache					
		Helicobacter Pylori					

Identification Test
Positive
Hypertension
Muscle Spasms
Nausea
Nervousness
Oesophageal Spasm
Oral Intake Reduced
Tardive Dyskinesia
Weight Increased

Date:12/20/02ISR Number: 4031287-5Report Type:Expedited (15-DaCompany Report #2002070768
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), Other ORAL		Medication Error Paraesthesia	Consumer	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/20/02ISR Number: 4031916-6Report Type:Expedited (15-DaCompany Report #2002067435
Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1800 MG, ORAL	Supraventricular Tachycardia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:12/20/02ISR Number: 4031918-XReport Type:Expedited (15-DaCompany Report #2002066679
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG, THREE TIMES DAILY, ORAL	Autonomic Nervous System Imbalance	Foreign Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
		Coordination Abnormal		Diclofenac Sodium	C		
		Dyspepsia					
		Paraesthesia					

Date:12/23/02ISR Number: 4032438-9Report Type:Expedited (15-DaCompany Report #2002054167
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	10 MG (DAILY), ORAL	Amnesia	Health Professional Company Representative	Lipitor (Atorvastatin)	PS		ORAL
Other	900 MG (TID), ORAL	Arthralgia		Neurontin (Gabapentin)	SS		ORAL
		Cognitive Disorder					
		Coordination Abnormal					
		Depression					
		Diabetes Mellitus					
		Inadequate Control		Insulin	C		
		Diabetic Neuropathy		Metformin			

Difficulty In Walking	Hydrochloride	C
Disturbance In Attention	Provella-14	C
Hypersensitivity	Fluoxetine	
Hypertension	Hydrochloride	C
Influenza Like Illness	Valsartan	C
Oedema Peripheral	Loratadine	C
Pain In Extremity	Irbesartan	C
Spinal Osteoarthritis		
Tarsal Tunnel Syndrome		

Date:12/23/02ISR Number: 4032821-1Report Type:Expedited (15-DaCompany Report #2002071322
Age:49 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 40 MG (BID), Initial or Prolonged ORAL	Disease Recurrence	Literature	Geodon (Ziprasidone)	PS		ORAL
Other 600 MG (BID), ORAL	Hyponatraemia	Health				
	Neuroleptic Malignant Syndrome	Professional	Gabapentin (Gabapentin)	SS		ORAL
	Neutrophilia					
	Polydipsia Psychogenic Psychotic Disorder		Venlafaxine Hydrochloride	SS		ORAL
300 MG (BID), ORAL	Rhabdomyolysis					
1.5 MG (DAILY), ORAL	Tardive Dyskinesia		Lorazepam	SS		ORAL
	White Blood Cell Count Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/23/02ISR Number: 4032850-8Report Type:Expedited (15-DaCompany Report #2002071087

Age:3 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2100 MG Other (Q8H), ORAL	Contusion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Atovaquone	SS		
			Prednisone	C		
			Metronidazole	C		
			Amlodipine	C		
			Potassium Chloride	C		
			Diphenhydramine Hydrochloride	C		
			Nystatin	C		
			Ergocalciferol	C		
			Cyproheptadine	C		
			Tacrolimus	C		
			Nystatin	C		
			Magnesium Gluconate	C		
			Lactobacillus			
			Acidophilus	C		
			Vitamins	C		
			Colistin Mesilate			
			Sodium	C		
			Ranitidine	C		
			Enalapril	C		
			Omeprazole	C		
			Atovaquone	C		

Date:12/23/02ISR Number: 4032864-8Report Type:Expedited (15-DaCompany Report #2002067264

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Drug Screen Positive Intentional Misuse Post Procedural Pain Urine Amphetamine Positive	Health Professional	Neurontin (Gabapentin)	PS		
			Nefazodone			
			Hydrochloride	SS		
			All Other Therapeutic Products	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agranulocytosis	Consumer	Neurontin			
(DAILY), ORAL		Bradycardia		(Gabapentin)	PS		ORAL
		Hypotension		Tiagabine			
8 MG (QID),		Thrombocytopenia		Hydrochloride	SS		ORAL
ORAL							
(DAILY), ORAL				Lamotrigine	SS		ORAL
				Lithium	C		
				Zaleplon	C		
				Loratadine	C		
				Multivitamins	C		
				Docusate Sodium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/23/02ISR Number: 4033109-5Report Type:Expedited (15-DaCompany Report #2002071441
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Colour Blindness	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Visual Field Defect	Professional Company Representative	Hormonin	C		

Date:12/23/02ISR Number: 4033137-XReport Type:Expedited (15-DaCompany Report #USA-2002-0003699
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Aspiration	Consumer	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		
Other		Brain Death	Other	Diazepam (Diazepam)	SS		
		Brain Herniation		Methocarbamol (Methocarbamol)	SS		
		Coma		Neurontin (Gabapentin)	SS		
		Drug Dependence					
		Gastrointestinal					
		Haemorrhage					
		Hydrocephalus					
		Loss Of Consciousness					

Date:12/23/02ISR Number: 4033150-2Report Type:Expedited (15-DaCompany Report #2002051749
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Constipation	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:12/24/02ISR Number: 4033968-6Report Type:Expedited (15-DaCompany Report #2002GB01238
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 40 MG DAILY		Blood Pressure Decreased	Foreign	Lisinopril	PS		ORAL
Initial or Prolonged PO		Dizziness	Health				
900 MG DAILY		Drug Interaction	Professional	Neurontin	SS		
			Other				

Date:12/24/02ISR Number: 4034289-8Report Type:Expedited (15-DaCompany Report #A0388885A
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation Memory Impairment	Health Professional	Lamictal Unspecified Tablet (Lamotrigine)	PS		ORAL
200 MG / PER DAY / ORAL			Company				
800 MG / PER DAY			Representative	Gabapentin (Gabapentin)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/26/02ISR Number: 4035189-XReport Type:Expedited (15-DaCompany Report #2002071937

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anaphylactic Reaction	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 (TID),			Professional				
ORAL				Diclofenac Sodium	SS		ORAL
100 MG							
(DAILY), ORAL							

Date:12/26/02ISR Number: 4035190-6Report Type:Expedited (15-DaCompany Report #2002068595

Age:91 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation	Foreign	Gabapentin (Gabapentin)	PS		ORAL
300 MG		Insomnia	Consumer				
(DAILY), ORAL		Meningioma Benign					
		Transient Ischaemic Attack		Phenobarbital	C		
		Urinary Tract Infection		Cinnarizine	C		
				Ginkgo Biloba	C		
				Dimenhydrinate	C		

Date:12/26/02ISR Number: 4035328-0Report Type:Expedited (15-DaCompany Report #2002071743

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG		Difficulty In Walking	Health				
Other		Hypoaesthesia	Professional				
(DAILY), ORAL		Intervertebral Disc Protrusion					
		Muscle Twitching					

Nausea
Paraesthesia

Date:12/26/02ISR Number: 4035329-2Report Type:Expedited (15-DaCompany Report #2002071949
Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cyst	Consumer	Neurontin			
900 MG (300		Drug Ineffective		(Gabapentin)	PS		ORAL
MG, THREE		Eating Disorder Symptom					
		Nausea					
TIMES DAILY),		Spinal Disorder					
ORAL		Throat Tightness					
		Vertigo					
		Weight Decreased					

Date:12/26/02ISR Number: 4035573-4Report Type:Expedited (15-DaCompany Report #2002071679
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disease Recurrence	Foreign	Neurontin			
UNKNOWN		Pemphigoid	Health	(Gabapentin)	PS		ORAL
(UNKNOWN),			Professional				
ORAL				Carbamazepine	C		

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Freedom Of Information (FOI) Report

Date:12/26/02ISR Number: 4035577-1Report Type:Expedited (15-DaCompany Report #2002069054
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urinary Retention	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
2700 MG		Urinary Tract Disorder	Professional				
(TID), ORAL			Company Representative	Phenytoin	C		
				Paroxetine Hydrochloride	C		
				Omeprazole Magnesium	C		
				Insulin Human	C		
				Lorazepam	C		
				Zolpidem	C		
				Paracetamol	C		

Date:12/26/02ISR Number: 4035709-5Report Type:Expedited (15-DaCompany Report #2002071934
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haemolytic Anaemia	Health Professional	Neurontin (Gabapentin)	PS		
				Omeprazole	C		
				Valproate Semisodium	C		
				Oxycocet	C		

Date:12/27/02ISR Number: 4034766-XReport Type:Expedited (15-DaCompany Report #2002070294
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Confusional State	Foreign Health	Neurontin (Tablets)			
Hospitalization -		Dysphagia	Health	(Gabapentin)	PS		ORAL
1600 MG							
Initial or Prolonged		Motor Dysfunction	Professional				
(BID), ORAL							
Other		Muscular Weakness	Company Representative	Valproate Sodium	C		
		Pharmaceutical Product Complaint					

Date:12/27/02ISR Number: 4034894-9Report Type:Expedited (15-DaCompany Report #2002050928

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chronic Fatigue Syndrome	Consumer	Neurontin			
Other		Decreased Activity	Health	(Gabapentin)	PS		ORAL
(QID), ORAL		Dental Caries	Professional	Venlafaxine			
		Face Oedema		Hydrochloride	C		
		Hypothyroidism		Paroxetine			
		Mental Disorder		Hydrochloride	C		
		Migraine		Tramadol			
		Oral Intake Reduced		Hydrochloride	C		
		Oral Pain		Clonazepam	C		
		Oral Soft Tissue Disorder		Carisoprodol	C		
		Pyrexia					
		Tooth Abscess					
		Tooth Discolouration					
		Tooth Disorder					
		Tooth Loss					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/30/02ISR Number: 4035822-2Report Type:Expedited (15-DaCompany Report #2002072250
Age:85 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (200 MG, TID), ORAL		Choking Condition Aggravated Dehydration Dysphagia Lipoma Oligodipsia	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Metoprolol Succinate	C		

Date:12/30/02ISR Number: 4035904-5Report Type:Expedited (15-DaCompany Report #2002072136
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dyspepsia Grand Mal Convulsion	Health Professional	Neurontin (Gabapentin) Dilantin Suspension (Phenytoin Sodium)	PS SS		

Date:12/30/02ISR Number: 4036623-1Report Type:Expedited (15-DaCompany Report #2002072435
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		White Blood Cell Count Decreased	Foreign Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL

Date:12/31/02ISR Number: 4036968-5Report Type:Expedited (15-DaCompany Report #2002061973
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Disturbance In Attention	Consumer	Neurontin			

300 MG	Dizziness	Health	(Gabapentin)	PS	ORAL
(ONCE), ORAL	Feeling Abnormal	Professional			
	Speech Disorder		Paroxetine		
	Syncope		Hydrochloride	C	
	Tongue Oedema		Levothyroxine Sodium	C	
			Metoprolol Succinate	C	
			Propacet	C	

Date:12/31/02ISR Number: 4038530-7Report Type:Expedited (15-DaCompany Report #2002072480
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neurotoxicity	Foreign	Neurontin			
		Renal Disorder	Health	(Gabapentin)	PS		ORAL
900 MG (TID),			Professional				
ORAL			Company				
			Representative				

Date:12/31/02ISR Number: 4040195-5Report Type:Expedited (15-DaCompany Report #UK025779
 Age:75 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Angioneurotic Oedema
Initial or Prolonged	Erythema

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pruritus

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
SUBCUTANEOUS	100 MG,	Foreign	Kineret	PS		
DAILY, SC		Health				
100 MG,		Professional	Gabapentin	SS		ORAL
DAILY, PO			Fentanyl	C		
			Celecoxib	C		
			Cloprednol	C		
			Methotrexate	C		
			Tilidine			
			Hydrochloride	C		
			Raloxifen Hcl	C		
			Hydrochlorothiazide/			
			Triamterene	C		
			Ranitidine			
			Hydrochloride	C		
			Gabapentin	C		

Date:01/01/03ISR Number: 4033667-0Report Type:Direct
Age:40 YR Gender:Female I/FU:I

Company Report #CTU 183672

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Failure Of Implant		Atenol 25 Mg Mylan	PS	Mylan	ORAL
1 DAILY ORAL				Neurontin 300 Mg Pd	SS	Pd	ORAL
Hospitalization -							
3 DAILY ORAL							
Initial or Prolonged							
Disability							
Required							
Intervention to							
Prevent Permanent							
Impairment/Damage							

Date:01/01/03ISR Number: 4033976-5Report Type:Direct
Age:73 YR Gender:Male I/FU:I

Company Report #CTU 183683

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Pain In Extremity		Neurontin 100mg Pfizer Us	PS	Pfizer Us	ORAL
THREE CAP							
THREE TIMES							
ORAL							

Date:01/03/03ISR Number: 4038684-2Report Type:Expedited (15-DaCompany Report #2002072453
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Condition Aggravated Hepatic Encephalopathy	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:01/03/03ISR Number: 4039076-2Report Type:Expedited (15-DaCompany Report #2002072779
Age:82 YR Gender:Female 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
1500 MG (TID)		Coma Drug Interaction	Health Professional	Neurontin (Gabapentin)	PS		
390 MG (PRN)		Drug Toxicity Hypotonia		Dextropropoxyphene Hydrochloride	SS		
		Lethargy Muscular Weakness Pneumonia		Combivent Prednisone Gliclazide Zolpidem Tartrate Paracetamol Pantoprazole Combivent Ergocalciferol Calcium Levothyroxine Sodium Calcitonin, Salmon	C C C C C C C C C C C		

Date:01/03/03ISR Number: 4039077-4Report Type:Expedited (15-DaCompany Report #2002073234
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Anger Drug Dependence Treatment Noncompliance	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:01/03/03ISR Number: 4039078-6Report Type:Expedited (15-DaCompany Report #2002062997
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 Disability MG, TID),		Anxiety Aphasia Cerebral Haemorrhage	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Other
ORAL

Cerebrovascular Accident

Complex Partial Seizures
Confusional State
Depression
Insomnia
Memory Impairment

Cytalopram
Hydrobromide C
Lotrel C
Peri-Colace C

Date:01/03/03ISR Number: 4039141-XReport Type:Expedited (15-DaCompany Report #2002072974
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 800 MG (DAILY), ORAL		Affective Disorder Anorexia Anxiety Drug Dependence Fatigue	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Bupropion Hydrochloride Thioridazine Hydrochloride Doxepin Hydrochloride	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/06/03ISR Number: 4039504-2Report Type:Expedited (15-DaCompany Report #ZONI000979

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anger	Literature	Zonisamide			
ORAL		Completed Suicide	Health	(Zonisamide)	PS		ORAL
ORAL		Gun Shot Wound	Professional	Gabapentin			
				(Gabapentin)	SS		ORAL
ORAL				Phenytoin			
				(Phenytoin)	SS		ORAL

Date:01/07/03ISR Number: 4039229-3Report Type:Expedited (15-DaCompany Report #2002066569

Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Diplopia	Foreign	Neurontin			
Other		Vitreous Detachment	Health	(Gabapentin)	PS		ORAL
900 MG (300			Professional				
MG, THREE				Tramadol			
TIMES DAILY),				Hydrochloride	C		
ORAL				Paramol-118	C		
				Amitriptyline	C		

Date:01/07/03ISR Number: 4039256-6Report Type:Expedited (15-DaCompany Report #2002069842

Age:47 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Drug Level Decreased	Foreign	Neurontin			
Initial or Prolonged			Health	(Gabapentin)	PS		
1800 MG							

(DAILY),

Ciclosporin C
 All Other
 Therapeutic Products C

Date:01/07/03ISR Number: 4039546-7Report Type:Expedited (15-DaCompany Report #2002050012

Age:83 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Coma	Literature	Gabapentin			
Other		Drug Level Above	Health	(Gabapentin)	PS		
200 MG (AFTER		Therapeutic	Professional				
EACH		Haemodialysis					
DIALYSIS)		Mental Status Changes		Amiodarone	C		
				Calcium Carbonate	C		
				Docusate Sodium	C		
				Erythropoietin	C		
				Folic Acid	C		
				Metoprolol	C		
				Tamsulosin	C		
				Lactulose	C		
				Promethazine	C		
				Amitriptyline	C		
				Carbamazepine	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/07/03ISR Number: 4039548-0Report Type:Expedited (15-DaCompany Report #2002059016

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetes Mellitus	Health	Neurontin			
		Grand Mal Convulsion	Professional	(Gabapentin)	PS		ORAL
		Hyperventilation					
				Amitriptyline	SS		
				Levothyroxine Sodium	C		
				Glipizide	C		
				Metformin			
				Hydrochloride	C		

Date:01/08/03ISR Number: 4038509-5Report Type:Direct

Company Report #CTU 184077

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain		Neurontin 300 Mg			
				Parke Davis	PS	Parke Davis	ORAL
				Carbitrol	C		
				Lescol Xl	C		
				Lexapro	C		
				Prevacid	C		
				Multivitamin	C		
				Aspirin	C		

Date:01/08/03ISR Number: 4040441-8Report Type:Expedited (15-DaCompany Report #2003000080

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Myeloid Metaplasia	Consumer	Neurontin			
		Paraesthesia		(Gabapentin)	PS		

Date:01/08/03ISR Number: 4040481-9Report Type:Expedited (15-DaCompany Report #2002066580
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Delirium	Consumer	Neurontin			
Initial or Prolonged	Hallucination	Health	(Gabapentin)	PS		
1800 MG (TID)						
Other		Professional	Famotidine	C		

Date:01/08/03ISR Number: 4040493-5Report Type:Expedited (15-DaCompany Report #2002073286
Age:35 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Anxiety
Initial or Prolonged	Asthenopia
Other	Clumsiness
	Depression
	Diplopia
	Eye Pain
	Fatigue
	Mental Disorder
	Muscle Twitching
	Oedema
	Pain In Extremity
	Suicidal Ideation

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tendonitis Tremor Vision Blurred	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG (300 MG, QID), ORAL		Weight Decreased		Promethazine Hydrochloride Hydrochlorothiazide Furosemide Librax Hyoscyamine Sulfate Rabeprazole Sodium Fexofenadine Hydrochloride All Other Therapeutic Products Fortagesic Fluoxetine Hydrochloride Bupropion Hydrochloride Trimethobenzamide Hydrochloride	SS C C C C C C C C C C C C C C C C C C		

Date:01/08/03ISR Number: 4040499-6Report Type:Expedited (15-DaCompany Report #001-0945-950171
Age:66 YR Gender:Female I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression Drug Interaction	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300MG Other 3 TIMES DAILY), ORAL (25MG ONE		Hypertensive Crisis Myocardial Ischaemia	Professional	Zoloft (Sertraline)	SS		ORAL

TIME DOSE),

ORAL

Decadron 4mg (Dexamethasone)	C
Decadron 1.5mg (Dexamethasone)	C
Decadron 1.5mg (Dexamethasone)	C
Midrin(Midrid)	C
Ativan 0.25mg (Lorazepam)	C
Ativan 0.5mg (Lorazepam)	C
Berocca Plus (Berocca /Old Form/)	C
Procardia (Nifedipine)	C
Zantac 150mg (Ranitidine Hydrochloride)	C
Compazine 10mg (Prochlorperazine Edisylate)	C
Demerol 75mg Im (Pethidine)	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C
 Vistaril 50mg
 (Hydroxyzine
 Embonate) C
 Serax 10mg(Oxazepam) C
 Milk Of Magnesia
 (Magnesium
 Hydroxide) C
 Ativan 0.5mg
 (Lorazepam) C

Date:01/09/03ISR Number: 4040792-7Report Type:Expedited (15-DaCompany Report #2002073243
 Age:67 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged (THREE TIMES Other DAILY), ORAL	Alanine Aminotransferase Increased	Foreign Health	Neurontin(Gabapentin)	PS		ORAL
ORAL	Blood Bilirubin Increased	Professional				
ORAL	Microcytic Anaemia		Tahor(Atorvastatin)	SS		ORAL
ORAL	Pyelonephritis		Asasantin	SS		ORAL
	Rectal Haemorrhage Thrombocytopenia					

Date:01/10/03ISR Number: 4041121-5Report Type:Expedited (15-DaCompany Report #2002066162
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other ORAL	Rhinorrhoea Surgery	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Terazosin Hydrochloride	C		
			Diltiazem	C		
			Finasteride	C		
			Oxycocet	C		

Date:01/10/03ISR Number: 4041148-3Report Type:Expedited (15-DaCompany Report #2002071293

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Fatigue	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Professional Company Representative	Mirtazapine	C		

Date:01/10/03ISR Number: 4041285-3Report Type:Expedited (15-DaCompany Report #2003000197

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG(300MG, TID), ORAL	Blood Bilirubin Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/10/03ISR Number: 4041300-7Report Type:Expedited (15-DaCompany Report #EMADSS2003000044
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Coma Hypoaesthesia Paraesthesia	Foreign Health Professional	Durogesic (100 Mcg/Hr Patch) (Fentanyl)			
TRANSDERMAL	100 MCG,				PS		
TRANSD				Gabapentin (Gabapentin)	SS		ORAL
2700 MG,							
DAILY, ORAL				Trandolapril (Trandolapril)	C		
				Diazepam (Diazepam)	C		
				Etodolac (Etodolac)	C		
				Beclomethasone (Beclometasone)	C		
				Salbutamol (Salbutamol)	C		
				Zydol (Tramadol Hydrochloride)	C		
				Pethidine (Pethidine)	C		

Date:01/10/03ISR Number: 4041333-0Report Type:Expedited (15-DaCompany Report #DE9388106JAN2003
Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Bowel Sounds Abnormal Intentional Misuse	Health Professional	Tavor (Lorazepam, Tablet, 0)	PS		ORAL
12 TABLETS (OVERDOSE AMOUNT 12 MG)	1 DAY	Miosis Stupor Suicide Attempt		Neurontin (Gabapentin, , 0)	SS		ORAL
5 TABLETS							

(OVERDOSE
AMOUNT 1500
MG) 1 DAY

Tafil (Alprazolam, ,
0) SS ORAL

10 TABLETS

(OVERDOSE
AMOUNT 10 MG) 1 DAY

Ximovan (Zopiclone,
,0) SS ORAL

20 TABLETS

(OVERDOSE
AMOUNT 150
MG) 1 DAY

Zolpidem (Zolpidem,
,0) SS ORAL

20 TABLETS

(OVERDOSE
AMOUNT 200
MG) 1 DAY

Zopiclone
(Zopiclone, , 0) SS ORAL

10 TABLETS

(OVERDOSE
AMOUNT 75 MG) 1 DAY

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/10/03ISR Number: 4041469-4Report Type:Expedited (15-DaCompany Report #2002071949
Age:76 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG, THREE TIMES DAILY) ORAL		Cyst Dizziness Drug Ineffective Dyspnoea Nausea Oral Intake Reduced Spinal Disorder Throat Tightness Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:01/10/03ISR Number: 4041471-2Report Type:Expedited (15-DaCompany Report #2002073396
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG Other (DAILY), ORAL		Anger Anxiety Bipolar Disorder Blood Glucose Abnormal Delusion Depression Diarrhoea Disease Recurrence Drug Effect Decreased Drug Interaction Dyspepsia Emotional Disorder Eye Pain Flatulence Flushing Gastrointestinal Disorder Hyperhidrosis Mental Disorder Migraine	Health Professional	Neurontin (Gabapentin) Venlafaxine Hydrochloride Topiramate Oxycocet Ibuprofen	PS SS SS C C		ORAL

Mydriasis
Nausea
Nervousness
Oral Intake Reduced
Suicidal Ideation
Suicide Attempt
Vision Blurred
Vomiting

Date:01/13/03ISR Number: 4040657-0Report Type:Direct Company Report #USP 55444
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Neurontin	PS	Parke Davis	
				Neurontin	SS	Parke Davis	

Date:01/13/03ISR Number: 4041685-1Report Type:Expedited (15-DaCompany Report #2002066041
Age:50 YR Gender:Female I/FU:I

Outcome	PT
Other	Anxiety
	Arthropathy

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Coordination Abnormal Depressed Level Of Consciousness	Report Source	Product	Role	Manufacturer	Route
2400 MG (TID), ORAL		Dry Mouth Dysarthria	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Emotional Disorder	Professional				
		Face Oedema Fatigue Feeling Abnormal Hypertension Insomnia Memory Impairment Nausea Nervousness Oedema Oedema Peripheral Palpitations Petit Mal Epilepsy Photopsia Scintillating Scotoma Somnolence Tremor Vision Blurred		Tramadol Hydrochloride	C		

Date:01/13/03ISR Number: 4041786-8Report Type:Expedited (15-DaCompany Report #2002129159FR
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour Confusional State	Foreign Health	Celebrx (Celecoxib) Capsule	PS		ORAL
200 MG, QD, ORAL		Diplopia	Professional				
1200 TID, ORAL		Gastric Perforation Haematoma	Other	Neurontin (Gabapentin)	SS		ORAL
		Overdose					
		Perforated Ulcer Visual Disturbance Vomiting		Mopral Lexomil Rythmodan	C C		

(Disopyramide) C
 Seresta C
 Lasilix C
 Estreva C
 Utrogestan C

Date:01/13/03ISR Number: 4041816-3Report Type:Expedited (15-DaCompany Report #2003000058
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Cholesterol Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
2700 MG		Blood Pressure Increased					
(THREE TIMES		Diabetes Mellitus					
DAILY), ORAL		Liver Disorder		Norvasc (Amlodipine)	SS		ORAL
5 MG (DAILY),		Muscle Disorder					
ORAL		Parkinson'S Disease Tremor		Lipitor (Atorvastatin)	SS		ORAL
20 MG							
(DAILY), ORAL				Pioglitazone Hydrochloride	C		
				Insulin Human			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Injection, Isophane C
 Insulin Human C
 Valsartan C

Date:01/13/03ISR Number: 4041903-XReport Type:Expedited (15-DaCompany Report #044-0945-M0100090
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cardiac Failure	Foreign	Gabapentin			
		Lower Respiratory Tract	Study	(Gabapentin)	PS		
		Infection	Health				
		Transient Ischaemic	Professional				
		Attack					

Date:01/16/03ISR Number: 4043707-0Report Type:Expedited (15-DaCompany Report #2002062399
 Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour	Foreign	Neurontin			
Other		Aggression	Health	(Gabapentin)	PS		ORAL
(TID), ORAL							
		Confusional State	Professional	Celecoxib	SS		ORAL
(DAILY), ORAL							
		Diplopia	Company	Antiulcerous			
		Drug Toxicity	Representative	Treatment)			
		Gastric Ulcer Perforation		(Omeprazole)	C		
		Haematoma		Disopyramide	C		
		Medication Error		Bromazepam	C		
		Visual Disturbance		Oxazepam	C		
				Progesterone	C		
				Estradiol	C		
				Furosemide	C		

Date:01/17/03ISR Number: 4044197-4Report Type:Expedited (15-DaCompany Report #2003001026
 Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Stupor	Foreign	Zoloft (Sertraline)	PS		ORAL
50 MG, ORAL							

Hospitalization - Initial or Prolonged 300 MG	Health Professional	Neurontin (Gabapentin)	SS	ORAL
(DAILY), ORAL				
TOPICAL	TOPICAL	Fentanyl	SS	
50 MG		Prednisone	SS	ORAL
(DAILY), ORAL				
		Carbamazepine	C	
		Metamizole Sodium	C	
		Baclofen	C	
		Metoprolol Succinate	C	
		Valsartan	C	
		Furosemide	C	
		Thiamazole	C	
		Valoron N	C	
		Amitriptyline	C	

Date:01/17/03ISR Number: 4044208-6Report Type:Expedited (15-DaCompany Report #2002062399
Age:53 YR Gender:Female I/FU:F

Outcome	PT
Death	Abdominal Pain Upper
Other	Abnormal Behaviour

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Compartment Syndrome Confusional State Diplopia	Report Source	Product	Role	Manufacturer	Route
UNKNOWN	(TID), ORAL	Drug Toxicity Gastric Perforation	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
UNKNOWN	(DAILY), ORAL	Haematoma Medication Error Visual Acuity Reduced	Professional Company Representative				ORAL
				(Antiulcerous Treatment) (Omeprazole) Disopyramide Bromazepam Oxazepam Progesterone Estradiol Furosemide	C C C C C C C		

Date:01/17/03ISR Number: 4044537-6Report Type:Expedited (15-DaCompany Report #2002071675
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Condition Aggravated Night Blindness	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:01/17/03ISR Number: 4044540-6Report Type:Expedited (15-DaCompany Report #2002064861
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, THREE		Bronchopneumonia Cervical Vertebral Fracture	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Joint Dislocation	Company		
Liver Function Test	Representative	Morphine	C
Abnormal		Cyclizine	C
Nerve Root Injury		Metoclopramide	C
Post Procedural		Ondansetron	C
Complication		Paracetamol	C
Pseudomonas Infection		Diclofenac	C
		Fentanyl	C
		Propofol	C
		Omeprazole	C
		Combivent	C
		Clonidine	C
		Amoxicillin/Clavulan	
		ate Acid	C
		Salbutamol	C
		Heparin-Fraction,	
		Sodium Salt	C
		E1-4	C
		Allopurinol	C
		Colchicine	C
		Ciprofloxacin	C
		Potassium Chloride	C
		Nifedipine	C
		Droperidol	C
		Ranitidine	C
		Morphine	C

Freedom Of Information (FOI) Report

Codeine Phosphate	C
Omeprazole	C
Ranitidine	C
Sodium Chloride	C
Fluticasone	
Propionate	C
Olive Oil	C
Coloxyl With Senna	C
Bendroflumethiazide	C
Eze Bar	C
Egopsoryl Ta	C
Clotrimazole	C
Chloramphenicol	C
Lactulose	C
Potassium Chloride	C
Cefradine	C
Clindamycin	C
Pip/Tazo	C
Lidocaine	C
Ciprofloxacin	C
Enalapril	C

Date:01/17/03ISR Number: 4044570-4Report Type:Expedited (15-DaCompany Report #2002068411

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 5600 MG (800 Other MG, 7 TIMES A DAY),		Unevaluable Event	Consumer	Neurontin (Gabapentin)	PS		
200 MG (100 MG, BID)				Zoloft (Sertraline)	SS		
200 MG (100 MG, BID)				Trazodone Hydrochloride	SS		
1 MG (0.25				Alprazolam	SS		

MG, QID)

Vicodin

SS

15/1500 MG

(DOSE

STRENGTH

5/500 MG)

(TID)

Ibuprofen

SS

1000 MG (200

MG, FIVE

TIMES A DAY)

Acetylsalicylic Acid SS

162 MG (81

MG, BID)

Date:01/21/03ISR Number: 4044000-2Report Type:Expedited (15-DaCompany Report #2003000420

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Congenital Cystic Kidney Disease	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID),		Maternal Drugs Affecting					
ORAL		Foetus		Fexofenadine Hydrochloride	C		
				Multivitamins	C		
				Folic Acid	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/21/03ISR Number: 4044014-2Report Type:Expedited (15-DaCompany Report #2002066912
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bronchospasm Drug Dependence	Consumer Health	Neurontin (Gabapentin)	PS		
900 MG (QD), UNKNOWN (TWICE DAILY IN MORNING)		Dry Mouth	Professional				
		Nausea		Lorazepam	SS		
		Pain					
		Speech Disorder Weight Increased		Paracetamol	C		

Date:01/21/03ISR Number: 4045587-6Report Type:Expedited (15-DaCompany Report #2003001595
 Age:16 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin (Gabapentin)	PS		

Date:01/21/03ISR Number: 4045641-9Report Type:Expedited (15-DaCompany Report #2003000948
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain Dizziness	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
500 MG (THREE TIMES DAILY), ORAL		Dyspepsia					
		Dyspnoea					
		Headache Heart Rate Decreased Renal Pain Syncope Syncope Vasovagal					

Date:01/21/03ISR Number: 4046137-0Report Type:Expedited (15-DaCompany Report #2002055658
Age:34 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG(400 Other MG, TID)	Abscess Limb Hepatitis	Foreign Health	Neurontin (Gabapentin)	PS		
1200 MG (400 MG , TID)	Leukocytosis Lung Disorder Overdose Pyrexia Systemic Inflammatory Response Syndrome	Professional Company Representative	Carbamazepine	SS		

Date:01/23/03ISR Number: 4046026-1Report Type:Expedited (15-DaCompany Report #2003001248
Age:55 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Feeling Abnormal Lethargy	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Sertraline Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/23/03ISR Number: 4046027-3Report Type:Expedited (15-DaCompany Report #2003001198

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dementia	Consumer	Neurontin			
		Diabetes Mellitus		(Gabapentin)	PS		
		Hypertension		Metoprolol Tartrate	SS		
				Digoxin	SS		
				Isosorbide Dinitrate	SS		
				Furosemide	SS		
				Captopril	SS		
				Warfarin Sodium	C		
				Lorazepam	C		
				Quetiapine Fumarate	C		

Date:01/23/03ISR Number: 4046186-2Report Type:Expedited (15-DaCompany Report #2003001247

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatitis Cholestatic	Foreign Health	Neurontin			
			Professional	(Gabapentin)	PS		
900 MG,							
(ORAL)			Company Representative	Aciclovir	C		
				Metamizole Sodium	C		

Date:01/23/03ISR Number: 4046187-4Report Type:Expedited (15-DaCompany Report #2003001003

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Epilepsy	Foreign Health	Neurontin			
		Gastric Ulcer Perforation		(Gabapentin)	PS		
1200 MG (400			Professional				
MG, TID)							

Outcome PT
Disability Abdominal Distension
Aggression
Asthenia
Back Pain
Blood Cholesterol
Increased
Blood Triglycerides
Increased
Bone Neoplasm
Cardiovascular Disorder
Chest Pain
Condition Aggravated
Constipation
Depression
Difficulty In Walking
Dizziness
Dyspnoea
Fall
Hair Growth Abnormal
Infection
Irritability
Nasal Congestion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
800 MG (400 MG, BID), ORAL		Nasal Disorder Neck Pain Nervous System Disorder Neuropathy Peripheral Oedema Oedema Peripheral Pain Psychotic Disorder Sciatica Tremor Weight Increased	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
				Amitriptyline Hydrochloride Diazepam Sertraline Venlafaxine Alprazolam Acetylsalicylic Acid	C C C C C		

Date:01/23/03ISR Number: 4047022-0Report Type:Expedited (15-DaCompany Report #2003001275
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Other		Blood Potassium Decreased Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Olanzapine Fluoxetine Hydrochloride Quetiapine Fumarate Rofecoxib	C C C C		

Date:01/23/03ISR Number: 4047057-8Report Type:Expedited (15-DaCompany Report #2002068085
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (QD), ORAL		Asthenia Depression Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL

Suicidal Ideation

Docetaxel	C
Zoledronic Acid	C
Terazosin	
Hydrochloride	C
Hyzaar	C
Potassium Chloride	C
Ranitidine	
Hydrochloride	C
Naproxen	C
Acetylsalicylic Acid	C
Vitamins	C
Amino Acids	C

Date:01/23/03ISR Number: 4047061-XReport Type:Expedited (15-DaCompany Report #2002068084
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Depression	Consumer	Neurontin			
Other		Drug Ineffective		(Gabapentin)	PS		
		Neoplasm Malignant					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/23/03ISR Number: 4047067-0Report Type:Expedited (15-DaCompany Report #2003001446
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Drug Interaction	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		International Normalised					
ORAL		Ratio Decreased	Professional	Warfarin	SS		ORAL
ORAL							

Date:01/23/03ISR Number: 4047069-4Report Type:Expedited (15-DaCompany Report #2002055082
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Narcolepsy Syncope	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (THREE TIMES DAILY)			Professional Company Representative	Serotonin Antagonists All Other Therapeutic Products Amitriptyline Hydrochloride Morphine Sulfate Levothyroxine Sodium Pindolol Diazepam Estradiol Valerate Provella-14 Nitrazepam Periciazine	C C C C C C C C C C C		
ORAL							

Date:01/23/03ISR Number: 4047179-1Report Type:Expedited (15-DaCompany Report #A108646
Age:25 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - ORAL	Drug Ineffective	Consumer	Geodon(Ziprasidone)	PS	ORAL
Initial or Prolonged	Mania	Health	Neurontin		
3600 MG	Medication Error	Professional	(Gabapentin)	SS	
(TID);	Schizophrenia				
UNKNOWN			Quetiapine Fumarate	SS	ORAL
300 MG					
(DAILY); ORAL			Ethanol	SS	ORAL
(UNKNOWN),					
ORAL			Valproate Semisodium	C	
			Clonazepam	C	

Date:01/23/03ISR Number: 4047181-XReport Type:Expedited (15-DaCompany Report #2003001096
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Angioneurotic Oedema	Consumer	Neurontin(Gabapentin			
Initial or Prolonged	Fibrin D Dimer Increased)	PS		ORAL
1200 MG (400	Hypothermia					
MG, TID);	Hypothyroidism					
ORAL	Monocyte Percentage		Triamcinolone			
	Increased		Acetonide	C		
	Pancytopenia		Benzatropine			
			Mesilate	C		

Freedom Of Information (FOI) Report

Valproate Semisodium C
Risperidone C

Date:01/23/03ISR Number: 4047246-2Report Type:Expedited (15-DaCompany Report #L03-USA-00138-09
Age:55 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Literature Health Professional	Diltiazem Atenolol (Atenolol) Gabapentin	PS SS SS		

Date:01/23/03ISR Number: 4047296-6Report Type:Expedited (15-DaCompany Report #L03-USA-00138-09
Age:55 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Literature Health Professional	Diltiazem Atenolol Gabapentin	PS SS SS		

Date:01/23/03ISR Number: 4047535-1Report Type:Expedited (15-DaCompany Report #2003001277
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3600 MG (DAILY), ORAL		Alanine Aminotransferase Increased Aspartate Aminotransferase Increased Blood Alkaline Phosphatase Increased Gamma-Glutamyltransferase Increased	Foreign Health Professional	Neurontin (Gabapentin) Clodronate Disodium Diclofenac Omeprazole Anastrozole Oxycodone Hydrochloride ..	PS C C C C C C		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Asthenia		Didanosine	PS	Bristol-Myers Squibb Company	ORAL
Hospitalization - Initial or Prolonged		Cyanosis		Stavudine	SS	Bristol-Myers Squibb Company	ORAL
		Dyspnoea		Abacavir	SS		ORAL
		Lactic Acidosis		T-20/Ro 29-9800	SS		
		Myalgia		Gabapentin	SS		
		Neuropathy Peripheral		Lopinavir	C		
				Ritonavir	C		
SUBCUTANEOUS							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dizziness	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (THREE		Orthostatic Hypotension					
Other		Pyrexia	Professional				
TIMES DAILY),		Syncope					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Insulin Human C
 Insulin Human
 Injection, Isophane C

Date:01/24/03ISR Number: 4047737-4Report Type:Expedited (15-DaCompany Report #2003000982
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain	Foreign	Neurontin			
		Disease Recurrence	Health	(Gabapentin)	PS		ORAL
ORAL		Drug Effect Decreased	Professional	Nsaid'S	SS		
		Dyspepsia					
		Headache					
		Influenza					
		Multiple Sclerosis					
		Trigeminal Neuralgia					
		Weight Increased					

Date:01/24/03ISR Number: 4047794-5Report Type:Expedited (15-DaCompany Report #2002063521
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Hypertension	Consumer	Neurontin			
Other			Health	(Gabapentin)	PS		ORAL
2400 MG							
(TWICE			Professional				
DAILY), ORAL							

All Other
 Therapeutic Products C
 Dextroprpoxyphene C
 Loratadine C
 Esomeprazole C
 Paroxetine
 Hydrochloride C
 Celecoxib C
 Clonazepam C
 Propacet C
 Tramadol

Date:01/27/03ISR Number: 4046823-2Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-11743036

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose		Glucovance Tabs	PS	Bristol-Myers Squibb Company	ORAL
Other unknown							
quantity				Trazodone Hcl Tabs	SS	Apothecon	ORAL
				Neurontin	SS		
				Xanax	SS		
				Darvocet	SS		
				Phenergan + Codeine	SS		

2 oz

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/27/03ISR Number: 4048097-5Report Type:Expedited (15-DaCompany Report #S02-FRA-01975-01
 Age:84 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 20 MG QD PO		Apraxia Balance Disorder	Foreign Health	Seropram (Citalopram Hydrobromide)	PS		ORAL
		Difficulty In Walking Fall Hypertonia	Professional Other	Neurontin (Gabapentin)	SS		
4 MG BID PO		Polyneuropathy		Galantamine	SS		ORAL
				Risperdal (Risperidone)	SS		
				Bi-Tildiem (Diltiazem Hydrochloride)	SS		
				Colchimax	C		
				Aricept (Donepezil Hydrochloride)	C		
				Anafranil (Clomipramine Hydrochloride)	C		
				Corvasal (Molsidomine)	C		
				Nitriderm Tts (Glyceryl Trinitrate)	C		
				Hept-A-Mil (Heptaminol Hydrochloride)	C		
				Temesta (Lorazepam)	C		
				Imovane (Zopiclone)	C		
				Dafalgan (Paracetamol)	C		
				Kardegic (Acetylsalicylate Lysine)	C		
				Stablon (Tianeptine)	C		

Date:01/27/03ISR Number: 4048294-9Report Type:Expedited (15-DaCompany Report #FR9402220JAN2003
 Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization - Initial or Prolonged ORAL	- YR	Eczema Prurigo	Health Professional	Temesta (Lorazepam, Tablet)	PS	ORAL
			Other	Coversyl (Perindopril, ,0)	SS	ORAL
2 MG 1X PER 1 DAY ORAL	YR			Elisor (Pravastatin Sodium, , 0)	SS	ORAL
20 MG 1X PER 1DAY ORAL	YR			Loxen (Nicardipine Hydrochloride, , 0)	SS	ORAL
100 MG 1X PER 1 DAY ORAL	YR			Mopral (Omeprazole, , 0)	SS	ORAL
20 MG 1X PER 1 DAY ORAL	YR			Neurontin (Gabapentin, , 0)	SS	ORAL
400 MG 3X PER 1 DAY ORAL	6 DAY			Neurontin (Gabapentin, , 0)	SS	ORAL
600 MG 3X PER 1 DAY ORAL	9 DAY					

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Freedom Of Information (FOI) Report

Mediatensyl
 (Urapidil) C
 Sotalex (Sotalol
 Hydrochloride) C
 Aricept (Donepezil
 Hydrochloride) C
 Plavix (Clopidogrel
 Sulfate) C

Date:01/27/03ISR Number: 4048620-0Report Type:Expedited (15-DaCompany Report #EMADSS2002005362
 Age:84 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 8 MG, DAILY, ORAL	Apraxia Areflexia Cauda Equina Syndrome	Foreign Health Professional	Reminyl (Tablet) (Galantamine)	PS		ORAL
0.5 MG, DAILY, ORAL	Cerebral Atrophy Dementia Fall Gait Disturbance		Risperdal (1 Mg Tablet) (Risperidone)	SS		ORAL
1000 MG, DAILY, ORAL	Hypertonia Lacunar Infarction Pain In Extremity		Neurontin (Gabapentin)	SS		ORAL
1 CAP, DAILY, ORAL	Paraesthesia Polyneuropathy Vulvovaginal Discomfort		Colchimax (Colchimax)	SS		ORAL
20 MG, DAILY, ORAL			Seropram (Citalopram Hydrobromide)	SS		ORAL
180 MG,			Bi-Tildiem (Diltiazem Hydrochloride)	SS		ORAL

DAILY, ORAL

Nitriderm Tts (Glyceryl Trinitrate)	C
Hept-A-Myl (Heptaminol Hydrochloride)	C
Temesta (Lorazepam)	C
Imovane (Zopiclone)	C
Dafalgan (Paracetamol)	C
Kardegic (Acetylsalicylate Lysine)	C

Date:01/27/03ISR Number: 4048624-8Report Type:Expedited (15-DaCompany Report #EMADSS2002005362
Age:84 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Apraxia
Initial or Prolonged	Areflexia
	Cauda Equina Syndrome
	Cerebral Atrophy
	Dementia
	Fall
	Gait Disturbance
	Hypertonia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Lacunar Infarction Pain In Extremity Paraesthesia Polyneuropathy Vulvovaginal Discomfort	Report Source	Product	Role	Manufacturer	Route
0.5 MG,			Foreign Health Professional	Risperdal (1 Mg Tablet) (Risperidone)	PS		ORAL
DAILY, ORAL				Reminyl (Tablet) (Galantamine)	SS		ORAL
8 MG, DAILY,				Neurontn (Gabapentin)	SS		ORAL
ORAL				Colchimax(Colchimax)	SS		ORAL
1000 MG,				Seropram (Citalopram Hydrobromide)	SS		ORAL
DAILY, ORAL				Bi-Tildiem (Diltiazem Hydrochloride)	SS		ORAL
1 CAP, DAILY,				Nitriderm Tts (Glyceryl Trin-Nitrate)	C		
ORAL				Hept-A-Myl (Heptaminol Hydrochloride)	C		
20 MG, DAILY,				Temesta (Lorazepam)	C		
ORAL				Imovane (Zopiclone)	C		
180 MG,				Dafalgan (Paracetamol)	C		
DAILY, ORAL				Kardegic (Acetylsalicylate)			

Lysine)

C

Date:01/28/03ISR Number: 4048762-XReport Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 185392

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Dizziness		Atenolol 100mg Qd	PS		
Initial or Prolonged		Fall		Lisinopril 10mg Qd	SS		
				Hydrochlorothiazide 25mg Qd	SS		
				Gabapentin 1200mg Tid	SS		
				Albuterol 90/Ipratrop	C		
				Allopurinol	C		
				Atenolol	C		
				Beclomethasone	C		
				Colchicine	C		
				Flunisolide	C		
				Folic Acid	C		
				Gabapentin	C		
				Hydrochlorothiazide	C		
				Rabeprazole Na	C		
				Ranitidine Hcl	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/03ISR Number: 4048850-8Report Type:Expedited (15-DaCompany Report #2003001458

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arrhythmia Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL
	3200 MG (FOUR TIMES DAILY), ORAL	Dysphonia Fatigue Muscle Spasticity Nausea		Vitamins	C		

Date:01/28/03ISR Number: 4048852-1Report Type:Expedited (15-DaCompany Report #2003001907

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Prothrombin Time Prolonged	Consumer	Neurontin (Gabapentin)	PS		
				Potassium Phosphate Dibasic	C		
				Potassium Chloride	C		
				Filgrastim	C		
				Ranitidine	C		
				Opium Alkaloids Total	C		
				Hydrocortisone	C		
				Aciclovir	C		
				Voriconazole	C		
				Metronidazole	C		
				Diphenoxylate W/Atropine Sulfate	C		
				Vancomycin	C		
				Cirprofloxacin	C		
				Fludrocortisone Acetate	C		
				Fluticasone	C		
				Salmeterol	C		
				Folic Acid	C		
				Ipratropium Bromide	C		
				Epoetin Alfa	C		
				Pyridoxine	C		

Thiamine	C
Fentanyl	C
Paracetamol	C
Anusol	C
Diphenhydramine	C
Promethazine	C
Lorazepam	C
Morphine Sulfate	C

Date:01/28/03ISR Number: 4049074-0Report Type:Expedited (15-DaCompany Report #2003001924
 Age:73 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (DAILY),	Depression	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/03ISR Number: 4049077-6Report Type:Expedited (15-DaCompany Report #2002072453

Age:76 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG, ORAL	Ammonia Increased Condition Aggravated	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other	Hepatic Encephalopathy	Professional	Acetylsalicylic Acid Metoprolol Succinate Salutec	C C C		

Date:01/29/03ISR Number: 4048913-7Report Type:Direct Company Report #CTU 185547

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300MG PO TID Initial or Prolonged PTA	Coma Lethargy		Neurontin	PS		ORAL
			Phenobarbital	C		
			Synthroid	C		
			Citracal-D	C		
			Zocor	C		
			Neurontin	C		
			Peri Colace	C		
			Senokot	C		
			Mom	C		
			Cascara	C		
			Dulcolax	C		
			Fleet	C		

Date:01/29/03ISR Number: 4050333-6Report Type:Expedited (15-DaCompany Report #2003001908

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged (DAILY), ORAL	Amnesia Inner Ear Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other	Pain		Levothyroxine Sodium Arthrotec	C C		

Date:01/29/03ISR Number: 4050337-3Report Type:Expedited (15-DaCompany Report #2003002471

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 10 MG (DAILY), ORAL	Mental Disorder	Consumer	Lipitor (Atorvastatin)	PS		ORAL
 (TID), ORAL			Neurontin (Gabapentin)	SS		ORAL
 (DAILY)			Quetiapine Fumarate	SS		
900 MG (TID)			Carbamazepine	SS		
			Fluphenazine Hydrochloride Valproate Semisodium	SS C		

Date:01/29/03ISR Number: 4050702-4Report Type:Periodic Company Report #DSA_80010_2002

Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 2.25 G TWICE NIGHTLY PO	Drug Effect Decreased Pain	Consumer	Xyrem	PS		ORAL

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Freedom Of Information (FOI) Report

Neurontin SS

Date:01/30/03ISR Number: 4050503-7Report Type:Expedited (15-DaCompany Report #2002057832

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 900 MG (TID), Initial or Prolonged ORAL		Agranulocytosis Bone Marrow Toxicity	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Cardiac Enzymes Increased Electrocardiogram St-T Segment Elevation Erythema Infectiosum Leukopenia Myocarditis Pericardial Effusion Splenomegaly Staphylococcal Bacteraemia	Health Professional	Cefotaxime Sodium	C		

Date:01/30/03ISR Number: 4050514-1Report Type:Expedited (15-DaCompany Report #2003000982

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Back Pain Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Dyspepsia Headache Influenza Multiple Sclerosis Pain Stress Trigeminal Neuralgia Weight Increased	Professional	Nsaid'S	SS		

Date:01/30/03ISR Number: 4050537-2Report Type:Expedited (15-DaCompany Report #2002071743

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG Other (DAILY), ORAL		Asthenia Difficulty In Walking Hypoaesthesia Intervertebral Disc Protrusion Muscle Twitching Nausea Paraesthesia Sensory Disturbance	Consumer Health Professional	Neurontin (Gabapentin) Tramadol Hydrochloride Metoprolol Arthrotec	PS C C C		ORAL

Date:01/30/03ISR Number: 4050538-4Report Type:Expedited (15-DaCompany Report #2003002566
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Drug Effect Decreased Ocular Neoplasm	Health Professional	Neurontin (Gabapentin) Phenytoin Sodium Dexamethasone Hydromorphone	PS C C		ORAL

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Freedom Of Information (FOI) Report

Hydrochloride C
Fentanyl C

Date:01/30/03ISR Number: 4050540-2Report Type:Expedited (15-DaCompany Report #2003002564
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 MG, TID), ORAL		Headache Shunt Occlusion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Clonazepam	C		

Date:01/30/03ISR Number: 4050803-0Report Type:Expedited (15-DaCompany Report #2002069208
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (200 MG, TID), ORAL		Asthma Condition Aggravated Contraception Depression Dizziness Drop Attacks Electroencephalogram Abnormal Epilepsy Head Injury Hyposthenuria Hypotonia Loss Of Consciousness Pain Petit Mal Epilepsy Syncope	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
				Fluoxetine	C		
				Dosulepin	C		
				Fluticasone			
				Propionate	C		
				Salmeterol Xinafoate	C		
				Normensal	C		
				Salbutamol	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300 MG	Alanine Aminotransferase Increased	Foreign Health	Neurontin (Gabapentin)	PS		
		Aspartate Aminotransferase Increased	Professional	All Other Therapeutic Products Amitriptyline	C C		
		Blood Alkaline Phosphatase Increased Blood Creatinine Increased Blood Potassium Increased Blood Sodium Decreased Diabetes Mellitus Diabetic Hyperosmolar Coma Dyspnoea Gamma-Glutamyltransferase Increased Liver Disorder Prothrombin Time Shortened					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/30/03ISR Number: 4053465-1Report Type:Expedited (15-DaCompany Report #2003001595
Age:16 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin (Gabapentin)	PS		

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
Death		Circulatory Collapse Drug Interaction Drug Level Increased Heat Stroke		Cogentin Wellbutrin Neurontin	PS SS SS	Merck & Co., Inc	

Date:01/31/03ISR Number: 4050450-0Report Type:Expedited (15-DaCompany Report #1667783A
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Condition Aggravated Leukopenia	Foreign Health Professional	Imodium (Loperamide Hydrochloride) Capsules	PS		ORAL
PO	15 DAY			Dolpirane (R) (Paracetamol)	SS		ORAL
PO	8 DAY			Neurontin (R) Zeclar (R) Myambutol (R) Rifater (R)	SS SS SS SS		

Date:01/31/03ISR Number: 4051158-8Report Type:Expedited (15-DaCompany Report #FR9402220JAN2003
Age:80 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Eczema	Health	Temesta (Lorazepam,			

Initial or Prolonged	Professional Other	Tablet, 0) Coversyl (Perindopril)	PS SS	ORAL ORAL
2 MG 1X PER 1 DAY				
20 MG 1X PER 1 DAY		Elisor (Pravastatin Sodium)	SS	ORAL
100 MG 1X PER 1 DAY		Loxen (Nicardipine Hydrochloride)	SS	ORAL
20 MG 1X PER 1 DAY		Mopral (Omeprazole)	SS	ORAL
SEE IMAGE 9 DAY		Neurontin (Gabapentin)	SS	ORAL
		Meditensyl (Urapidil)	C	
		Sotalex (Sotalol Hydrochloride)	C	
		Aricept (Donepezil Hydrochloride)	C	
		Plavix (Clopidogrel Sulfate)	C	

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Date:01/31/03ISR Number: 4051219-3Report Type:Expedited (15-DaCompany Report #001-0945-M0100590

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 MG, TID)	Hypercapnia Lethargy Po2 Increased Respiratory Distress Respiratory Failure Wheezing	Literature Health Professional	Neurontin (Gabapentin)	PS		
			Albuterol (Salbutamol)	C		
			Ipratropium Bromide (Ipratropium Bromide)	C		
			Clonazepam (Clonazepam)	C		
			Zolpidem (Zolpidem)	C		

Date:01/31/03ISR Number: 4051306-XReport Type:Expedited (15-DaCompany Report #2003001003

Age:88 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Other 1200 MG (400 MG, TID) ORAL	Cardiac Failure Epilepsy Gastric Ulcer Perforation Malnutrition Multi-Organ Failure Pneumonia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Metoprolol	C		
			Furosemide	C		
			Allopurinol	C		
			Potassium Chloride	C		
			Levothyroxine Sodium	C		
			Clopidogrel Sulfate	C		
			Isosorbide			
			Mononitrate	C		
			Colecalciferol	C		

Date:02/04/03ISR Number: 4052260-7Report Type:Expedited (15-DaCompany Report #2002056983

Age:84 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Apraxia Fall	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other 8 MG (4 MG, TWICE DAILY), ORAL (DAILY), ORAL		Gait Disturbance Hypertonia Polyneuropathy	Professional	Galantamine	SS		ORAL
20 MG (DAILY), ORAL				Colchimax	SS		ORAL
0.5 MG (DAILY), ORAL				Citalopram Hydrobromide	SS		ORAL
180 MG (90 MG, TWICE DAILY), ORAL				Risperidone	SS		ORAL
				Diltiazem Hydrochloride	SS		ORAL
				Glyceryl Trinitrate	C		
				Heptaminol Hydrochloride	C		
				Lorazepam	C		
				Zopiclone	C		
				Paracetamol	C		
				Acetylsalicylate	C		
				Lysine	C		

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Clomipramine
 Hydrochloride C
 Molsidomine C
 Donepezil
 Hydrochloride C
 Tianeptine C

Date:02/04/03ISR Number: 4052264-4Report Type:Expedited (15-DaCompany Report #2002056669
 Age:43 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG Other (DAILY), ORAL	Coma Hepatitis Leukopenia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL (BID), ORAL (BID), ORAL	Neutropenia Tuberculosis		Ganciclovir Nevirapine Kaletra	SS SS SS		ORAL ORAL ORAL
(DAILY), ORAL (BID), ORAL			Tenofovir Disoproxil Fumarate Valganciclovir	SS SS		ORAL ORAL
			Paracetamol Loperamide Rifampicin Ethambutol Dihydrochloride Clarithromycin Bactrim Calcium Folate Filgrastim	C C C C C C C C		

Date:02/04/03ISR Number: 4052349-2Report Type:Expedited (15-DaCompany Report #2003002927
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Other Drug Abuser Health Professional Neurontin (Gabapentin) PS ORAL

2100 MG

(DAILY), ORAL

Olanzapine C
Paroxetine C
Hydrochloride C
Valdecoxib C

Date:02/04/03ISR Number: 4052405-9Report Type:Expedited (15-DaCompany Report #2003002971

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Gingivitis	Foreign	Neurontin			
		Pyrexia	Consumer	(Gabapentin)	PS		ORAL
300 MG (100		Shock					
MG,TID), ORAL							

Date:02/04/03ISR Number: 4052413-8Report Type:Expedited (15-DaCompany Report #2003002866

Age:46 YR Gender:Female I/FU:I

Outcome PT
Other Finger Amputation
Liver Function Test
Abnormal

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Platelet Count Increased
Thrombosis

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/04/03ISR Number: 4052648-4Report Type:Expedited (15-DaCompany Report #2003001498
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG (DAILY), ORAL		Back Pain Drug Intolerance Eyelid Oedema Food Allergy Malaise Spinal Column Stenosis	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:02/04/03ISR Number: 4052741-6Report Type:Expedited (15-DaCompany Report #2003002473
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiac Arrest	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Paroxetine Hydrochloride Lorazepam Levothyroxine Sodium	PS C C C		

Date:02/04/03ISR Number: 4052949-XReport Type:Expedited (15-DaCompany Report #2003003300
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged ORAL	Cytolytic Hepatitis Depression	Foreign Health	Neurontin (Gabapentin)	PS	ORAL
Other INTRAVENOUS	Disease Recurrence 175 MG/M2	Professional	Paclitaxel	SS	
(EVERY THREE WEEKS),	Frequent Bowel Movements Gastrointestinal Disorder				
INTRAVENOUS	General Physical Health Deterioration Hypokalaemia Liver Disorder Neutropenia Orthostatic Hypotension Paraesthesia Pericardial Effusion Pleural Effusion Renal Failure Acute Suicide Attempt Treatment Noncompliance				

Freedom Of Information (FOI) Report

Date:02/04/03ISR Number: 4052954-3Report Type:Expedited (15-DaCompany Report #001-0719-M0100407

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 1200 MG (600 Initial or Prolonged MG, BID), Disability ORAL Other 1600 MG (400 MG, 2 CAPSULES IN AM AND PM) ONE EVERY 5 MIN., MAX: 3 (PRN) 0.4 MG (DAILY)	Anhedonia Anxiety Atherosclerosis Back Pain Blood Cholesterol Increased Blood Pressure Increased Blood Triglycerides Increased Blood Urea Increased Body Temperature Increased Cardiac Disorder Cerebrovascular Accident Condyloma Acuminatum Coronary Artery Disease Coronary Artery Stenosis Diabetes Mellitus Drug Interaction Headache Heart Rate Increased High Density Lipoprotein Decreased Hypercholesterolaemia Hyperlipidaemia Hypertension Hypothyroidism Intermittent Claudication Ischaemia Lipids Increased	Consumer Health Professional	Lopid (Gemfibrozil) Neurontin (Gabapentin) Nitroglycerin (Glyceryl Trinitrate) Cerivastatin (Cerivastatin)	PS SS SS		ORAL

Myocardial Infarction
 Neuropathy Peripheral
 Palpitations
 Peptic Ulcer
 Peripheral Vascular
 Disorder
 Pneumonia
 Pruritus
 Pulse Abnormal
 Rhabdomyolysis
 Sinusitis
 Ventricular Extrasystoles

Date:02/05/03ISR Number: 4053372-4Report Type:Expedited (15-DaCompany Report #2003003392
 Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Upper Defaecation Urgency	Health Professional	Neurontin (Gabapentin)	PS		ORAL
6000 MG (DAILY), ORAL		Diarrhoea					
37.5 MG (DAILY), ORAL		Drug Effect Decreased Frequent Bowel Movements		Venlafaxine Hydrochloride	SS		ORAL
				Vicodin Carisoprodol	C C		

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Date:02/05/03ISR Number: 4053792-8Report Type:Expedited (15-DaCompany Report #2002072435

Age:36 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 Other MG, THREE TIMES DAILY), ORAL	White Blood Cell Count Decreased	Foreign Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL

Date:02/05/03ISR Number: 4053821-1Report Type:Expedited (15-DaCompany Report #2003003481

Age:48 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 80 MG Other (DAILY), ORAL 150 MG (DAILY), ORAL 900 MG (DAILY), ORAL 800 MG (DAILY), ORAL	Aspiration Bradyphrenia Coronary Artery Disease Depressed Level Of Consciousness Depression Disturbance In Attention Drug Interaction Drug Level Above Therapeutic General Physical Health Deterioration Lobar Pneumonia Neuropathy Peripheral Pulmonary Embolism Somnolence	Foreign Health Professional Company Representative	Lipitor (Atorvastatin) Zoloft (Sertraline) Gabapentin (Gabapentin) Carbamazepine Metformin Gliclazide Diltiazem Candesartan Atenolol Isosorbide	PS SS SS C C C C C C		ORAL ORAL ORAL ORAL

Date:02/06/03ISR Number: 4055737-3Report Type:Expedited (15-DaCompany Report #2002CG00895
 Age:91 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 25 MG DAILY	Hepatitis Cholestatic	Foreign	Tenormine	PS		ORAL
Initial or Prolonged PO	Hepatocellular Damage	Health				
600 MG DAILY	Pleural Effusion	Professional	Neurontin	SS		ORAL
PO		Other				
			Coumadine	SS		
25 MG DAILY			Aldactone	SS		ORAL
PO						
0.125 MG			Hemigoxine Nativelle	SS		ORAL
DAILY PO						
20 MG DAILY			Lasilix	SS		ORAL
PO						

Date:02/07/03ISR Number: 4053840-5Report Type:Periodic Company Report #2002107546US
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 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			

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one Hydrochloride) SS
 Paxil(Paroxetine
 Hydrochloride) SS
 Klonopin(Clonazepam) SS
 Oxycodone
 Hydrochloride
 (Oxycodone
 Hydrochloride) C
 Progesterone C
 Salbutamol(Salbutamo
 l) C
 Pentosan
 Polysulfate(Pentosan
 Polysulfate) C
 Hydroxyzine(Hydroxyz
 ine) C
 Amitriptyline C

Date:02/07/03ISR Number: 4053895-8Report Type:Expedited (15-DaCompany Report #2002071335
 Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (QD), Other ORAL		Ascites Asthenia Biliary Cirrhosis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Blood Pressure Decreased Cholelithiasis Coordination Abnormal Dizziness Faeces Discoloured Fall Gait Disturbance Gallbladder Disorder Gastrointestinal Haemorrhage Haematocrit Decreased Haemoglobin Decreased Head Injury Heart Rate Increased Hepatitis C Laceration Occult Blood Positive Platelet Count Decreased		Amitriptyline Hydrochloride Glibomet Thioctic Acid	C C C		

Portal Hypertension
Red Blood Cell Count
Decreased
Splenomegaly
Syncope
Varices Oesophageal
Vomiting
Weight Decreased

Date:02/07/03ISR Number: 4053897-1Report Type:Expedited (15-DaCompany Report #2002068084

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Depression	Consumer	Neurontin			
Other		Drug Ineffective		(Gabapentin)	PS		

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Date:02/07/03ISR Number: 4053900-9Report Type:Expedited (15-DaCompany Report #2003004307

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coma	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/07/03ISR Number: 4053955-1Report Type:Expedited (15-DaCompany Report #2003004705

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Disorder	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:02/07/03ISR Number: 4053969-1Report Type:Expedited (15-DaCompany Report #2003003630

Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bladder Cancer Kidney Infection	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (TID),							
ORAL							

Amlodipine Besilate	C
Simvastatin	C
Lorazepam	C
Hydrochlorothiazide	C
Acetylsalicylic Acid	C
Tocopherol	C
Ascorbic Acid	C
Retinol	C
Calcium	C
Lactobacillus Acidophilus	C

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Acute Sinusitis
Initial or Prolonged	Anxiety
Disability	Back Pain
Other	Bladder Disorder
	Blood Alkaline
	Phosphatase Increased
	Brain Neoplasm
	Bronchitis Acute
	Cholelithiasis
	Colitis
	Depression
	Diverticulum Intestinal
	Ear Pain
	Electroencephalogram
	Abnormal
	Fatigue
	Gastrointestinal Disorder
	Hepatic Steatosis
	Insomnia

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (BID),		Monocyte Count Increased Myalgia Obstructive Airways	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Plantar Fasciitis	Professional				
		Scar Pain	Company Representative	Fluoxetine Hydrochloride	C		
				Omeprazole	C		
				Capsaicin	C		
				Bupirone			
				Hydrochloride	C		

Date:02/07/03ISR Number: 4054183-6Report Type:Expedited (15-DaCompany Report #033-0945-M0100075
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE IMAGE		Asthenia Cholecystitis	Foreign Literature	Neurontin(Gabapentin)	PS		
Other		Dyspnoea Encephalitis Eosinophilia Epilepsy Fatigue Hepatic Trauma Hypersensitivity Monoparesis Oedema Peripheral	Health Professional				

Date:02/07/03ISR Number: 4054188-5Report Type:Expedited (15-DaCompany Report #2003002866
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arterial Thrombosis Limb Finger Amputation	Foreign Health	Neurontin(Gabapentin)	PS		
Other		Liver Disorder Liver Function Test	Professional Company				

Abnormal
Platelet Count Increased

Representative

Date:02/07/03ISR Number: 4054205-2Report Type:Expedited (15-DaCompany Report #2003004340
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea	Foreign	Neurontin(Gabapentin			
600 MG		Therapeutic Response	Health)	PS		ORAL
(DAILY), ORAL		Increased	Professional				
		Tremor		Lithium	SS		
				Lithium Carbonate	C		

Date:02/07/03ISR Number: 4054210-6Report Type:Expedited (15-DaCompany Report #2003004707
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Disorder	Foreign	Neurontin			
ORAL			Health	(Gabapentin)	PS		ORAL
			Professional				

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Date:02/07/03ISR Number: 4054213-1Report Type:Expedited (15-DaCompany Report #2003004704

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Disorder	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Professional				

Date:02/07/03ISR Number: 4054238-6Report Type:Expedited (15-DaCompany Report #033-0945-M0200094

Age:91 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Hepatitis Cholestatic	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Hospitalization - 600 MG		Hepatocellular Damage					
Initial or Prolonged (DAILY), ORAL		Organ Failure	Professional				
Other				Spironolactone	SS		ORAL
25 MG							
(DAILY), ORAL							
ORAL				Digoxin	SS		ORAL
ORAL				Warfarin	SS		ORAL
ORAL				Furosemide	SS		ORAL
ORAL				Atenolol	SS		ORAL

Date:02/07/03ISR Number: 4054241-6Report Type:Expedited (15-DaCompany Report #2003002473

Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Pericardial Haemorrhage	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Hospitalization - 300 MG		Torsade De Pointes					
Initial or Prolonged (DAILY), ORAL			Professional				

Other

Company
Representative

Paroxetine
Hydrochloride C
Lorazepam C
Levothyroxine Sodium C
Lactobacillus
Acidophilus C

Date:02/07/03ISR Number: 4054243-XReport Type:Expedited (15-DaCompany Report #044-0945-M0100088

Age: Gender:Not Specified/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Hypoglycaemia	Foreign Study Health Professional	Gabapentin (Gabapentin)	PS		

Date:02/07/03ISR Number: 4054246-5Report Type:Expedited (15-DaCompany Report #2003004338

Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 300 MG (DAILY), ORAL		Clonic Convulsion Lower Respiratory Tract Infection Respiratory Distress Sedation Tremor	Foreign Health Professional	Gabapentin (Gabapentin) Morphine Sulfate Fentanyl Paracetamol Domperidone Benzatropine Mesilate	PS C C C C C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/07/03ISR Number: 4068957-9Report Type:Periodic
Age:48 YR Gender:Male I/FU:I

Company Report #2002135208US

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorgasmia	Health	Celebrex (Celecoxib)			
ORAL		Constipation	Professional	Capsule	PS		ORAL
		Drug Ineffective		Neurontin (Gabapentin)	SS		ORAL
600 MG, BID,							
ORAL; SEE							
IMAGE							

Date:02/10/03ISR Number: 4055087-5Report Type:Expedited (15-DaCompany Report #2003003075
Age:50 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Completed Suicide	Literature	Gabapentin			
Death		Intentional Misuse	Health	(Gabapentin)	PS		ORAL
ORAL			Professional	Tramadol	SS		ORAL
ORAL				Rofecoxib	SS		ORAL
ORAL							

Date:02/10/03ISR Number: 4055100-5Report Type:Expedited (15-DaCompany Report #2003003069
Age:52 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Completed Suicide	Literature	Gabapentin(Gabapenti			
Death		Intentional Misuse	Health	n)	PS		ORAL
ORAL			Professional	Oxycodone	SS		ORAL
ORAL							

Date:02/10/03ISR Number: 4055101-7Report Type:Expedited (15-DaCompany Report #2003003079

Age:36 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature Health	Gabapentin(Gabapentin)	PS		ORAL
ORAL		Intentional Misuse	Professional	Chloral Hydrate	SS		ORAL
ORAL				Codeine	SS		ORAL

Date:02/10/03ISR Number: 4055102-9Report Type:Expedited (15-DaCompany Report #2003003076

Age:38 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health	Gabapentin(Gabapentin)	PS		ORAL
ORAL		Intentional Misuse	Professional	Amoxapine	SS		ORAL
ORAL				Quetiapine	SS		ORAL

Date:02/10/03ISR Number: 4055103-0Report Type:Expedited (15-DaCompany Report #2002065565

Age:55 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Consumer	Gabapentin(Gabapentin)	PS		ORAL
ORAL		Intentional Misuse	Health	Diltiazem(Diltiazem)	SS		ORAL
ORAL			Professional	Atenolol	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/10/03ISR Number: 4055287-4Report Type:Expedited (15-DaCompany Report #2002070374

Age:87 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG (100 Other MG, TID), ORAL	Neutropenia White Blood Cell Count Decreased	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
			Atorvastatin	C		
			Quinine Bisulfate	C		
			Tramadol	C		
			Flucloxacillin	C		
			Gliclazide	C		
			Furosemide	C		
			Diltiazem	C		
			Acetylsalicylic Acid	C		
			Prochlorperazine	C		
			Temazepam	C		
			Predonium	C		
			Amoxicillin	C		
			Azithromycin	C		
			Cefalotin	C		
			Metformin			
			Hydrochloride	C		
			Gliclazide	C		
			Tramadol			
			Hydrochloride	C		

Date:02/11/03ISR Number: 4055900-1Report Type:Expedited (15-DaCompany Report #2003003667

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (TWICE DAILY), ORAL	Convulsion Drug Ineffective Increased Appetite	Consumer	Neurontin (Gabapentin)	PS		ORAL
	Nervousness Weight Increased		Carbamazepine	C		

Date:02/11/03ISR Number: 4056205-5Report Type:Expedited (15-DaCompany Report #2003003067
Age:20 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Gabapentin			
ORAL		Completed Suicide	Health	(Gabapentin)	PS		ORAL
		Intentional Misuse	Professional	Fentanyl	SS		
ORAL				Dextropropoxyphene	SS		ORAL

Date:02/11/03ISR Number: 4056295-XReport Type:Expedited (15-DaCompany Report #2003003074
Age:31 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Gabapentin			
ORAL		Overdose	Health	(Gabapentin)	PS		ORAL
			Professional	Risperidone	SS		ORAL
ORAL				Benzatropine			
ORAL				Mesilate	SS		ORAL
ORAL				All Other			
				Therapeutic Products	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/11/03ISR Number: 4056644-2Report Type:Expedited (15-DaCompany Report #2003002971

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Foreign	Neurontin(Gabapentin			
300 MG (100		Gingivitis	Consumer)	PS		ORAL
MG, TID),		Pyrexia					
ORAL		Shock					

Date:02/12/03ISR Number: 4054729-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0386384A

Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agranulocytosis		Lamictal	PS	Glaxosmithkline	ORAL
2MG Four		Bradycardia		Gabitril	SS		ORAL
times per day		Hypotension					
		Thrombocytopenia		Neurontin	SS		ORAL
				Lithium	C	Glaxosmithkline	
				Sonata	C		
				Claritin	C		
				Multivitamins	C		
				Colace	C		

Date:02/12/03ISR Number: 4055694-XReport Type:Direct

Company Report #CTU 186504

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Apnoea		Neurontin Toxicity	PS		
Hospitalization -		Bradycardia					
Initial or Prolonged		Diabetes Mellitus					
Required		Dialysis					
Intervention to		Drug Level Increased					
Prevent Permanent		Hypoxia					
Impairment/Damage		Renal Failure Chronic					

Date:02/12/03ISR Number: 4056876-3Report Type:Expedited (15-DaCompany Report #2003004387
Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Affective Disorder	Foreign	Neurontin			
Initial or Prolonged	Anxiety Disorder	Health	(Gabapentin)	PS		ORAL
300 MG, ORAL						
Disability		Professional Company Representative				

Date:02/13/03ISR Number: 4057456-6Report Type:Expedited (15-DaCompany Report #PHHO2002PL09384
Age:58 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Dizziness
Disability	Drug Ineffective Dyskinesia Extrapyramidal Disorder Hypotonia Malignant Neoplasm Progression Purpura

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Rash Syncope Tremor	Report Source	Product	Role	Manufacturer	Route
400 MG, QD, ORAL; 300 MG,BID, UNK			Foreign Study Health Professional Other	Glivec(Sti571/Cgp571 48 T35717+Caps) Capusle	PS		ORAL
30 MG, TID, ORAL				Durogesic (Fentanyl) Mst(Morphine Sulfate)	SS SS		ORAL
300 MG, TID, ORAL				Neurontin (Gabapentin)	SS		ORAL
2 X 5MG DAILY				Transene (Clorazepate Dipotassium)	SS		

Date:02/13/03ISR Number: 4058127-2Report Type:Expedited (15-DaCompany Report #2003005167
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Ammonia Increased Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other		Encephalopathy Oesophageal Varices Haemorrhage		Beta Blocking Agents	C		

Date:02/14/03ISR Number: 4060505-2Report Type:Expedited (15-DaCompany Report #331015
Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Eczema Pruritus Rash Erythematous	Foreign Other	Loxen Lp (Nicardipine Hydrochloride)	PS		ORAL
100 MG DAILY							
ORAL				Neurontin (Gabapentin)	SS		ORAL
400 MG 3 PER DAY ORAL							
ORAL				Temesta (Lorazepam)	SS		ORAL
20 MG DAILY				Mopral (Omeprazole)	SS		ORAL
ORAL							
2 MG DAILY				Coversyl (Perindopril Erbumine)	SS		ORAL
ORAL							
20 MG DAILY				Drug Uae For Unknown Indication	SS		ORAL
ORAL							
				Urapidil (Urapidil)	C		
				Sotalol Hydrochloride (Sotalol Hydrochloride)	C		
				Donepezil Hydrochloride (Donepezil)	C		
				Clopidogrel			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Bisulfate
(Clopidogrel
Bisulfate) C

Date:02/14/03ISR Number: 4060549-0Report Type:Expedited (15-DaCompany Report #2003005556
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL			Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:02/14/03ISR Number: 4060550-7Report Type:Expedited (15-DaCompany Report #2003005558
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL			Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:02/14/03ISR Number: 4061030-5Report Type:Expedited (15-DaCompany Report #2003005280
Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
Other				Escitalopram	SS		
				Oxcarbazepine	SS		
				Trazodone	SS		
				Fluoxetine Hydrochloride	SS		
				Insulin	C		

Date:02/14/03ISR Number: 4061118-9Report Type:Expedited (15-DaCompany Report #2002070768

Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG (300 Other MD, TID), ORAL	Drug Effect Decreased Treatment Noncompliance	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Pancrelipase Citalopram Hydrobromide	C C		

Date:02/18/03ISR Number: 4056341-3Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-12176418

Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	400mg three times a day	Eczema Pruritus Rash Erythematous		Elisor Tabs 20 Mg Neurontin	PS SS	Bristol-Myers Squibb Company	ORAL ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

30-Nov-2002

through

02-Dec-2002 12 DAY

Temesta	SS	ORAL
Mopral	SS	ORAL
Coversyl	SS	ORAL
Loxen	SS	ORAL
Depakine Chrono	C	
Plavix	C	Regulatory Health Authority South Africa
Mediatensyl	C	
Sotalex	C	
Aricept	C	

Date:02/19/03ISR Number: 4056649-1Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-11937562
Age:91 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hepatitis Cholestatic		Coumadine	PS	Bristol-Myers Squibb Company	ORAL
Hospitalization - Dosage Form: Initial or Prolonged Tablet. Stop Other date given as both 05 Jun 2001 and 05 Stop date given as both 25 and 28 May 2002.		Infection Pleural Effusion		Tenormine	SS		ORAL
Treatment began in 1999	35	MON		Aldactone	SS		ORAL

or Jun 2001

or Mar 2002.

Dose: 0.125

mg.

17 DAY

Hemigoxine Nativelle SS ORAL

Lasilix SS
Neurontin SS

Date:02/19/03ISR Number: 4059317-5Report Type:Expedited (15-DaCompany Report #2003004961

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Gastric Neoplasm Syncope	Consumer	Neurontin (Gabapentin) Geodon (Ziprasidone)	PS SS		ORAL
ORAL				Lamotrigine Antihypertensives Diuretics Arpiprazole	SS C C C		

Date:02/19/03ISR Number: 4059629-5Report Type:Expedited (15-DaCompany Report #2003005473

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG		Coronary Artery Disease Myocardial Infarction	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
(UNKNOWN)				Chloral Hydrate	SS		ORAL

ORAL

250 MG

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(UNKNOWN)

ORAL

Mirtazapine

SS

ORAL

30 MG

(UNKNOWN)

ORAL

Metamizole Sodium

SS

OROPHARINGEAL 120 DROP

(UNKNOWN),

ORAL

Date:02/19/03ISR Number: 4059632-5Report Type:Expedited (15-DaCompany Report #2003001986

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG (THREE TIMES DAILY),	Chills Dizziness	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		Orthostatic Hypotension Pyrexia	Professional				
ORAL		Syncope		Insulin Human Insulin Human Injection, Isophane	C C		

Date:02/19/03ISR Number: 4059638-6Report Type:Expedited (15-DaCompany Report #061-0945-M0100035

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 1200 MG Initial or Prolonged (DAILY), ORAL		Oedema Peripheral Sepsis	Foreign Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other			Professional	Celecoxib			

200 MG

(Celecoxib)

SS

ORAL

(DAILY), ORAL

Sinemet	C
All Other	
Therapeutic Products	C
Furosemide	C
Potassium Chloride	C
Atorvastatin	C
Karvezide (Karvea	
Hct)	C
Ranitidine	
Hydrochloride	C
Clopidogrel Sulfate	C
Doxepin	
Hydrochloride	C

Date:02/19/03ISR Number: 4059822-1Report Type:Expedited (15-DaCompany Report #2002056983

Age:84 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Apraxia Cerebral Atrophy	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other 8 MG (4 MG, TWICE DAILY), ORAL	Dementia Difficulty In Walking Fall	Professional	Galantamine	SS		ORAL
(DAILY), ORAL	Gait Disturbance		Colchimax	SS		ORAL
20 MG (DAILY), ORAL	Hypertonia Hyporeflexia Pain In Extremity		Citalopram Hydrochloride	SS		ORAL
0.5 MG	Polyneuropathy		Risperidone	SS		ORAL

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Freedom Of Information (FOI) Report

(DAILY), ORAL

180 MG (90

MG, TWICE

DAILY), ORAL

Diltiazem			
Hydrochloride	SS		ORAL

Glyceryl Trinitrate	C
Heptaminol	
Hydrochloride	C
Lorazepam	C
Zopiclone	C
Paracetamol	C
Acetylsalicylate	
Lysine	C
Clomipramine	
Hydrochloride	C
Molsidomine	C
Donepezil	
Hydrochloride	C
Tianeptine	C

Date:02/19/03ISR Number: 4060471-XReport Type:Expedited (15-DaCompany Report #2003005368

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams	Consumer	Neurontin			
		Cataract		(Gabapentin)	PS		ORAL
400 MG (100		Memory Impairment					
MG, FOUR							

TIMES DAILY),

ORAL

Acetylsalicylic Acid	C
Calcium	C
Multivitamins	C

Date:02/19/03ISR Number: 4060473-3Report Type:Expedited (15-DaCompany Report #2003005367

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Unevaluable Event	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG							
(DAILY), ORAL							

Date:02/19/03ISR Number: 4060519-2Report Type:Expedited (15-DaCompany Report #2003005471
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Blindness Unilateral	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Cataract					
ORAL		Glaucoma		Pioglitazone	C		
		Lens Dislocation		Simvastatin	C		
				Amitriptyline	C		
				All Other Therapeutic Products	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/20/03ISR Number: 4057462-1Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0388885A
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	200MG Per day	Agitation		Lamictal	PS	Glaxosmithkline	ORAL
	800MG Per day	Irritability		Neurontin	SS		
		Memory Impairment		Unspecified Medications	C		

Date:02/20/03ISR Number: 4060718-XReport Type:Expedited (15-DaCompany Report #2003006077
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	48.9 GRAM	Diarrhoea Dizziness	Literature Health	Neurontin (Gabapentin)	PS		ORAL
(ONCE), ORAL		Drug Screen Positive	Professional				
INHALATION		Dysphoria		Cocaine	SS		NASAL
		Emotional Disorder Faecal Incontinence Intentional Misuse Lethargy Medication Error					

Date:02/20/03ISR Number: 4060724-5Report Type:Expedited (15-DaCompany Report #2003006078
Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	(ONCE) ORAL	Aggression Convulsion	Literature Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other	25 GRAM	Electrocardiogram Qrs	Professional	Lamotrigine	SS		ORAL
(ONCE), ORAL		Complex Abnormal					
		Electrocardiogram Qrs					

Complex Prolonged
Hypertension
Suicide Attempt
Tachycardia

Date:02/20/03ISR Number: 4060725-7Report Type:Expedited (15-DaCompany Report #2003002566

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Health	Neurontin			
		Ocular Neoplasm	Professional	(Gabapentin)	PS		ORAL
ORAL				Phenytoin Sodium	C		
				Dexamethasone	C		
				Hydromorphone			
				Hydrochloride	C		
				Fentanyl	C		

Date:02/20/03ISR Number: 4060733-6Report Type:Expedited (15-DaCompany Report #2003006079

Age:44 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Aphasia
Initial or Prolonged	Blood Alcohol Increased
	Blood Glucose Decreased
	Confusional State
	Depressed Level Of

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Consciousness Drug Level Increased Drug Screen Positive Dysarthria					
ORAL		Hyporeflexia Lethargy	Literature Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
ORAL		Medication Error	Professional	Valproic Acid	SS		ORAL
ORAL		Platelet Count Abnormal		Mexiletine	SS		ORAL
ORAL		Pupillary Reflex Impaired		Ethanol	SS		ORAL
		Pupils Unequal Somnolence Speech Disorder White Blood Cell Count Decreased					

Date:02/20/03ISR Number: 4060887-1Report Type:Expedited (15-DaCompany Report #2003001908
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Amnesia Balance Disorder	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other		Inner Ear Disorder Post Procedural Pain Surgical Procedure Repeated	Professional	Sildenafil Citrate Levothyroxine Sodium Arthrotec Testosterone	C C C C		

Date:02/20/03ISR Number: 4061182-7Report Type:Expedited (15-DaCompany Report #2003006044
Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Coma		Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL				Carbamazepine	C		

Hypnotics And
 Sedatives C
 Zopiclone C
 Citalopram
 Hydrobromide C
 Antihypertensives C
 Antibiotics C

Date:02/20/03ISR Number: 4061185-2Report Type:Expedited (15-DaCompany Report #2003002452
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 120 MG (TID), Other ORAL		Drug Interaction Neoplasm Prostate Oedema Peripheral Peripheral Vascular Disorder Prostatic Specific Antigen Increased	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Amlodipine (Amlodipine) Nicardipine Hydrochloride	PS SS SS		ORAL ORAL ORAL
10 MG (5 MG, DAILY), ORAL							
100 MG (50 MG, BID), ORAL							
				Baclofen Troxeutin Aporex	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/20/03ISR Number: 4061235-3Report Type:Expedited (15-DaCompany Report #2003006202
Age:24 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1500 MG , ORAL		Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus	Foreign Health Professional	Neurontin (Gabapentin) Amitriptyline	PS C		ORAL

Date:02/21/03ISR Number: 4060067-XReport Type:Direct Company Report #CTU 187128
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG PO TID Initial or Prolonged		Blood Creatinine Increased Clonic Convulsion Dizziness Drug Toxicity Muscle Twitching		Neurontin 300mg Mscontin Synthroid Spronolactone Zanaflex K-Dur Singuan Zoloft Ambien Uripus Detrol La Hyosycamine Prilosec Toprol Xl	PS C C C C C C C C C C C C		ORAL

Date:02/21/03ISR Number: 4062113-6Report Type:Expedited (15-DaCompany Report #PHBS2003CA01501
Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 425 MG/D		Condition Aggravated Drug Interaction	Foreign Literature	Clozapine (Clozapine) Unknown	PS		

300 MG/D

Hallucination, Auditory Insomnia	Health Professional	Gabapentin (Gabapentin)	SS
Paranoia Psychotic Disorder	Other	Procyclidine (Procyclidine) Divalproex Sodium (Valproate Semisodium) Fluoxetine (Fluoxetine)	C C C

Date:02/21/03ISR Number: 4062791-1Report Type:Expedited (15-DaCompany Report #DSA_22367_2003
Age:84 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Apraxia Dementia Difficulty In Walking Fall Gait Disturbance Hypertonia Hyporeflexia Lacunar Infarction Pain In Extremity

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Polyneuropathy

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
180 MG	PER24HR PO	Foreign Health	Bi-Tildiem	PS		ORAL
1 G PER24HR	PO	Professional	Neurontin	SS		ORAL
		Other				
8 MG PER24HR	PO		Reminyl	SS		ORAL
1 U PER24HR	PO		Colchimax	SS		ORAL
20 MG PER24HR	PO		Seropram	SS		ORAL
0.5 MG	PER24HR PO		Risperdal	SS		ORAL
			Temesta	C		
			Imovane	C		
			Dafalgan	C		
			Aricept	C		
			Anafranil	C		
			Corvasal	C		
			Nitriderm	C		
			Heptamyl	C		
			Kardegic	C		
			Stablon	C		

Date:02/24/03ISR Number: 4064995-0Report Type:Expedited (15-DaCompany Report #2003001285

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hepatic Steatosis	Foreign	Neurontin			

Pain In Extremity Health (Gabapentin) PS ORAL
(THREE TIMES DAILY), ORAL Professional

Date:02/24/03ISR Number: 4065044-0Report Type:Expedited (15-DaCompany Report #2003006318
Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Toxic Epidermal Necrolysis	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG							
(DAILY), ORAL							
INTRAVENOUS	20 MG			Methylprednisolone	SS		
(DAILY),							
INTRAVENOUS							

Date:02/24/03ISR Number: 4065045-2Report Type:Expedited (15-DaCompany Report #2002058564
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Neutropenia White Blood Cell Count Decreased	Foreign Health Professional Company Representative	Neurontin (Gabapentin) All Other Therapeutic Products	PS C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/24/03ISR Number: 4065098-1Report Type:Expedited (15-DaCompany Report #2003006153

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNKNOWN		Liver Disorder	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

(UNKNOWN),

ORAL

Date:02/24/03ISR Number: 4065404-8Report Type:Expedited (15-DaCompany Report #2003002927

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2100 MG DAILY		Drug Abuser	Health Professional	Neurontin (Gabapentin)	PS		ORAL

ORAL

Olanzapine	C
Paroxetine Hydrochloride	C
Valdecoxib	C

Date:02/25/03ISR Number: 4065221-9Report Type:Expedited (15-DaCompany Report #2003006796

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Condition Aggravated Drug Withdrawal Syndrome Epilepsy	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/25/03ISR Number: 4065223-2Report Type:Expedited (15-DaCompany Report #2003001247

Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other 900 MG (UNKNOWN), ORAL		Hepatitis Cholestatic	Foreign Health Professional Company Representative	Neurontin (Gabapetin) Aciclovir Metamizole Sodium	PS C C		ORAL

Date:02/25/03ISR Number: 4065373-0Report Type:Expedited (15-DaCompany Report #PHNU2003DE00883
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG, BID; Other 200 MG, QD		Abdominal Abscess Hypothermia Polyneuropathy Respiratory Failure	Foreign Health Professional Other	Tegretal (Carbamazepi ne) Neurontin(Gabapentin) Duragesic(Fentanyl) Patch Novalgin(Metamizole Sodium) Drops Antidepressants(No Ingredients/Substanc es)	PS SS SS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/25/03ISR Number: 4065659-XReport Type:Expedited (15-DaCompany Report #2003006906

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aphonia	Consumer	Neurontin			
ORAL		Blood Pressure Decreased		(Gabapentin)	PS		ORAL
		Cold Sweat		Topiramate	SS		
ORAL		Condition Aggravated		Naproxen	SS		ORAL
		Convulsion		Oxcarbazepine	C		
		Dizziness		Pravastatin Sodium	C		
		Dyskinesia		All Other			
		Fall		Therapeutic Products	C		
		Ill-Defined Disorder		Hydrocortisone	C		
		Limb Injury		Alendornate Sodium	C		
		Muscle Twitching		Levothyroxine Sodium	C		
		Oedema Peripheral		Lansoprazole	C		
		Painful Respiration					
		Tremor					
		Viral Infection					
		Vomiting					
		Weight Increased					

Date:02/25/03ISR Number: 4065756-9Report Type:Expedited (15-DaCompany Report #2003007155

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Health	Neurontin			
			Professional	(Gabapentin)	PS		
			Company	All Other			
			Representative	Therapeutic Products	SS		

Date:02/25/03ISR Number: 4066563-3Report Type:Expedited (15-DaCompany Report #2003006860

Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetes Mellitus	Foreign	Neurontin (Tablets)			
		Inadequate Control	Consumer	(Gabapentin)	PS		ORAL

3200 MG (1600

MG, BID), Difficulty In Walking

Headache

ORAL

Medication Error
Overdose

Metformin
Hydrochloride C
Atorvastatin C
Mirtazapine C
Glimepiride C
Asasantin C

Date:02/26/03ISR Number: 4067065-0Report Type:Expedited (15-DaCompany Report #2002060915
Age:78 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Disorder	Foreign Health	Neurontin (Tablets)(Gabapentin)			
Life-Threatening Hospitalization - 1200 MG		Grand Mal Convulsion	Professional		PS		ORAL
Initial or Prolonged (DAILY), ORAL		Hemiparesis	Company				
Disability			Representative	Glimepiride	C		
Other				Carvedilol	C		
				Furosemide	C		
				Trimipramine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/27/03ISR Number: 4067439-8Report Type:Expedited (15-DaCompany Report #2003007169

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG, TID), ORAL		Double Vessel Bypass Graft	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Dry Mouth					
		Dry Throat					
		Glossodynia Neuropathy Peripheral		All Other Non-Therapeutic Products	SS		
				All Other Non-Therapeutic Products	SS		
				All Other Therapeutic Products	C		
				Warfarin Sodium	C		

Date:02/27/03ISR Number: 4067665-8Report Type:Expedited (15-DaCompany Report #2003002927

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2100 MG (DAILY), ORAL		Drug Abuser	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Olanzapine	C		
				Paroxetine Hydrochloride	C		
				Valdecoxib	C		

Date:02/27/03ISR Number: 4067667-1Report Type:Expedited (15-DaCompany Report #2002072755

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		High Density Lipoprotein	Health	Lipitor			

Initial or Prolonged	Decreased	Professional	(Atorvastatin)	PS	
	Thrombosis	Company	Neurontin		
		Representative	(Gabapentin)	SS	ORAL
900 MG (TID),					
ORAL					
Tramadol					
Hydrochloride					
Capecitabine					
C					
C					

Date:02/27/03ISR Number: 4068219-XReport Type:Expedited (15-DaCompany Report #2003005569
 Age:87 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Confusional State	Foreign	Neurontin			
Initial or Prolonged		Fall	Health	(Gapabentin)	PS		ORAL
ORAL							
Professional							

Date:02/27/03ISR Number: 4068221-8Report Type:Expedited (15-DaCompany Report #2003007170
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source
Other	Parkinson'S Disease	Foreign
		Health
		Professional

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
4800 MG (400 MG, 12 TIMES A DAY), ORAL			Neurontin (Gabapentin)	PS		ORAL

Date:02/27/03ISR Number: 4068235-8Report Type:Expedited (15-DaCompany Report #2003005182
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Hepatitis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other			Professional	Clomipramine	C		

Date:02/28/03ISR Number: 4067411-8Report Type:Expedited (15-DaCompany Report #2003007195
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL	900 MG (TID),	Pruritus Rash Macular	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
Other			Professional	Amlodipine	C		
				Senna	C		
				Warfarin	C		
				Lansoprazole	C		
				Amitriptyline	C		

Date:02/28/03ISR Number: 4067853-0Report Type:Expedited (15-DaCompany Report #2003007189
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Meningioma Muscle Twitching	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other ORAL		Neuralgia					
300 MG (TID), ORAL		Paraesthesia Pruritus		Dilantin(Phenytoin Sodium)	SS		ORAL
				Levothyroxine Sodium	C		
				Warfarin Sodium	C		
				Candesartan			
				Cilexetil	C		

Date:02/28/03ISR Number: 4068033-5Report Type:Expedited (15-DaCompany Report #2002062771
Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Alcohol Interaction Aphasia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other (DAILY), ORAL		Dysphagia	Professional				
ORAL		Transient Ischaemic Attack		Ethanol	SS		ORAL
				Valaciclovir Hydrochloride	C		
				Diclofenac Sodium	C		

Freedom Of Information (FOI) Report

Date:03/03/03ISR Number: 4069102-6Report Type:Expedited (15-DaCompany Report #033-0945-980038

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abortion Spontaneous	Foreign	Neurontin			
Life-Threatening		Cardiac Disorder	Health	(Gabapentin)	PS		
TRANSPLACENTAL	1600 MG						
Congenital Anomaly		Complications Of Maternal	Professional				
(DAILY),		Exposure To Therapeutic					
PLACENTAL							
		Drugs		Tegretol			
		Congenital Anomaly		(Carbamazepine)	SS		
TRANSPLACENTAL	600 MG						
Congenital Nose							
(DAILY),		Malformation					
PLACENTAL							
		Congenital Teratoma					
		Foetal Disorder					
		Generalised Oedema					
		Intra-Uterine Death					
		Maternal Drugs Affecting					
		Foetus					
		Pregnancy					

Date:03/03/03ISR Number: 4069209-3Report Type:Expedited (15-DaCompany Report #2003006596

Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Alcohol Withdrawal	Foreign	Neurontin			
Initial or Prolonged		Syndrome	Health	(Gabapentin)	PS		ORAL
900 MG(300							
Other		Alcoholic Liver Disease	Professional				
MG, THREE		Alcoholism					
TIMES DAILY),		Gastrointestinal Disorder					
ORAL							
		Medication Error		Keltican /Old Form/	C		
		Pancreatic Disorder		Mebeverine	C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Distension	Consumer	Neurontin			
		Endoscopy Abnormal		(Gabapentin)	PS		ORAL
DAILY, ORAL							
ORAL				Methadone	SS		ORAL
				Methotrexate	C		
				Prednisone	C		
				Levothyroxine Sodium	C		
				Estropipate	C		
				Medroxyprogesterone			
				Acetate	C		
				Lansoprazole	C		
				Paroxetine			
				Hydrochloride	C		
				Multivitamins	C		
				Cisapride	C		

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Status Epilepticus	Foreign	Gabapentin			
Initial or Prolonged			Health	(Gabapentin)	PS		ORAL
UNKNOWN							
(DAILY) ,			Professional				
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/03/03ISR Number: 4069828-4Report Type:Expedited (15-DaCompany Report #2003007174
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG (DAILY), ORAL		Bladder Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Thyroid	C		
				Amitriptyline	C		
				Estrogen Nos	C		
				Cyanocobalamin	C		

Date:03/04/03ISR Number: 4069924-1Report Type:Expedited (15-DaCompany Report #PHBS2003CA01501
Age:23 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 425MG/D 300 MG/D		Hallucination Paranoia Thinking Abnormal	Foreign Literature Health	Clozapine(Clozapine) Gabapentin (Gabapentin)	PS SS		
			Professional Other	Procyclidine (Procyclidine)	C		
				Divalproex Sodium (Valproate Semisodium)	C		
				Fluoxetine (Fluoxetine)	C		

Date:03/05/03ISR Number: 4066258-6Report Type:Expedited (15-DaCompany Report #US-BRISTOL-MYERS SQUIBB COMPANY-12183901
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged INTRAVENOUS		Abdominal Pain Anaemia Asthenia	Health Professional	Taxol	PS	Bristol-Myers Squibb Company	

mg/m2 then 90
 mg/m2 27 DAY
 62 DAY

Biliary Tract Disorder
 Cholestasis
 Dyspnoea
 Frequent Bowel Movements
 Hepatitis
 Hypokalaemia
 Neutropenia
 Paraesthesia
 Peritoneal Carcinoma
 Pleural Fibrosis

Neurontin SS ORAL

Date:03/05/03ISR Number: 4066308-7Report Type:Expedited (15-DaCompany Report #CH-GLAXOSMITHKLINE-B0293213A
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	20MG Per day 57 DAY	Fall	Consumer	Deroxat	PS	Glaxosmithkline	ORAL
Initial or Prolonged	19 DAY	Tremor		Neurontin	SS		ORAL

Date:03/05/03ISR Number: 4067895-5Report Type:Direct Company Report #USP 55485
 Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Medication Error		Neurontin 600 (Gabapentin)	PS	Pfizer Uspg	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Motrin 600
(Ibuprofen 600) SS Mcneil Cons

Date:03/05/03ISR Number: 4071290-2Report Type:Expedited (15-DaCompany Report #2003007704
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL				Cosopt Paracetamol	C C		

Date:03/05/03ISR Number: 4071326-9Report Type:Expedited (15-DaCompany Report #2003008062
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (30 MG, THREE TIMES DAILY),		Clonic Convulsion Muscle Twitching	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL		Somnolence	Professional	Human Mixtard Furosemide Enalapril Acetylsalicylic Acid	C C C C		

Date:03/05/03ISR Number: 4071362-2Report Type:Expedited (15-DaCompany Report #2003005182
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Hepatic Steatosis Hepatitis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL

Other	Professional	Clomipramine	C
		Blopress Plus	C
		Pravastatin	C
		Atenolol	C
		Clopidogrel	C
		Urapidil	C

Date:03/05/03ISR Number: 4071396-8Report Type:Expedited (15-DaCompany Report #2003149054IT
 Age:80 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Disorientation Hypoxia Restlessness	Foreign Health Professional	Fragmin (Dalteparin Sodium) Solution, Sterile	PS		
SUBCUTANEOUS	2500 IU, QD, Tachypnoea	Other				
SUBCUTANEOUS			Catapres (Clonidine) Patch	SS		
TRANSDERMAL	2. 5MG, WEEKLY,					
TRANSDERMAL			Norvasc (Amlodipine Besilate) Tablet	SS		ORAL
5 MG, QD, ORAL						
2 MG, QD, ORAL			Cardura (Doxazosin Mesilate) Tablet	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150 MG, QD, ORAL	Neurontin (Gabapentin) Capsule	SS	ORAL
1 SACHET, QD, ORAL	Cardirene (Lysin Aspirin)	SS	ORAL
10 GTT, TID, ORAL	Contramal (Tramadol Hydrochloride)	SS	ORAL
INTRAVENOUS 400 MG, QD, IV	Ciproxin (Ciprofloxacin) Solution	SS	
	Morphine	C	

Date:03/06/03ISR Number: 4071758-9Report Type:Expedited (15-DaCompany Report #2003008068
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Neurontin			
Disability		Balance Disorder		(Gabapentin)	PS		
Other		Difficulty In Walking					
		Dry Mouth					
		Fall					
		Pain					
		Pain In Extremity					
		Paralysis					

Date:03/06/03ISR Number: 4072083-2Report Type:Expedited (15-DaCompany Report #2003007170
Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain	Foreign	Neurontin			
Other							

4800 MG	Headache	Health	(Gabapentin)	PS	ORAL
(DAILY), ORAL	Parkinsonism	Professional			
		Company	Paramol-118	C	
		Representative	Primidone	C	
			Ibuprofen	C	
			Atorvastatin	C	

Date:03/06/03ISR Number: 4072086-8Report Type:Expedited (15-DaCompany Report #2003007708
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG (200 MG, TID), ORAL	Blood Potassium Increased	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
			Vancomycin	C		

Date:03/06/03ISR Number: 4072088-1Report Type:Expedited (15-DaCompany Report #2003008093
Age: Gender:Female I/FU:I

Outcome Hospitalization - Initial or Prolonged Other	PT Coordination Abnormal Disturbance In Attention Drug Interaction
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Fall Parkinsonism	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
	ORAL			Professional	Paroxetine Hydrochloride	SS		ORAL
	20 MG (PER DAY), ORAL				Clomethiazole Edisilate	C		

Date:03/06/03ISR Number: 4072122-9Report Type:Expedited (15-DaCompany Report #2003008079
Age:68 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Hepatitis Cholestatic	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
	2700 MG (900 MG, TID), ORAL			Professional	Seretide "Allen & Hanburys Ltd" Citalopram Calamine/Camphor/Dip henhydramine Ampicillin Ferrous Sulfate	C C C C C		

Date:03/07/03ISR Number: 4072570-7Report Type:Expedited (15-DaCompany Report #200310616GDS
Age:80 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
			Agitation Disorientation	Foreign Health	Ciproxin (Ciprofloxacin)	PS		
	INTRAVENOUS	200 MG, BID,						

			Hypoxia	Professional		
INTRAVENOUS			Tachypnoea	Other	Fragmin (Heparin-Fraction, Sodium Salt)	SS
SUBCUTANEOUS	2500 IU,					
TOTAL DAILY, SUBCUTANEOUS						
TRANSDERMAL	2.5 MG, TOTAL				Catapresan (Clonidine)	SS
DAILY, TRANSDERMAL (TRANSCUTANEO US)	7	YR				
5 MG, TOTAL DAILY, ORAL	5	YR			Norvasc (Amlodipine Besilate)	SS ORAL
2 MG, TOTAL DAILY, ORAL	4	YR			Cardura (Doxazosin Mesilate)	SS ORAL
300 MG, TOTAL DAILY, ORAL	7	MON			Neurontin (Gabapentin)	SS ORAL
ORAL	4	YR			Cardirene (Acetylsalicylate Lysine)	SS ORAL
ORAL					Contramal (Tramadol Hydrochloride)	SS ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/10/03ISR Number: 4073663-0Report Type:Expedited (15-DaCompany Report #2003008240

Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (TID), ORAL		Csf Pressure Increased Headache	Health Professional	Neurontin(Gabapentin)	PS		ORAL
Other		Multiple Sclerosis Muscle Rigidity Nuclear Magnetic Resonance Imaging Abnormal					

Date:03/10/03ISR Number: 4073699-XReport Type:Expedited (15-DaCompany Report #2003008392

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 900 MG (DAILY), ORAL		Nausea Vitreous Detachment	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Amitriptyline Tramadol Paramol-118 Lansoprazole	C C C C		

Date:03/10/03ISR Number: 4073730-1Report Type:Expedited (15-DaCompany Report #2003007508

Age:80 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 5 MG (DAILY), Initial or Prolonged ORAL		Agitation	Foreign	Norvasc (Amlodipine)	PS		ORAL
		Disorientation	Health				
		Hypoxia Pain	Professional	Cardura (Doxazosin Mesilate)	SS		ORAL
2 MG (DAILY), ORAL		Tachypnoea					

1.5 MG, ORAL		Neurontin (Gabapentin)	SS	ORAL
TRANSDERMAL	5 (DAILY),	Clonidine Hydrochloride	SS	
TRANSDERMAL				
SUBCUTANEOUS	2500 UNITS	Heparin-Fraction, Sodium Salt	SS	
(DAILY),				
SUBCUTANEOUS				
		Ciprofloxacin	C	
		Acetylsalicylate		
		Lysine	C	
		Morphine		
		Hydrochloride	C	
		Tramadol		
		Hydrochloride	C	

Date:03/10/03ISR Number: 4074007-0Report Type:Expedited (15-DaCompany Report #2003008536
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Blood Pressure Decreased
Other	Bronchitis
	Convulsion
	Dehydration

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Level Fluctuating Drug Toxicity Rhinorrhoea	Report Source	Product	Role	Manufacturer	Route
ORAL		Somnolence Urinary Tract Infection	Consumer	Dilantin Infatabs (Phenytoin Sodium)	PS		ORAL
300 MG (100 MG, THREE TIMES DAILY), ORAL		Vaginal Infection		Neurontin (Gabapentin)	SS		ORAL
INTRADERMAL	INTRADERMAL			Glyceryl Trinitrate	SS		
				Acetylsalicylic Acid	C		
				Estrogens Conjugated	C		
				Hydrochlorothiazide	C		
				Potassium	C		
				Donepezil			
				Hydrochloride	C		

Date:03/10/03ISR Number: 4074022-7Report Type:Expedited (15-DaCompany Report #1667783A
Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Condition Aggravated Leukopenia	Foreign Health Professional	Imodium (Loperamide Hydrochloride) Capsules	PS		ORAL
PO	15 DAY			Doliprane (Paracetamol)	SS		ORAL
PO	8 DAY			Neurontin	SS		
				Zelcar	SS		
				Myabumbutol Tb	SS		
				Rifater Tb	SS		

Date:03/10/03ISR Number: 4074332-3Report Type:Expedited (15-DaCompany Report #2003006796
Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Other		Drug Withdrawal Syndrome	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Professional Company Representative	Diclofenac Rabeprazole Panadeine Co Ergotamine Tartrate	C C C C		

Date:03/10/03ISR Number: 4074339-6Report Type:Expedited (15-DaCompany Report #2003008542
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthralgia Oedema Oliguria	Foreign Health Professional	Neurontin (Gabapentin) All Other Therapeutic Agents	PS C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/10/03ISR Number: 4074342-6Report Type:Expedited (15-DaCompany Report #2003005551
Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	900 MG (TID), ORAL	Visual Acuity Reduced	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
				Tramadol	C		
				Paracetamol	C		
				Mianserin	C		
				Mepronizine	C		

Date:03/11/03ISR Number: 4073203-6Report Type:Expedited (15-DaCompany Report #2003008839
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	600 MG (BID), ORAL	Insomnia Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Pharmaceutical Product					
		Complaint		Alendronate Sodium	C		
		Treatment Noncompliance		Ascorbic Acid	C		
				Tocopherol	C		
				Calcium With Vitamin D	C		

Date:03/11/03ISR Number: 4074268-8Report Type:Expedited (15-DaCompany Report #2003-IT-00018IT(1)
Age:80 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	TRANSDERMAL 0.1 MG (2.5	Agitation Disorientation Hypoxia Tachypnoea	Other	Catapresan Tts (Clonidine Hydrochloride) (Tts) (Clonidine-Hcl)	PS		

MG, 1 IN 1

WK)	7	YR	Norvasc (Amlodipine Besilate) (Ta)	SS	ORAL
5 MG (5 MG, 1 TA DAY)	5	YR	Cardura (Doxazosin Mesilate) (Ta)	SS	ORAL
2 MG (2 MG, 1 TA DAY)	4	YR	Neurontin (Gabapentin) (Ka)	SS	ORAL
300 MG	7	YR	Cardirene (Acetylsalicylate Lysine) (Grb)	SS	ORAL
1 GRB DAY			Contramal (Tramadol Hydrochloride) (Tr)	SS	ORAL
75 MG (2.5 MG, 10 GTT TID)			Ciproxin (Ciprofloxacin) (Loi)	SS	
INTRAVENOUS	400 MG (400 MG)		Fragmin (Heparin Fraction, Sodium Salt)	SS	
SUBCUTANEOUS	2500 U				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/11/03ISR Number: 4074424-9Report Type:Expedited (15-DaCompany Report #2003008819
Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 Other MG(DAILY), ORAL	Alopecia Anaemia Vitamin B6 Deficiency Blood Iron Decreased Catheter Related Infection Insomnia Pneumonia Therapeutic Response Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Hydrocodone	C		

Date:03/11/03ISR Number: 4074670-4Report Type:Expedited (15-DaCompany Report #2003008333
Age:11 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1500 MG Other (DAILY), ORAL	Aggression Disturbance In Attention	Foreign Health Professional	Gabapentin(Gabapenti n)	PS		ORAL
			Carbamazepine	C		
			Oxcarbazepine	C		
			Primidone	C		

Date:03/12/03ISR Number: 4075137-XReport Type:Expedited (15-DaCompany Report #2003008266
Age:30 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged (DAILY), ORAL	Condition Aggravated Head Injury Status Epilepticus	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
			Valproate Sodium	C		
			Topiramate	C		

Chlorpromazine
Hydrochloride C

Date:03/12/03ISR Number: 4075356-2Report Type:Expedited (15-DaCompany Report #2003006318
Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Renal Failure Acute Shock	Foreign Health	Neurontin(Gabapentin)	PS		ORAL
600 MG		Staphylococcal Sepsis	Professional				
(DAILY), ORAL		Toxic Epidermal Necrolysis		Methylprednisolone	SS		
INTRAVENOUS	20 MG						
(DAILY),							
INTRAVENOUS							
INTRAMUSCULAR	10 MG			Methotrexate	SS		
(WEEKLY),							
INTRAMUSCULAR							

Date:03/12/03ISR Number: 4075509-3Report Type:Expedited (15-DaCompany Report #2003008843
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Glucose Increased Convulsion	Health Professional	Neurontin (Gabapentin)	PS		
700 MG		Rhabdomyolysis					
Other							
(DAILY)							

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Freedom Of Information (FOI) Report

Acetylsalicyclic Acid	C
Warfarin Sodium	C
Iron	C
Folic Acid	C
Docusate Sodium	C
Ergocalciferol	C

Date:03/13/03ISR Number: 4075604-9Report Type:Expedited (15-DaCompany Report #2003003300
 Age:47 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Abdominal Pain Anaemia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other INTRAVENOUS (EVERY THREE WEEKS) , INTRAVENOUS	Ascites Asthenia Biliary Tract Disorder Blood Creatinine Increased Cholestasis Diarrhoea Gastrointestinal Disorder Hepatic Trauma Hypokalaemia Nausea Neutropenia Paraesthesia Peritoneal Carcinoma Pleural Effusion Pleural Fibrosis Vomiting White Blood Cell Count Decreased	Professional	Paclitaxel	SS		

Date:03/13/03ISR Number: 4075626-8Report Type:Expedited (15-DaCompany Report #2003009270
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Drug Interaction	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL		Respiratory Depression	Professional	Morphine	SS		

Date:03/13/03ISR Number: 4075805-XReport Type:Expedited (15-DaCompany Report #FRWYE050807MAR03
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2.5 MG 2X PER	Hyponatraemia	Health Professional	Temesta (Lorazepam, Tablet, 0)	PS		ORAL
1 DAY			Other				
1 DOSE 1X PER				Atenolol (Atenolol, , 0)	SS		ORAL
1 DAY							
30 MG 1X PER				Athymil (Mianserin Hydrochloride, , 0)	SS		ORAL
1 DAY							
5 DAY				Depamide (Valpromide, , 0)	SS		ORAL
				Neurontin			

Freedom Of Information (FOI) Report

(Gabapentin, , 0) SS

ORAL

300 MG 2X PER

1 DAY

Date:03/13/03ISR Number: 4076082-6Report Type:Expedited (15-DaCompany Report #2003001198

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased	Consumer	Neurontin			
		Cardiac Operation		(Gabapentin)	PS		
		Dementia		Metoprolol Tartrate	SS		
		Diabetes Mellitus		Digoxin	SS		
				Isosorbide Dinitrate	SS		
				Furosemide	SS		
				Captopril	SS		
				Warfarin Sodium	C		
				Lorazepam	C		
				Quetiapine Fumarate	C		

Date:03/13/03ISR Number: 4076156-XReport Type:Expedited (15-DaCompany Report #PHNU2003DE00883

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dialysis	Foreign	Tegretal(Carbamazepi			
Initial or Prolonged		Drug Interaction	Health	ne) Unknown	PS		
SEE IMAGE							
Other		Hypothermia	Professional	Neurontin(Gabapentin			
		Loss Of Consciousness	Other)	SS		ORAL
SEE IMAGE							
		Respiratory Failure		Durogesic(Fentanyl)			
				Patch	SS		
				Novalgine(Metamizole			
				Sodium) Drop	SS		
				Cipramil(Citalopram			
				Hydrobromide)	SS		

Date:03/13/03ISR Number: 4076277-1Report Type:Expedited (15-DaCompany Report #2003009224

Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	230 MG	Arthralgia Clumsiness	Health Professional	Dilantin Kapseals (Phenytoin Sodium)	PS		ORAL
Other (DAILY), ORAL		Electrocardiogram					
	1500 MG (300 MG, TWICE DAILY), ORAL	Abnormal Fatigue Oral Intake Reduced Post Procedural Complication Vision Blurred Weight Decreased		Neurontin (Gabapentin)	SS		ORAL
				Estrogens Conjugated Vitamins Acetylsalicylic Acid Zolpidem Tartrate	C C C C		

Date:03/17/03ISR Number: 4077346-2Report Type:Expedited (15-DaCompany Report #2003007189
Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Burning Sensation Drug Interaction
Other	Facial Pain Meningioma Muscle Twitching

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Paraesthesia Pruritus	Report Source	Product	Role	Manufacturer	Route
300 MG (TID), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (TID), ORAL				Dilantin (Phenytoin Sodium)	SS		ORAL
ORAL				Warfarin Sodium	SS		ORAL
				Levothyroxine Sodium	C		
				Candesartan			
				Cilexetil	C		

Date:03/17/03
Age:20 YR
Gender:Male
ISR Number: 4077372-3
Report Type:Expedited (15-DaCompany Report #2003009221
I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (DAILY), ORAL		Convulsion Essential Tremor Eye Disorder Memory Impairment Visual Disturbance	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Didanosine	C		
				Stavudine	C		
				Saquinavir	C		
				Kaletra	C		
				Bactrim	C		
				Azithromycin	C		
				Fluconazole	C		
				Levothyroxine Sodium	C		
				Paroxetine Hydrochloride	C		
				Diazepam	C		
				Oxycodone	C		
				Dronabinol	C		
				Nystatin	C		
				Benzonatate	C		

Phenytoin Sodium C
 Cetirizine
 Hydrochloride C
 Celecoxib C

Date:03/17/03ISR Number: 4077393-0Report Type:Expedited (15-DaCompany Report #2002068411

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 5600 MG (800 Other MG, 7 TIMES A DAY)		Unevaluable Event	Consumer Health Professional	Neurontin (Gabapentin)	PS		
200 MG (100 MG,BID)				Zoloft (Sertraline)	SS		
200 MG (100 MG, BID)				Trazodone Hydrochloride	SS		
1 MG (0.25 MG, QID)				Alprazolam	SS		
15/500 MG (DOSE STRENGTH				Vicodin	SS		

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Freedom Of Information (FOI) Report

5/500 MG)
 (TID)
 1000 MG (200
 MG, FIVE
 TIMES A DAY)
 162 MG (81
 MG, BID)

Ibuprofen SS

Acetylsalicylic Acid SS

Date:03/17/03ISR Number: 4077395-4Report Type:Expedited (15-DaCompany Report #2003007189
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG (TID),	Burning Sensation Facial Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other ORAL		Meningioma					
	300 MG (TID),	Muscle Twitching Paraesthesia		Dilantin (Phenytoin Sodium)	SS		ORAL
ORAL		Pruritus					
				Levothyroxine Sodium	C		
				Warfarin Sodium	C		
				Candesartan			
				Cilexetil	C		

Date:03/17/03ISR Number: 4077427-3Report Type:Expedited (15-DaCompany Report #2003009356
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	800 MG (400	Breast Pain Constipation	Consumer	Neurontin (Gabapentin)	PS		ORAL

MG BID) ORAL

Diarrhoea

Fibroadenoma Of Breast

Flatulence

Insomnia

Pharyngolaryngeal Pain

Stress

Suicidal Ideation

Weight Increased

Fluoxetine Hydrochloride C

Date:03/17/03ISR Number: 4077531-XReport Type:Expedited (15-DaCompany Report #2003005569
 Age:87 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (DAILY), ORAL		Confusional State Fall	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:03/17/03ISR Number: 4077604-1Report Type:Expedited (15-DaCompany Report #EMADSS2003002034
 Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged TRANSDERMAL	TRANSD	International Normalised Ratio Decreased	Foreign Health Professional	Durogesic (Patch) (Fentanyl)	PS		
200 MG, 3 IN				Neurontin (Gabapentin)	SS		ORAL

1 DAY(S),

ORAL

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Freedom Of Information (FOI) Report

10 MG DAILY,
ORAL

Previscan
(Fluindione) SS ORAL

Date:03/17/03ISR Number: 4077693-4Report Type:Expedited (15-DaCompany Report #2003001247
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Hepatitis Cholestatic	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other			Professional Company Representative	Aciclovir Metamizole Sodium	C C		

900 MG, ORAL

Date:03/17/03ISR Number: 4077746-0Report Type:Expedited (15-DaCompany Report #2003005168
Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Delirium Hallucination	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other			Professional Company Representative	Tramadol	SS		ORAL
				All Other Therapeutic Products	C		

600 MG (TID),
ORAL
300 MG (BID),
ORAL

Date:03/17/03ISR Number: 4077748-4Report Type:Expedited (15-DaCompany Report #2003006044
Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Coma	Foreign	Neurontin			

900 MG (TID),	Hypothermia	Health	(Gabapentin)	PS	ORAL
ORAL	Loss Of Consciousness	Professional			
200 MG, ORAL	Respiratory Failure		Carbamazepine	SS	ORAL
			Hypnotics And Sedatives	C	
			Zopiclone	C	
			Citalopram		
			Hydrobromide	C	
			Antihypertensives	C	

Date:03/17/03ISR Number: 4077770-8Report Type:Expedited (15-DaCompany Report #2003009487
Age: Gender:Female I/FU:I

Outcome	PT
Other	Alopecia
	Arrhythmia
	Blood Pressure Increased
	Cardiac Failure
	Dizziness
	Drug Hypersensitivity
	Heart Rate Increased
	Hypotension
	Memory Impairment
	Myocardial Infarction
	Pain
	Pulmonary Hypertension

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Restlessness Somnolence Thinking Abnormal					
50 MG, ORAL		Tricuspid Valve Disease Weight Decreased Weight Increased	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
				Nitrostat (Glyceryl Trinitrate)	SS		
				Lidocaine	SS		
				Baclofen	SS		
				K-Lyte	C		
				Irbesartan	C		
				Spiroinolactone	C		
				Oxygen	C		
				Taurine	C		
				All Other Therapeutic Products	C		
				Atenolol	C		

Date:03/17/03ISR Number: 4077771-XReport Type:Expedited (15-DaCompany Report #2003009903
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG, TID), ORAL		Jaundice Cholestatic	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
				Insulin Porcine	C		
				Ramipril	C		
				Metformin	C		
				Dihydrocodeine	C		

Date:03/17/03ISR Number: 4077774-5Report Type:Expedited (15-DaCompany Report #2002072263
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Anorexia	Foreign	Neurontin			

Initial or Prolonged ORAL	Inflammation	Health	(Gabapentin)	PS	ORAL
	Malaise	Professional	Simvastatin	SS	ORAL
ORAL	Myalgia		Acetylsalicylic Acid	C	
	Red Blood Cell Sedimentation Rate Increased		Anastrozole	C	
	Thyroid Neoplasm				

Date:03/17/03ISR Number: 4077777-0Report Type:Expedited (15-DaCompany Report #2003009896

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 500 MG	Cardiac Failure Endocarditis Bacterial	Foreign Health	Neurontin (Gabapentin)	PS		
Initial or Prolonged (DAILY), Other	Hyponatraemia	Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/18/03ISR Number: 4073578-8Report Type:Expedited (15-DaCompany Report #WAES 0209USA02503

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Enzymes Increased		Vioxx	PS	Merck & Co., Inc	ORAL
Hospitalization -		Completed Suicide		Gabapentin	SS		
Initial or Prolonged		Coordination Abnormal		Ultram	SS		ORAL
Other		Cyanosis					
		Dehydration					
		Electromechanical					
		Dissociation					
		Heart Rate Decreased					
		Hypotension					
		Mental Status Changes					
		Myoglobin Urine Present					
		Renal Failure					
		Somnolence					
		White Blood Cell Count					
		Increased					

Date:03/18/03ISR Number: 4078376-7Report Type:Expedited (15-DaCompany Report #EMADSS2003002034

Age:76 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		International Normalised	Foreign	Durogesic			
Initial or Prolonged		Ratio Decreased	Health	(Fentanyl)	PS		
TRANSDERMAL	TRANSD	International Normalised	Professional	Neurontin			
		Ratio Increased		(Gabapentin)	SS		ORAL
200 MG, 3 IN							
1 DAY(S),							
ORAL							
				Previscan			
				(Fluindione)	SS		ORAL
10 MG, DAILY,							
ORAL							

Date:03/18/03ISR Number: 4078899-0Report Type:Expedited (15-DaCompany Report #2002-04879
Age:43 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300 MG QD PO	Hepatitis Infectious	Foreign	Viread	PS		ORAL
Initial or Prolonged 3 U BID PO	Leukopenia	Health	Kaletra	SS		ORAL
1 U BID PO	Neutropenia	Professional	Viramune	SS		ORAL
1800 MG QD PO	Tuberculosis	Other	Cymevan Neurontin	SS SS		ORAL
			Valcyte	SS		
			Bactrim	C		
			Osfolate	C		
			Paracetamol	C		
			Imodium	C		

Date:03/19/03ISR Number: 4076549-0Report Type:Direct Company Report #CTU 189141
Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Muscle Rigidity Neuropathy Peripheral		Gabapentin Diazepam	PS C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/19/03ISR Number: 4077417-0Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #USP 55632

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error Somnolence		Neurontin 100mg Gabapentin Parke Davis Capsule 100mg Neurontin Gabapentin Parke Davis Capsule 300mg	PS SS	Parke Davis Parke Davis	

Date:03/19/03ISR Number: 4078565-1Report Type:Expedited (15-DaCompany Report #2003009796
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Breast Cancer Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
400 MG (BID), ORAL		Diabetes Mellitus					
2.5 MG (DAILY), ORAL		Mitral Valve Disease		Glucotrol (Glipizide)	SS		ORAL
				Digoxin	C		
				Ramipril	C		
				Pramipexole	C		
				Entacapone	C		
				Sinemet	C		
				Levothyroxine Sodium	C		

Date:03/19/03ISR Number: 4078566-3Report Type:Expedited (15-DaCompany Report #2003009893
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged INTRAVENOUS	(ONCE),	Blister Erythema	Consumer	Benadryl Injection (Diphenhydramine)	PS		

Other	Face Oedema				
INTRAVENOUS					
	Pain In Extremity	Neurontin			
	Pruritus	(Gabapentin)	SS		ORAL
500 MG (TID),					
ORAL	Rash				
	Tongue Oedema	Oxycodone			
	Urticaria	Hydrochloride	C		
		Ibuprofen	C		

Date:03/19/03ISR Number: 4078760-1Report Type:Expedited (15-DaCompany Report #2003010332

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Serum Sickness	Health Professional	Neurontin (Gabapentin)	PS		

Date:03/19/03ISR Number: 4078769-8Report Type:Expedited (15-DaCompany Report #2003010329

Age:62 YR Gender:Female I/FU:I

Outcome	PT
Other	Blood Pressure Increased
	Condition Aggravated
	Drug Effect Decreased
	Eye Haemorrhage
	Feeling Abnormal

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Freedom Of Information (FOI) Report

Dose	Duration	Pain In Extremity Transient Ischaemic Attack	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Clonidine	C		
				Methadone	C		
				Diphenhydramine Hydrochloride	C		
				Panadeine Co	C		

Date:03/19/03ISR Number: 4078770-4Report Type:Expedited (15-DaCompany Report #2003010331
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Nail Disorder	Health Professional	Neurontin (Gabapentin)	PS		

Date:03/19/03ISR Number: 4078771-6Report Type:Expedited (15-DaCompany Report #2003009975
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Abnormal Sensation In Eye Constipation	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Fatigue Impaired Driving Ability Increased Appetite Intervertebral Disc Protrusion Oliguria Weight Increased		Valdecoxib	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged INTRAVENOUS		Body Temperature Decreased	Health Professional	Paclitaxel	PS	Bristol-Myers Squibb Company	
		Haematocrit Decreased Haemoglobin Decreased Pancreatitis White Blood Cell Count Decreased		Gabapentin	SS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG DAILY		Inappropriate Antidiuretic Hormone Secretion	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Furosemide	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/24/03ISR Number: 4082223-7Report Type:Expedited (15-DaCompany Report #2003010602

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Convulsion Medication Error	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:03/24/03ISR Number: 4082233-XReport Type:Expedited (15-DaCompany Report #2003007169

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (300 MG, TID), ORAL UNKNOWN (UNKNOWN), UNKNOWN UNKNOWN (UNKNOWN), UNKNOWN		Double Vessel Bypass Graft Dry Mouth Dry Throat Glossodynia Neuropathy Peripheral	Consumer Health Professional	Neurontin (Gabapentin) All Other Non-Therapeutic Products All Other Non-Therapeutic Products All Other Therapeutic Products Warfarin Sodium	PS SS SS C C		ORAL

Date:03/24/03ISR Number: 4082265-1Report Type:Expedited (15-DaCompany Report #2003010615

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bone Marrow Depression	Foreign Health	Neurontin (Gabapentin)	PS		
400 MG			Professional				
(DAILY)			Company Representative				

Date:03/24/03ISR Number: 4082296-1Report Type:Expedited (15-DaCompany Report #2003010319

Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Drug Interaction International Normalised	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL			Professional				
Other		Ratio		Fentanyl	SS		
INTRADERMAL	INTRADERMAL			Fluindione	SS		ORAL
(DAILY), ORAL							

Date:03/24/03ISR Number: 4082306-1Report Type:Expedited (15-DaCompany Report #2003010534

Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Blood Sodium Decreased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Professional	Valpromide	SS		ORAL
ORAL				Atenolol	SS		ORAL
ORAL							

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL				Mianserin Hydrochloride	SS		ORAL
ORAL				Lorazepam	SS		ORAL
Date:03/24/03ISR Number: 4082341-3Report Type:Expedited (15-DaCompany Report #2003008542 Age:62 YR Gender:Female I/FU:F							
Hospitalization - Initial or Prolonged	900 MG (300 MG, THREE TIMES DAILY),	Arthralgia Oedema Oedema Peripheral Oliguria	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL	200 MG (DAILY), ORAL			Fenofibrate	SS		ORAL
ORAL	50 MG (DAILY), ORAL			Anethole Trithione	SS		ORAL
ORAL	25 MG (DAILY), ORAL			Clomipramine Hydrochloride	SS		ORAL
ORAL	1 MG (DAILY),			Clonazepam	SS		ORAL
ORAL				Endotelon	SS		ORAL
ORAL				Cefpodoxime Proxetil	C		
				Carbocisteine	C		
				Furosemide	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Agitation		Gabapentin	PS		
Hospitalization - Initial or Prolonged		Lethargy Somnolence					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Renal Failure Schizophrenia	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Condition Aggravated Dermatosis	Foreign Study	Neurontin (Gabapentin)	PS		ORAL
600 MG (DAILY), ORAL		Toxic Skin Eruption	Health Professional	Triatec (Ramipril) Bromazepam Clonazepam Perindopril Esberiven Forte Metformin Embonate	C C C C C C		

Freedom Of Information (FOI) Report

Metformin
Hydrochloride C

Date:03/26/03ISR Number: 4083816-3Report Type:Expedited (15-DaCompany Report #2003009224
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 230 MG Other (DAILY) ORAL		Arthralgia Biliary Tract Disorder Clumsiness	Health Professional	Dilantin Kapseals (Phenytoin Sodium)	PS		ORAL
1500 MG (300 MG TWICE DAILY) ORAL		Electrocardiogram Abnormal Faeces Pale Fatigue		Neurontin (Gabapentin)	SS		ORAL
		Gastritis Oral Intake Reduced Post Procedural Complication Vision Blurred Weight Decreased		Estrogens Conjugated Vitamins Acetylsalicylic Acid Zolpidem Tartrate	C C C C		

Date:03/26/03ISR Number: 4083861-8Report Type:Expedited (15-DaCompany Report #2003007174
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100 MG (DAILY), ORAL		Bladder Pain Bladder Spasm Irritable Bowel Syndrome	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Pelvic Pain Pollakiuria Pruritus Urinary Retention		Zolpidem Tartrate Amitriptyline Hydrochloride Amitriptyline Thyroid Estradiol	C C C C C		

Date:03/26/03ISR Number: 4084299-XReport Type:Expedited (15-DaCompany Report #EMADSS2003002034
Age:76 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged TRANSDERMAL	TRANS	International Normalised Ratio Decreased	Foreign Health Professional	Durogesic (Patch) (Fentanyl)	PS		
200 MG, 3 IN				Neurontin (Gabapentin)	SS		ORAL
1 DAY, ORAL				Previscan (Fluindione)	SS		ORAL
10 MG, DAILY, ORAL							

Date:03/27/03ISR Number: 4084807-9Report Type:Expedited (15-DaCompany Report #2003004307
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Coma	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:03/27/03ISR Number: 4084809-2Report Type:Expedited (15-DaCompany Report #2003011961
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Bronchiolitis Dyspnoea	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1800 MG (900 MG, BID), ORAL				Bupropion Hydrochloride Seretide Mite Fluticasone Propionate Salbutamol	C C C C		

Date:03/27/03ISR Number: 4085869-5Report Type:Expedited (15-DaCompany Report #2002072567
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Fatigue Malaise Somnolence	Health Professional	Neurontin (Gabapentin) All Other Therapeutic Products	PS C		

Date:03/27/03ISR Number: 4086575-3Report Type:Expedited (15-DaCompany Report #PHFR2003GB00605
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Interaction Metastasis Salivary Gland Cancer	Foreign Health Professional Other	Tegretal (Carbamazepine) Diamorphine (Diamorphine) Gabapentin (Gabapentin)	PS SS SS		

Date:03/27/03ISR Number: 4086788-0Report Type:Expedited (15-DaCompany Report #2003011833
Age:80 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Fall Femur Fracture	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:03/28/03ISR Number: 4083667-XReport Type:Direct Company Report #CTU 189682
Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration MUSCLE SPASUM	Asthenia Circulatory Collapse Convulsion Feeling Abnormal Vision Blurred		Neurontin	PS		

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Freedom Of Information (FOI) Report

Date:03/28/03ISR Number: 4086758-2Report Type:Expedited (15-DaCompany Report #2003009224

Age:63 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 230 MG Other (DAILY), ORAL	Arthralgia Biliary Tract Disorder	Health Professional	Dilantin Kapseals (Phenytoin Sodium)	PS		ORAL
1500 MG (TWICE DAILY), ORAL	Clumsiness Dyskinesia Electrocardiogram Abnormal Faeces Discoloured		Neurontin (Gabapentin)	SS		ORAL
	Fatigue Gastritis Hypersensitivity Insomnia Muscle Injury Nerve Injury Oral Intake Reduced Post Procedural Complication Vision Blurred Weight Decreased		Estrogens Conjugated Vitamins Acetylsalicylic Acid Zolpidem Tartrate	C C C C		

Date:03/28/03ISR Number: 4087010-1Report Type:Expedited (15-DaCompany Report #2003010424

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG (600 MG, TID), ORAL	Inappropriate Antidiuretic Hormone Secretion	Foreign Health Professional	Neurontin(Gabapentin)	PS		ORAL
			Furosemide Amiodarone Hydrochloride Warfarin	C C C C		

Digoxin C
Tramadol C
Valsartan C

Date:03/28/03ISR Number: 4087018-6Report Type:Expedited (15-DaCompany Report #2003009975
Age:28 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Constipation	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Dizziness					
		Dysarthria		Valdecoxib	SS		
		Eye Disorder					
		Fatigue					
		Feeling Abnormal					
		Increased Appetite					
		Intervertebral Disc Protrusion					
		Oliguria					
		Sciatica					
		Weight Increased					

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Freedom Of Information (FOI) Report

Date:03/28/03ISR Number: 4087020-4Report Type:Expedited (15-DaCompany Report #2003008240

Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Csf Pressure Increased Headache	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other		Musculoskeletal Stiffness Nuclear Magnetic Resonance Imaging Abnormal					

Date:03/28/03ISR Number: 4087060-5Report Type:Expedited (15-DaCompany Report #2003011978

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abnormal Behaviour Anxiety	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Drug Dependence Irritability Thinking Abnormal		Bupropion Hydrochloride Doxycycline	C C		

Date:03/31/03ISR Number: 4087535-9Report Type:Expedited (15-DaCompany Report #2001-07-0134

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Anaemia Arthralgia	Health Professional Company	Thalomid (Thalidomide 50 Mg) Capsules	PS		ORAL
400 MG QD		Arthritis	Representative				
ORAL		Asthenia		Aredia	SS		
INTRAVENOUS	90 MG	Blood Creatinine					
INTRAVENOUS		Increased					
NOS							

300 MG BID

Constipation	Neurontin	SS
Dehydration	Arava	SS
Diarrhoea	Bicnu	C
Dyspnoea Exertional	Dexamethasone	C
Ecchymosis	Cytoxan	C
Flushing	Biaxin	C
Generalised Oedema		
Hypoproteinaemia		
Leukopenia		
Mucosal Inflammation		
Nephrotic Syndrome		
Orthostatic Hypotension		
Pain In Extremity		
Pitting Oedema		
Red Blood Cell		
Sedimentation Rate		
Increased		
Renal Failure Acute		
Syncope		
Thrombocytopenia		
Tremor		

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Date:04/01/03 ISR Number: 4085587-3 Report Type:Expedited (15-DaCompany Report #2003012199

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract	Consumer	Neurontin			
300 MG		Condition Aggravated		(Gabepentin)	PS		ORAL
(DAILY), ORAL		Discomfort					
		Pain		Pentosan Polysulfate			
		Pharmaceutical Product		Sodium (Pentosan	C		
		Complaint		Polysulfate Sodium)			
		Pollakiuria		Amitriptyline			
				Hydrochloride			
				(Amitriptyline			
				Hydrochloride)	C		
				Clonazepam			
				(Clonazepam)	C		
				Rofecoxib			
				(Rofecoxib)	C		
				Simvastatin			
				(Simvastatin)	C		
				Estradiol			
				(Estradiol)	C		
				Oxycocet			
				(Paracetamol,			
				Oxycodone			
				Hydrochloride)	C		

Date:04/01/03 ISR Number: 4085671-4 Report Type:Expedited (15-DaCompany Report #2003005197

Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain	Consumer	Neurontin			
900 MG (300,		Cholelithiasis		(Gabapentin)	PS		ORAL
TID), ORAL		Diarrhoea					
ORAL		Difficulty In Walking		Norvasc (Amlodipine)	SS		ORAL
		Drug Ineffective		Meloxicam			
		Joint Stiffness		(Meloxicam)	C		
		Joint Swelling		Atenolol (Atenolol)	C		

Somnolence

Vitamins

C

Date:04/02/03ISR Number: 4087855-8Report Type:Expedited (15-DaCompany Report #2002068428

Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dehydration	Foreign	Neurontin			
Other		Dyspnoea	Health	(Gabapentin)	PS		ORAL
900 MG (THREE		Haematochezia	Professional				
TIMES DAILY),		Oedema					
ORAL		Shock		All Other			
				Therapeutic Products	SS		ORAL
				Valaciclovir			
				Hydrochloride			
				(Valaciclovir			
				Hydrochloride)	C		
				Ciprofloxacin			
				(Ciprofloxacin)	C		
				All Other			
				Therapeutic Products	C		
				Ketosteril (Amino			
				Acids Nos)	C		

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All Other
 Therapeutic Products C
 Furosemide
 (Furosemide) C
 Cefoperazone Sodium
 (Cefoperazone
 Sodium) C
 All Other
 Therapeutic Products C

Date:04/02/03ISR Number: 4088112-6Report Type:Expedited (15-DaCompany Report #2003012207
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Condition Aggravated Drug Tolerance	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL		Neuralgia Therapeutic Response Decreased		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride) Hydroxychloroquine Phosphate Glucosamine (Glucosamine) Soy Isoflavones (Soy Isoflavones) Ginseng (Ginseng)	C C C C		

Date:04/02/03ISR Number: 4088344-7Report Type:Expedited (15-DaCompany Report #USA-2002-0001677
 Age:41 YR Gender:Female I/FU:I

Outcome	PT
Death	Alopecia
Hospitalization - Initial or Prolonged	Anxiety Arterial Occlusive Disease Arthralgia Asthma Back Pain

Carpal Tunnel Syndrome
Coma
Depression
Disturbance In Attention
Drug Ineffective
Emphysema
Fatigue
Feeling Abnormal
Flushing
Hilar Lymphadenopathy
Influenza
Insomnia
Irritability
Libido Decreased
Memory Impairment
Nausea
Nervousness
Night Sweats
Overdose

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Dose	Duration	Adverse Event	Report Source	Product	Role	Manufacturer	Route
		Pain Pain In Extremity Pneumonia					
10 MG		Pulmonary Oedema Rash Generalised Rash Pruritic Rash Scaly	Consumer Health Professional Other	Oxycontin Tablets 10 Mg(Oxycodone Hydrochloride)Cr Tablet			PS
20 MG	1008 DAY	Skin Nodule Somnolence Upper Respiratory Tract Infection		Oxycontin Tablets 20 Mg(Oxycodone Hydrochloride)Cr Tablet			SS
		Urinary Incontinence Weight Increased		Hydrocodone Bitartrate(Similar To Ind 59,175)(Hydrocodone Bitartrate)Unknown			SS
1008 DAY				Alprazolam (Alprazolam)			SS
1008 DAY				Ephedrine(Epedrine)			SS
1008 DAY				Pseudoephedrine (Pseudoephedrine)			SS
75 MG, DAILY	1008 DAY			Effexor(Venlafaxine Hydrochloride)			SS
1008 DAY				Codeine(Codeine)			SS
100 MG, TID	1008 DAY			Neutrontin(Gabapenti n)			SS
1008 DAY				Metoclopramide (Metoclopramide)			SS
1008 DAY				Quetiapine(Quetiapin e)			SS
				Claritin (Loratadine)			C
				Vitamin C (Ascorbic Acid)			C
				Celebrex(Celecoxib)			C

Prilosec(Omeprazole)	C
Daypro(Oxaproxin)	C
Trandate(Labetalol Hydrochloride)	C
Axid(Nizatidine)	C
Paxil(Paroxetine Hydrochloride)	C
Medrol (Methylprednisone)	C
Macro Antioxidant(Ascorbic Acid, Cystine, Tocopherol, Calcium Ascorbate,	C
Zocor (Simvastatin)	C
Risperdal (Risperidone)	C
Fioricet(Butalbital)	C
Flonase (Fluticasone Propionate)	C
Phenobarbital (Phenobarbital)	C
Donnatal (Atropine Sulfate,Hyoscine Hydrobromide, Hyoscyamine Sulfate,	

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Phenobarbital)	C
Bellergal-S	
(Belladonna	
Alkaloids,	
Erogotamine	
Tartrate,	C
Weight Loss	
Supplment	C
Ibuprofen	C
Ambien (Zolpidem	
Tartrate)	C
Baclofen (Baclfoen)	C
(Labetalol	
Hydrochloride)	C
Celexa (Cialopram	
Hydrobromide)	C
Cortisone	
(Cortisone)	C
Atarax (Hydroxyzine	
Hydrochloride)	C
Proventil Tablet	
(Salbutamol Sulfate)	C
Amitrptyline	
(Amitriptyline)	C
Cyclobenzaprine	
(Cyclobenzaprine)	C

Date:04/03/03ISR Number: 4087549-9Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 190185

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 300MG BID PO	Confusional State		Neurontin 300mg	PS		ORAL
Initial or Prolonged 7.5MG Q4H PRN	Hallucination		Hydrocodone 7.5mg	SS		ORAL
PO	Mental Status Changes					

Maxzide	C
Atenolol	C
Lisinopril	C
Mom	C
Cascara	C
Vioxx	C

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Transient Ischaemic Attack	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (300, THREE TIMES DAILY), ORAL		Drug Effect Decreased Dysaesthesia Nerve Injury Paraesthesia	Consumer	Neurontin (Gabapentin)	PS		ORAL

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Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C
 Ibuprofen
 (Ibuprofen) C
 Allegra-D
 (Pseudoephedrine
 Hydrochloride,
 Fexofenadine) C
 Mometasone Furoate
 (Mometasone Furoate) C
 Cyclobenzaprine
 Hydrochloride
 (Cyclobenzaprine
 Hydrochloride) C
 Esomeprazole
 (Esomeprazole) C
 Iron (Iron) C
 Vicodin
 (Paracetamol,
 Hydrocodone
 Bitartrate) C

Date:04/03/03ISR Number: 4089714-3Report Type:Expedited (15-DaCompany Report #2003012703
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Health	Neurontin	PS		ORAL
Other		Anger	Professional	(Gabapentin)			
ORAL		Drug Dependence		Oxycodone			
		Imprisonment		Hydrochloride			
				(Oxycodone			
				Hydrochloride)	C		
				Oxycocet			
				(Paracetamol,			
				Oxycodone			
				Hydrochloride)	C		
				Panadeine Co			
				(Codeine Phosphate,			
				Paracetamol)	C		
				Cyclobenzaprine			
				Hydrochloride			
				(Cyclobenzaprine			

Date:04/03/03ISR Number: 4091525-XReport Type:Expedited (15-DaCompany Report #2003008542
Age:62 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Arthralgia
Initial or Prolonged	Drug Ineffective
	Dyspnoea
	Gamma-Glutamyltransferase
	Increased
	Hyponatraemia
	Leukocytosis
	Oedema
	Oliguria
	Urine Sodium Increased

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Weight Decreased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, THREE TIMES)		Foreign Health Professional	Neurontin (Gabapentin)	PS		
200 MG (DAILY), ORAL			Fenofibrate (Fenofibrate)	SS		ORAL
50 MG (DAILY), ORAL			Anethole Trithione (Anethole Trithione)	SS		ORAL
25 MG (DAILY), ORAL			Clomipramine Hydrochloride (Clomipramine Hydrochloride)	SS		ORAL
1 MG (DAILY), ORAL			Clonazepam (Clonazepam)	SS		ORAL
ORAL			Endotelon (Vitis Vinifera, Herbal Extracts Nos)	SS		ORAL
			Cefpodoxime Proxetil (Cefpodoxime Proxetil)	C		
			Carbocisteine (Carbocisteine)	C		
			Dantrolene Sodium (Dantrolene Sodium)	C		
			Lansoprazole (Lansoprazole)	C		

Date:04/03/03ISR Number: 4091533-9Report Type:Expedited (15-DaCompany Report #2003000197
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged 900 MG (300 MG, TID) ORAL	Blood Bilirubin Increased Liver Function Test Abnormal	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:04/03/03ISR Number: 4091535-2Report Type:Expedited (15-DaCompany Report #2003012697
Age:28 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening 1800 MG (TID), ORAL	Electrocardiogram Qt Prolonged Ventricular Tachycardia	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
			Bisoprolol (Bisoprolol)	C		
			Lansoprazole (Lansoprazole)	C		
			Methadone (Methadone)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/04/03ISR Number: 4090087-0Report Type:Expedited (15-DaCompany Report #PHNU2003DE01380

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 50 MG/DAY, ORAL		Drug Interaction Fatigue Pain In Extremity Weight Increased	Foreign Health Professional Other	Lioresal (Baclofen) Tablet, 10mg Neurontin (Gabapentin) Saroten "Bayer Vital" (Amitriptyline Hydrochloride) L-Thyroxin "Henning Berlin" Candesartan (Candesartan) Oxybutynin (Oxybutynin)	PS SS SS C C C	 Bayer Vital	ORAL
1800MG/DAY							

Date:04/08/03ISR Number: 4091110-XReport Type:Expedited (15-DaCompany Report #2003013056

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 800 MG (DAILY), UNKNOWN		Agitation Back Pain Irritability Memory Impairment	Consumer	Neurontin (Gabapentin) Lamotrigine (Lamotrigine)	PS SS		
200 MG (DAILY), ORAL				All Other Therapeutic Products	C		ORAL

Age: Gender:Female I/FU:F

Outcome	PT
Disability	Abdominal Distension
	Aggression
	Asthenia
	Blood Cholesterol
	Increased
	Blood Triglycerides
	Increased
	Cardiovascular Disorder
	Chest Pain
	Constipation
	Depression
	Difficulty In Walking
	Dizziness
	Dyspnoea
	Fall
	Ganglion
	Hair Growth Abnormal
	Insomnia
	Irritability
	Mass
	Nasal Congestion
	Nasal Disorder

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Neuropathy Peripheral Oedema Oedema Peripheral	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
2000 MG (400 MG, 5X/DAY); ORAL (SEE IMAGE)		Pain Peripheral Coldness Psychotic Disorder Sciatica Spinal Disorder Treatment Noncompliance Tremor Weight Increased		Amitriptyline Hydrochloride Diazepam Sertraline Venlafaxine Alprazolam Acetylsalicylic Acid	C C C C C		

Date:04/08/03ISR Number: 4091178-0Report Type:Expedited (15-DaCompany Report #2003012425
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG Other (TID), ORAL		Adrenal Insufficiency	Foreign Health Professional	Neurontin (Gabapentin) Clomipramine Hydrochloride (Clomipramine Hydrochloride)	PS C		ORAL

Date:04/08/03ISR Number: 4091185-8Report Type:Expedited (15-DaCompany Report #2003002971
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Foreign	Neurontin			

300 MG (100	Gingivitis	Consumer	(Gabapentin)	PS	ORAL
MG, TID),	Pyrexia	Health			
ORAL	Shock	Professional			
	Therapeutic Response				
	Unexpected				

Date:04/08/03ISR Number: 4091186-XReport Type:Expedited (15-DaCompany Report #2003013019
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthropathy	Foreign	Neurontin			
1500 MG		Osteonecrosis	Health	(Gabapentin)	PS		ORAL
(UNKNOWN),			Professional				
ORAL				Valporate Sodium	C		

Date:04/08/03ISR Number: 4091195-0Report Type:Expedited (15-DaCompany Report #2003013664
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mental Disorder	Literature	Gabapentin			
			Health	(Gabapentin)	PS		
			Professional				

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Freedom Of Information (FOI) Report

Date:04/08/03ISR Number: 4091287-6Report Type:Expedited (15-DaCompany Report #2003013663

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Unevaluable Event	Literature Health Professional	Gabapentin (Gabapentin)	PS		

Date:04/08/03ISR Number: 4091305-5Report Type:Expedited (15-DaCompany Report #2003013662

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mental Disorder	Literature Health Professional	Gabapentin (Gabapentin)	PS		

Date:04/08/03ISR Number: 4091307-9Report Type:Expedited (15-DaCompany Report #2003013660

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Unevaluable Event	Literature Health Professional	Gabapentin (Gabapentin)	PS		

Date:04/08/03ISR Number: 4091333-XReport Type:Expedited (15-DaCompany Report #2003013658

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Literature Health Professional	Gabapentin (Gabapentin)	PS		
Other							

Date:04/08/03ISR Number: 4091338-9Report Type:Expedited (15-DaCompany Report #2003013657

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Gabapentin (Gabapentin)	PS		

Date:04/08/03ISR Number: 4091516-9Report Type:Expedited (15-DaCompany Report #2003013656
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Disability		Drug Toxicity	Literature Health Professional	Gabapentin (Gabapentin)	PS		
Other				All Other Therapeutic Products	SS		

Date:04/08/03ISR Number: 4091557-1Report Type:Expedited (15-DaCompany Report #2003009893
 Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Blister Dysphagia
Other	Erythema

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Face Oedema Pain In Extremity Pruritus					
		Rash Tongue Oedema	Consumer Health	Benadryl Injection (Diphenhydramine)	PS		
INTRAVENOUS	(ONCE),	Urticaria	Professional				
INTRAVENOUS				Neurontin (Gabapentin)	SS		ORAL
500 MG							
(TID),	ORAL						
				Ibuprofen (Ibuprofen)	SS		ORAL
ORAL							
				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		

Date:04/08/03ISR Number: 4092174-XReport Type:Expedited (15-DaCompany Report #USA-2002-0001677
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alopecia	Consumer	Oxycontin Tablets 10			
Hospitalization - Initial or Prolonged		Amnesia Angiopathy Anxiety	Health Professional Other	Mg(Oxycodone Hydrochloride) Cr Tablet	PS		
10 MG							
		Arterial Bruit Arterial Occlusive Disease Arthritis		Oxycontin Tablets 20mg(Oxycodone Hydrochloride) Cr Tablet	SS		
20 MG	1008 DAY						
		Back Pain Carpal Tunnel Syndrome Chest Pain Coma Depression		Hydrocodone Bitartrate (Similar To Ind 59, 175)(Hydrocodone Bitartrate) Unknown	SS		
1008 DAY							
		Disturbance In Attention		Alprazolam(Alprazola			

1008 DAY	Drug Ineffective	m)	SS	
1008 DAY	Emphysema	Ephedrine (Ephedrine)	SS	
1008 DAY	Fatigue Flushing	Pseudoephedrine (Pseudoephedrine)	SS	
75 MG, DAILY	1008 DAY	Influenza Insomnia	Effexor (Venlafaxine Hydrochloride)	SS
1008 DAY	Irritability	Codeine (Codeine)	SS	
100 MG, TID	1008 DAY	Libido Decreased Lymphadenopathy	Neurontin (Gabapentin)	SS
1008 DAY	Nausea Nervousness	Metoclopramide (Metoclopramide)	SS	
1008 DAY	Night Sweats Overdose	Quetiapine (Quetiapine)	SS	
1008 DAY	Pneumonia Pulmonary Oedema Pulse Absent Rash Generalised Rash Pruritic Rash Scaly Somnolence Upper Respiratory Tract Infection Urinary Incontinence Weight Increased	Claritin (Loratadine) Vitamin C (Ascorbic Acid) Celebrex (Celecoxib) Prilosec (Omeprazole) Daypro (Oxaprozin) Trandate (Labetalol Hydrochloride) Axid (Nizatidine) Paxil (Paroxetine Hydrochloride) Medrol	C C C C C C C C C	

Freedom Of Information (FOI) Report

(Methylprednisolone)	C
Macro Antioxidant	
(Ascorbic Acid,	
Cystine, Tocopherol,	
Calcium Ascorbate,	
Beta Carotene,	C
Zocor (Simvastatin)	C
Risperdal	
(Risperidone)	C
Fioricet	
(Butalbital)	C
Phenobarbital	
(Phenobarbital)	C
Donnatal (Atropine	
Sulfate, Hyoscine	
Hydrobromide,	
Hyoscyamine Sulfate,	
Phenobarbital)	C
Bellergal-S	
(Belldadonna	
Alkaloids,	
Erogotamine	
Tartrate,	C
Weight Loss	
Supplement (Does Not	
Code)	C
Ibuprofen	C
Ambien (Zolpidem	
Tartrate)	C
Baclofen(Baclofen)	C
Trandate (Labetalol	
Hydrochloride)	C
Celexa (Citalopram	
Hydrobromide)	C
Cortisone	
(Cortisone)	C
Atarax (Hydroxyzine	
Hydrochloride)	C
Proventil Tablet	
(Salbutamol Sulfate)	C
Amitriptyline	
(Amitriptyline)	C
Cyclobenzaprine	
(Cyclobenzaprine)	C
Augmentin	
(Amoxicillin	
Trihydrate,	
Clavulanate	

Potassium) C
Ergobel
(Nicergoline) C

Date:04/08/03ISR Number: 4092328-2Report Type:Expedited (15-DaCompany Report #USA-2003-0005143
Age:40 YR Gender:Male I/FU:I

Outcome PT
Death Accidental Overdose
Brain Herniation
Brain Oedema
Cardiomegaly
Coma

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Freedom Of Information (FOI) Report

Dose	Duration	Coronary Artery Atherosclerosis Drug Toxicity	Report Source	Product	Role	Manufacturer	Route
		Emphysema	Consumer	Oxycontin Tablets			
		Hepatic Cirrhosis	Health	(Oxycodone			
		Hepatic Steatosis	Professional	Hydrochloride) Cr			
		Hepatosplenomegaly	Other	Tablets	PS		
		Myocardial Ischaemia		Neurontin			
		Pulmonary Oedema		(Gabapentin)	SS		
		Skin Discolouration		Chlordiazepoxide	SS		
		Toxicologic Test Abnormal		Nordazepam			
				(Nordazepam)	SS		
				Nicotine (Nicotine)	SS		
				Depakote (Valproate			
				Semisodium)	C		

Date:04/09/03ISR Number: 4092451-2Report Type:Expedited (15-DaCompany Report #DEU-2002-0000223
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Insomnia	Foreign	Oxygesic 40			
		Tooth Disorder	Other	Mg(Oxycodone			
				Hydrochloride) Cr			
				Tablet	PS		ORAL
40 MG, TID,							
ORAL							
				Neurontin(Gabapentin			
)	SS		ORAL
600 MG, Q6H,							
ORAL							
				Baclofen(Baclofen)	SS		
30 MG, DAILY,							
ORAL							
				Tramal	C		

Date:04/09/03ISR Number: 4092825-XReport Type:Expedited (15-DaCompany Report #2003014093
 Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Epilepsy	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
1200 MG (400, THREE TIMES DAILY), ORAL				Lamotrigine (Lamotrigine)	C		

Date:04/09/03ISR Number: 4093217-XReport Type:Expedited (15-DaCompany Report #2003006906
Age: Gender:Female I/FU:F

Outcome	PT
Other	Aphonia Blood Pressure Decreased Bronchitis Cold Sweat Convulsion Dizziness Epilepsy Fall Gait Disturbance Joint Sprain Movement Disorder

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FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Muscle Twitching Oedema Peripheral Painful Respiration			
ORAL		Consumer Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Professional	Topiramate (Topiramate)	SS		
			Weight Increased Naproxen (Naproxen)	SS		ORAL
			White Blood Cell Count Increased			
			Oxcarbazepine (Oxcarbazepine)	C		
			Pravastatin Sodium (Pravastatin Sodium)	C		
			All Other Therapeutic Products	C		
			Hydrocortisone (Hydrocortisone)	C		
			Alendronate Sodium (Alendronate Sodium)	C		
			Levothyroxine Sodium (Levothyroxine Sodium)	C		
			Lansoprazole (Lansoprazole)	C		

Date:04/09/03ISR Number: 4096576-7Report Type:Expedited (15-DaCompany Report #US-SHR-03-002855
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Apnoea Blood Cholesterol Increased	Consumer Health Professional	Betaseron (Interferon Beta - 1b) Injection	PS		
SUBCUTANEOUS	8 MIU, EVERY 2D,	Bradycardia					
SUBCUTANEOUS		Coronary Artery Occlusion					
SEE IMAGE		Somnolence		Klonopin (Clonazepam)	SS		ORAL
				Neurontin			

(Gabapentin)	SS
Effexor - Slow Release (Venlafaxine Hydrochloride)	C
Levothyroxine (Levothyroxine)	C
Potassium Chloride	C
Zocor "Msd" (Simvastatin)	C
Evista (Raloxifene Hydrochloride)	C
Baclofen	C
Morphine (Morphine)	C
Nexium (Esomeprazole)	C
Aminopyridine (Fampridine)	C
Nasacort Ao (Triamcinolone Acetonide)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/10/03ISR Number: 4093546-XReport Type:Expedited (15-DaCompany Report #2003014027

Age:23 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Diabetes Mellitus	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
Hospitalization - 1800 MG, ORAL		Rhabdomyolysis					
Initial or Prolonged		Sepsis	Professional	Loxapine Hydrochloride (Loxapine Hydrochloride)	C		
Other				Citalopram Hydrobromide (Citalopram Hydrobromide)	C		
				Valproic Acid (Valproic Acid)	C		
				Procyclidine (Procyclidine)	C		
				Methylphenidate Hydrochloride (Methylphenidate Hydrochloride)	C		

Date:04/10/03ISR Number: 4093901-8Report Type:Expedited (15-DaCompany Report #A0401949A

Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Aggression Convulsion	Literature Health	Lamictal Tablet (Lamotrigine)	PS		ORAL
25000 MG/ SINGLE DOSE/ ORAL		Electrocardiogram Pr Prolongation	Professional				
30000 MG/ SINGLE DOSE/ ORAL		Electrocardiogram Qrs Complex Prolonged		Gabapentin Tablet (Gabapentin)	SS		ORAL
		Electrocardiogram Qt Corrected Interval Prolonged					

Hypertension
Overdose
Suicide Attempt
Tachycardia

Date:04/10/03ISR Number: 4094076-1Report Type:Expedited (15-DaCompany Report #2003003425

Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Pressure Increased Difficulty In Walking	Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG (BID), ORAL		Diplopia Headache Medication Error Paralysis		Dyazide (Hydrochlorothiazide , Triamterene) Caffeine (Caffeine) Paracetamol (Paracetamol) Butalbital W/Aspirin, Caffeine (Acetylsalicylic Acid, Caffeine, Butalbital) Axotal (Old Form)	C C C C		

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Freedom Of Information (FOI) Report

(Caffeine,
Butalbital,
Paracetamol) C

Date:04/11/03ISR Number: 4094705-2Report Type:Expedited (15-DaCompany Report #2003013019
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Osteonecrosis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1500 MG, ORAL			Professional	Valproate Sodium (Valproate Sodium)	C		

Date:04/14/03ISR Number: 4094770-2Report Type:Expedited (15-DaCompany Report #2003014027
Age:22 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Diabetes Mellitus	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
Hospitalization - 1800 MG		Insulin-Dependent					
Initial or Prolonged (TID), ORAL		Diabetic Hyperosmolar	Professional				
Other		Coma		Citalopram (Citalopram0	SS		ORAL
20 MG		Haemodialysis					
(DAILY), ORAL		Rhabdomyolysis					
		Sepsis		Loxapine Hydrochloride (Loxapine Hydrochloride)	C		
		Streptococcal Infection		Valproic Acid (Valproic Acid)	C		
				Procyclidine (Procyclidine)	C		
				Methylphenidate Hydrochloride (Methylphenidate Hydrochloride)	C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Foreign	Neurontin			
		Epilepsy	Consumer	(Gabapentin)	PS		ORAL
2300 MG, ORAL							
				Magnesium			
				(Magnesium)	SS		ORAL
450 MG (TID),							
ORAL							

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Trigeminal Neuralgia	Consumer	Neurontin			
				(Gabapentin)	PS		

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Date:04/15/03ISR Number: 4095607-8Report Type:Expedited (15-DaCompany Report #03P-076-0215619-00

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Abdominal Pain Blood Amylase Increased Chest Pain	Foreign Health Professional	Klacid (Biaxin) (Clarithromycin) (Clarithromycin)	PS		ORAL
SEE IMAGE	Chromaturia Flatulence Headache Lipase Increased Pancreatic Disorder Porphyria Acute Pyrexia	Company Representative	Gabapentin Famotidine Drotaverine Tramadol Potassium Chloride Metamizole Benzcyclane Sodium Chloride Glucose Alprazolam Dimeticone, Activated Gabapentin Clonazepam Mixtura Pectoralis Fono	SS C C C C C C C C C C C C C C C C		

Date:04/15/03ISR Number: 4095717-5Report Type:Expedited (15-DaCompany Report #2003014428

Age:76 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN	Prostatic Operation Pruritus 900 MG (TID),	Foreign Health Professional	Neurontin (Gabapentin)	PS		
UNKNOWN			Insulin (Insulin) Lorazepam (Lorazepam) Acetylsalicylate Lysine (Acetylsalicylate Lysine) Buflomedil (Buflomedil)	C C C C C C		

Mirtazapine
(Mirtazapine) C
Vitamin B1 And B6
(Thiamine,
Pyridoxine) C
Tamsulosin
(Tamsulosin) C

Date:04/15/03ISR Number: 4095720-5Report Type:Expedited (15-DaCompany Report #2003003153
Age:62 YR Gender:Male I/FU:F

Outcome PT
Hospitalization - Dizziness
Initial or Prolonged Fibromyalgia
Haemoglobin Decreased
Headache
Hepatic Cirrhosis
Hepatomegaly
Hepatotoxicity
Joint Range Of Motion

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Dose	Duration	Decreased Nausea Pancreatic Disorder Transferrin Saturation	Report Source	Product	Role	Manufacturer	Route
1200 MG, (TID) , ORAL		Increased Vomiting	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Ethanol (Ethanol) Flupirtine Maleate (Flupirtine Maleate)	SS C		

Date:04/16/03ISR Number: 4093696-8Report Type:Direct
Age:72 YR Gender:Male I/FU:I Company Report #CTU 191085

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Screen Positive Intentional Misuse		Gabapentin	PS		

Date:04/16/03ISR Number: 4097164-9Report Type:Expedited (15-DaCompany Report #2003015241
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Convulsion	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Amitriptyline (Amitriptyline) All Other Therapeutic Products	PS C C		ORAL

Date:04/16/03ISR Number: 4097316-8Report Type:Expedited (15-DaCompany Report #2003014735
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL		Cerebral Fungal Infection	Consumer	Zoloft (Sertraline)	PS		ORAL

Initial or Prolonged	Confusional State	Health	Neurontin	
Other		Professional	(Gabapentin)	SS
			Aciclovir	
			(Aciclovir)	SS
			Anabolic Steroids	SS
			Valaciclovir	
			Hydrochloride	
			(Valaciclovir	
			Hydrochloride)	SS
			Clonazepam	
			(Clonazepam)	SS
			Digoxin	C

Date:04/16/03ISR Number: 4097319-3Report Type:Expedited (15-DaCompany Report #2003010329
Age:62 YR Gender:Female I/FU:F

Outcome	PT
Other	Back Pain
	Blood Pressure Increased
	Dizziness
	Drug Ineffective
	Eye Haemorrhage
	Malaise
	Ocular Vascular Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pain In Extremity

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), ORAL		Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Clonidine (Clonidine)	C		
			Methadone (Methadone)	C		
			Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride)	C		
			Panadeine Co (Codeine Phosphate, Paracetamol)	C		

Date:04/16/03ISR Number: 4097329-6Report Type:Expedited (15-DaCompany Report #2003014853
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Congestive Cardiomyopathy	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:04/16/03ISR Number: 4097331-4Report Type:Expedited (15-DaCompany Report #2003004961
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Benign Gastric Neoplasm Bipolar Disorder Drug Ineffective	Consumer Health Professional	Neurontin(Gabapentin) Geodon (Ziprasidone)	PS SS		ORAL
ORAL		Syncope		Lamotrigine			

(Lamotrigine) SS
 Antihypertensives C
 Diuretics C
 Aripiprazole
 (Aripiprazole) C

Date:04/16/03ISR Number: 4097332-6Report Type:Expedited (15-DaCompany Report #2002063521
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Disorder	Consumer	Neurontin			
Other		Hypertension	Health	(Gabapentin)	PS		ORAL
2400 MG		Medication Error	Professional				
(TWICE		Sudden Death					
DAILY), ORAL							

All Other
 Therapeutic Products C
 Dextropropoxyphene
 (Dextropropoxyphene) C
 Loratadine
 (Loratadine) C
 Esomeprazole
 (Esomeprazole) C

Freedom Of Information (FOI) Report

Paroxetine
 Hydrochloride
 (Paroxetine
 Hydrochloride) C
 Celecoxib
 (Celecoxib) C
 Clonazepam
 (Clonazepam) C
 Propacet
 (Paracetamol,
 Dextropropoxyphene
 Napsilate) C
 Tramadol
 Hydrochloride
 (Tramadol
 Hydrochloride) C

Date:04/16/03ISR Number: 4097333-8Report Type:Expedited (15-DaCompany Report #2003014665
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiac Operation	Consumer	Neurontin	PS		
Other		Drug Effect Decreased					
		Gallbladder Operation					
		Paraesthesia					
		Road Traffic Accident					
		Spinal Cord Injury					

Date:04/16/03ISR Number: 4097985-2Report Type:Expedited (15-DaCompany Report #2003014429
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Blood Electrolytes	Foreign	Neurontin			
Hospitalization -		Abnormal	Health	(Gabapentin)	PS		ORAL
900 MG (TID),							
Initial or Prolonged		Diabetes Mellitus	Professional				
ORAL							
		Inadequate Control		Amitriptyline			
		Drug Interaction		Hydrochloride			
		Renal Failure Acute		(Amitriptyline			

75 MG (TID),

ORAL

25 MG

(DAILY), ORAL

Hydrochloride)	SS	ORAL
Rofecoxib (Rofecoxib)	SS	ORAL
Xipamide (Xipamide)	C	
Benzbromarone (Benzbromarone)	C	
Benazepril Hydrochloride (Benazepril Hydrochloride)	C	
Carvedilol (Carvedilol)	C	
Furosemide (Furosemide)	C	
Insulin Lispro (Insulin Lispro)	C	
Morphine Sulfate (Morphine Sulfate)	C	
Pantoprazole Sodium		

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Freedom Of Information (FOI) Report

(Pantoprazole Sodium) C
 Phenytoin (Phenytoin) C
 Lorazepam (Lorazepam) C

Date:04/16/03ISR Number: 4097987-6Report Type:Expedited (15-DaCompany Report #USA-2002-0001677
 Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alopecia	Consumer	Oxycontin Tablet 10			
Hospitalization - Initial or Prolonged		Angiopathy	Health Professional	Mg(Oxycodoen Hydrochloride) Cr			
10 MG ,		Apnoea	Other	Tablet	PS		
		Arterial Occlusive Disease		Oxycontin Tablet 20			
20 MG	1008 DAY	Arthropathy		Mg(Oxycodoen Hydrochloride) Cr			
		Asthma		Tablet	SS		
		Atherosclerosis		Hydrocodone			
		Back Pain		Bitartrate Similar			
		Cardiac Arrest		To Ind			
1008 DAY		Carpal Tunnel Syndrome		59,175)(Hydrocodone			
		Chest Pain		Betartrate)Unknown	SS		
		Coma		Alprazolam			
1008 DAY		Depression		(Alprazolam)	SS		
		Disturbance In Attention		Ephedrine (Epedrine)	SS		
1008 DAY		Drug Ineffective		Pseudoephedrine			
		Fatigue		(Pseudoephedrine)	SS		
		Finger Deformity		Effexor (Venlafaxine			
75 MG, DAILY	1008 DAY	Flushing		Hydrochloride)	SS		
		Hilar Lymphadenopathy		Codeine (Codeine)	SS		
1008 DAY		Hypoaesthesia		Neurontin			
		Influenza Like Illness		(Gabapetin)	SS		
100 MG, TID	1008 DAY	Insomnia		Metoclopramide			

1008 DAY	Libido Decreased	(Metoclopramide)	SS
	Liver Disorder	Quetiapine	
	Memory Impairment	(Quetiapine)	SS
1008 DAY	Myofascial Pain Syndrome	Claritin	
	Nausea	(Loratadine)	C
	Nervousness	Vitamin C (Ascorbic	
	Night Sweats	Acid)	C
	Osteoarthritis	Celebrex (Celecoxib)	C
	Overdose	Prilosec	
	Paraesthesia	(Omeprazole)	C
	Peripheral Vascular	Daypro (Oxaprozin)	C
	Disorder	Trandate (Labetalol	
	Pneumonia	Hydrochloride)	C
	Pulmonary Interstitial	Axid (Nizatidine)	C
	Emphysema Syndrome	Paxil (Paroxetine	
	Pulmonary Oedema	Hydrochloride)	C
	Pulse Abnormal	Medrol	
	Rash Pruritic	(Methylprednisolone)	C
	Rash Scaly	Marco Antioxidant	
	Shock	(Ascorbic Acid,	
	Somnolence	Cystine, Tocopherol,	
	Upper Respiratory Tract	Calcium Ascorgate,	
	Infection	Betacarotene, Manganese	C
	Urinary Incontinence	Zocor (Simvastatin)	C
	Weight Increased	Risperdal	
		(Risperidone)	C

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Fioricet (Butalbital)	C
Flonase (Fluticasone Propionate)	C
Phenobarbital (Phenobarbital)	C
Donnatal (Atropine Sulfate, Hyoscine Hydrobromide, Hyoscyamine Sulfate, Phenobarbital)	C
Bellergal-S (Belladonna Alkaloids, Ergotamine Tartrate, Phenobarbital)	C
Weight Loss Supplement (Does Not Code)	C
Ibuprofen	C
Ambien(Zolpidem Tartrate)	C
Balcofen (Balcofen)	C
Trandate (Labetalol Hydrochloride)	C
Celexa (Citalopram Hydrobromide)	C
Cortisone (Cortisone)	C
Atarax (Hydroxyzine Hydrochloride)	C
Proventil Tablet (Salbutamol Sulfate)	C
Amitriptyline (Amitriptyline)	C
Cyclobenzaprine (Cyclobenzaprine)	C
Aaugmentin (Amoxicillin Trihydrate, Clavulanate Potassium)	C
Ergobel (Nicergoline)	C

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Cancer	Consumer	Neurontin			
Other		Condition Aggravated	Health	(Gabapentin)	PS		ORAL
400 MG (BID),		Diabetes Mellitus	Professional				
ORAL		Mitral Valve Disease		Glucotrol			
2.5 MG				(Glipizide)	SS		ORAL
(DAILY), ORAL				Digoxin (Digoxin)	C		
				Ramipril (Ramipril)	C		
				Pramipexole			
				(Pramipexole)	C		
				Entacapone			

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Freedom Of Information (FOI) Report

(Entacapone) C
 Sinemet (Levodopa,
 Carbidopa) C
 Levothyroxine Sodium
 (Levothyroxine
 Sodium) C

Date:04/17/03ISR Number: 4098232-8Report Type:Expedited (15-DaCompany Report #EMADSS2003002954
 Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchopneumonia	Foreign	Durogesic (100			
Hospitalization -		Confusional State	Health	Mcg/Hr Patch)			
Initial or Prolonged		Formication	Professional	(Fentanyl)	PS		
TRANSDERMAL	100 MCG/H, 1	Hallucination					
IN 72		Headache					
HOURS(S)		Pain					
TRANSD		Restlessness		Finlepsin (Carbamazepine)	SS		ORAL
600 MG, 2 IN							
1 DAILY, ORAL				Neurontin(Gabapentin)	SS		ORAL
300 MG, 3 IN							
1 DAILY, ORAL				Saroten (Amitriptyline Hydrochloride)	SS		ORAL
10 MG, 3 IN 1							
DAILY, ORAL				Lasix (Furosemide)	C		
				Novalgin (Metamizole Sodium)	C		
				Rytmonorm (Propafenone) Hydrochloride	C		

Date:04/17/03ISR Number: 4098276-6Report Type:Expedited (15-DaCompany Report #2003015129
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Decreased Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Feeling Abnormal Formication Heart Rate Decreased Syncope		Trazodone (Trazodone) Clopidogrel Sulfate (Clopidogrel Sulfate)	C C		

Date:04/17/03ISR Number: 4098277-8Report Type:Expedited (15-DaCompany Report #2003015349
Age: Gender:Female I/FU:I

Outcome	PT
Other	Abnormal Behaviour Anhedonia Autism Cognitive Disorder Crying Drug Level Increased Drug Toxicity Mutism Nightmare

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
300 MG	(DAILY), ORAL	Oligodipsia Oral Intake Reduced Respiratory Arrest	Consumer	Neurontin (Gabapentin)	PS		ORAL
750 MG	(DAILY), ORAL	Screaming Weight Gain Poor		Valproate Semisodium (Valproate Semisodium)	SS		ORAL
				Lamotrigine (Lamotrigine)	SS		
				Topiramate (Topiramate)	SS		

Date:04/17/03
 Age: 31
 Gender:Male
 I/FU:F
 ISR Number: 4098292-4
 Report Type:Expedited (15-Da
 Company Report #DEU-2002-0000223

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
40 MG, TID,	ORAL	Insomnia Sleep Disorder Tooth Disorder	Foreign Other	Oxygesic 40 Mg (Oxycodone Hydrochloride) Cr Tablet	PS		ORAL
600 MG, Q6H,	ORAL			Neurontin (Gabapentin)	SS		ORAL
30 MG. DAILY	ORAL			Baclofen	SS		ORAL
				Tramal	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchopneumonia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Hospitalization - 900 MG (THREE Initial or Prolonged TIMES DAILY), Other ORAL		Deterioration	Professional				
TOPICAL	100 MCG/H),	Hallucination		Fentanyl (Fentanyl)	SS		
TOPICAL		Headache Pain					
1200 MG, (TWICE DAILY), ORAL		Paraesthesia Restlessness		Carbamazepine (Carbamazepine)	SS		ORAL
30 MG (THREE TIMES DAILY), ORAL				Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		ORAL
				Furosemide (Furosemide)	C		
				Metamizole Sodium (Metamizole Sodium)	C		
				Propafenone Hydrochloride (Propafenone Hydrochloride)	C		

Freedom Of Information (FOI) Report

All Other
Therapeutic Products C

Date:04/18/03ISR Number: 4095706-0Report Type:Direct
Age:41 YR Gender:Female I/FU:I

Company Report #CTU 191223

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	6ML 2X /DAY	Allergy To Chemicals Alopecia		Neurontin 250ml Parke Davis	PS	Parke Davis	ORAL
ORAL		Apathy					
		Asthenia Blood Glucose Decreased Condition Aggravated Depression Disturbance In Attention Drug Withdrawal Syndrome Dysphonia Haemorrhagic Diathesis Increased Tendency To Bruise Mouth Plaque Muscle Spasms Obsessive-Compulsive Disorder Oedema Personality Change Suicidal Ideation Visual Disturbance Weight Decreased		Zoloft	C		

Date:04/18/03ISR Number: 4095834-XReport Type:Direct
Age:30 YR Gender:Female I/FU:I

Company Report #CTU 191252

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 600MG 1 DAY Initial or Prolonged ORAL Disability		Convulsion		Neurontin 300mg	PS		ORAL

Other

Date:04/18/03ISR Number: 4096044-2Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 191259

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Confusional State		Hydrocodone 7.5/650	PS		ORAL
1 TAB PO QID						
Initial or Prolonged	Hallucination		Neurontin 300mg	SS		ORAL
1TAB PO BID						
	Mental Status Changes		Atenolol	C		
	Nausea		Lisinopril	C		
	Vomiting		Hydrochlorothiazide	C		

Date:04/18/03ISR Number: 4098862-3Report Type:Expedited (15-DaCompany Report #2002068857
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Drug Effect Decreased	Consumer	Neurontin			
900 MG (UNK),	Pain	Health	(Gabapentin)	PS		ORAL
ORAL		Professional				
			Vicodin			

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FDA - Adverse Event Reporting System (AERS)

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
UNK (UNK),				(Paracetamol, Hydrocodone Bitartrate)	SS		ORAL
ORAL							
UNKNOWN		UNK (UNK),		Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		
UNKNOWN							
				Paroxetine Hydrochloride Baclofen	C C		

Date:04/18/03ISR Number: 4099053-2Report Type:Expedited (15-DaCompany Report #2003015897
Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Extrapyramidal Disorder	Foreign Health	Neurontin (Gabapentin)	PS		
1600 MG			Professional Company Representative	Sinemet (Levodopa, Carbidopa) Pravastatin Sodium (Pravastatin Sodium) Venlafaxine (Venlafaxine) Sertraline (Sertraline) Morphine Sulfate (Morphine Sulfate) Paracetamol (Paracetamol)	C C C C C		

Date:04/21/03ISR Number: 4099329-9Report Type:Expedited (15-DaCompany Report #2003015172
Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abdominal Distension	Consumer	Neurontin			

Initial or Prolonged	Asthenia	(Gabapentin)	PS	ORAL
300 MG				
Other	Blood Pressure Diastolic			
(DAILY), ORAL				
	Decreased	Lisinopril		
	Dizziness	(Lisinopril)	C	
	Dyspnoea	Antihypertensives	C	
	Fatigue	Verapamil		
	Gait Disturbance	(Verapamil)	C	
	Heart Rate Decreased	Estrogens Conjugated		
	Heart Rate Irregular	(Estrogens		
	Muscular Weakness	Conjugated)	C	
	Oedema			
	Oedema Peripheral			
	Renal Failure Acute			
	Sedation			

Date:04/21/03ISR Number: 4099468-2Report Type:Expedited (15-DaCompany Report #2003015736
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Parkinson'S Disease	Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG,							
(DAILY), ORAL							

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Freedom Of Information (FOI) Report

Tolterodine
 L-Tartrate
 (Tolterodine L-Tartrate) C
 Digoxin (Digoxin) C
 Furosemide
 (Furosemide) C
 Metoprolol Succinate
 (Metoprolol Succinate) C
 Metolazone
 (Metolazone) C
 Hydrocodone
 (Hydrocodone) C
 Metformin
 (Metformin) C
 Potassium Chloride
 (Potassium Chloride) C
 Neutra-Phos (Sodium Phosphate Dibasic,
 Sodium Phosphate Monobasic
 (Anhydrate), C
 Pantoprazole
 (Pantoprazole) C
 Trazodone
 (Trazodone) C
 Atorvastatin
 (Atorvastatin) C
 Insulin Human
 (Insulin Human) C
 Insulin Glargine
 (Insulin Glargine) C

Date:04/21/03ISR Number: 4099469-4Report Type:Expedited (15-DaCompany Report #001-0073-M0100134
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Burning Sensation	Consumer	Dilantin (Phenytoin Sodium)	PS		ORAL
Other		Convulsion					
300 MG		Drug Effect Decreased					
(DAILY), ORAL		Drug Interaction		Neurontin			

300 MG, ORAL	Drug Level Increased	(Gabapentin)	SS	ORAL
	Ear Pain	Zyrtec (Tablets)		
	Eye Irritation	(Cetirizine)	SS	ORAL
5 MG (2.5,	Headache			
TWICE DAILY),	Hypersensitivity			
ORAL	Hypothyroidism	Levothyroxine Sodium		
	Medication Error	(Levothyroxine		
	Memory Impairment	Sodium)	SS	ORAL
(TID), ORAL	Mouth Haemorrhage	Zinc Gluconate (Zinc		
	Oedema Peripheral	Gluconate)	SS	
	Pruritus	Centrum (Vitamins		
	Sleep Disorder	Nos, Minerals Nos)	C	
	Somnolence	Calcium Carbonate		
	Tremor	(Calcium Carbonate)	C	
	Urticaria	Levetiracetam		
	Vision Blurred	(Levetiracetam)	C	
	Weight Decreased			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/21/03ISR Number: 4099587-0Report Type:Expedited (15-DaCompany Report #2003015146

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Epilepsy	Foreign	Neurontin			
		Weight Increased	Health	(Gabapentin)	PS		
			Professional				
			Company				
			Representative				

Date:04/22/03ISR Number: 4099875-8Report Type:Expedited (15-DaCompany Report #03P-056-0216336-00

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Intentional Misuse	Foreign	Depakine (Depakene)			
Initial or Prolonged		Loss Of Consciousness	Other	(Sodium			
				Valproate/Valproic			
				Acid) (Sodium			
				Valproate/Valproic	PS		ORAL
ORAL							
				Zolpidem	SS		ORAL
10 MG, 30 IN							
ONCE, ORAL							
				Tri-Therapy Drugs	SS		
				Bromazepam	SS		ORAL
6 MG, ORAL							
				Hydroxyzine			
				Hydrochloride	SS		ORAL
ORAL							
				Gabapentin	SS		ORAL
ORAL							
				Ibuprofen	SS		ORAL
ORAL							

Date:04/22/03ISR Number: 4099925-9Report Type:Expedited (15-DaCompany Report #2003015534

Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Pressure Increased	Foreign	Neurontin			

Initial or Prolonged 300 MG, (DAILY), ORAL	Constipation Neuropathy Peripheral Oedema	Health Professional	(Gabapentin)	PS	ORAL
UNKNOWN (DAILY)	0.5 MG		Clonazepam (Clonazepam)	SS	
			Panadeine Co (Codeine Phosphate, Paracetamol) Omeprazole(Omeprazole) Zopiclone(Zopiclone) Verapamil(Verapamil) Insulin(Insulin) Metformin(Metformin) Simvastatin(Simvastatin) Hydrochlorothiazide(hydrochlorothiazide) Enalapril(Enalapril) Polyvinyl Alcohol(Polyvinyl Alcohol)	C C C C C C C C C C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/22/03ISR Number: 4099952-1Report Type:Expedited (15-DaCompany Report #2003015955

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anticonvulsant Drug Level Below Therapeutic	Foreign Health	Epanutin Suspension (Phenytoin Sodium)	PS		ORAL
ORAL		Drug Interaction Epilepsy	Professional	Neurontin (Gabapentin)	SS		ORAL
600 MG (300							
BID), ORAL							

Date:04/22/03ISR Number: 4099953-3Report Type:Expedited (15-DaCompany Report #2003016209

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased Dry Mouth	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Neuropathy Peripheral	Professional				

Date:04/22/03ISR Number: 4100862-1Report Type:Expedited (15-DaCompany Report #2003005368

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams Cataract	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
400 MG (100							
MG, FOUR		Cognitive Disorder	Professional				
TIMES), DAILY,		Memory Impairment					
ORAL							

Acetylsalicylic Acid (Acetylsalicylic Acid) C
 Calcium (Calcium) C
 Multivitamins

(Ergocalciferol,
Ascorbic Acid, Folic
Acid, Thiamine
Hydrochloride, C

Date:04/22/03ISR Number: 4100864-5Report Type:Expedited (15-DaCompany Report #2003015764
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG (600, Other QID)		Appendicectomy Atrioventricular Block Intestinal Obstruction	Health Professional	Neurontin (Gabapentin) All Other Therapeutic Products	PS C		

Date:04/22/03ISR Number: 4100867-0Report Type:Expedited (15-DaCompany Report #2003015585
Age:43 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other 3200 MG (DAILY)		Cardiomyopathy	Health Professional	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/23/03ISR Number: 4097451-4Report Type:Direct
Age: Gender: I/FU:I

Company Report #CTU 191535

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	25MG 1X DAY	3 YR	Confusional State		Topamax	PS	Ortho Mcneil	
	300MG 3X DAY	10 YR	Dehydration		Neurontin 300mg	SS	Park Davis	
	0.2MG X1		Suicidal Ideation		Clonidine Hcl 0.2mg	SS	Mylan Pharmaceutical	
			Tongue Disorder		Triamterine Hctz			
			Tremor		95/50mg	SS	Gineva Generic	

Date:04/23/03ISR Number: 4101879-3Report Type:Expedited (15-DaCompany Report #03P-056-0216333-00
Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Intentional Misuse Loss Of Consciousness	Foreign Other	Depakine Tablets (Sodium Valproate) (Sodium Valproate) (Sodium Valproate)	PS		ORAL
	PER ORAL				Zolpidem	SS		ORAL
	10 MG, 30 IN							
	ONCE, PER							
	ORAL				Bromazepam	SS		ORAL
	6 MG, ONCE,							
	PER ORAL							
					Hydroxyzine Hydrochloride	SS		ORAL
	ONCE, PER							
	ORAL				Gabapentin	SS		ORAL
	ONCE, PER							
	ORAL							
	ONCE, PER				Ibuprofen	SS		ORAL

ORAL

Date:04/23/03ISR Number: 4102144-0Report Type:Expedited (15-DaCompany Report #2003015898

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG		Blood Prolactin Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL				All Other Therapeutic Products	C		

Date:04/24/03ISR Number: 4102503-6Report Type:Expedited (15-DaCompany Report #2003008809

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Convulsion Pharmaceutical Product	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
		Complaint	Professional	Topiramate (Topiramate) Pramipexole (Pramipexole) Furosemide (Furosemide) Potassium (Potassium) Methylphenidate Hydrochloride (Methylphenidate Hydrochloride)	SS C C C		

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Freedom Of Information (FOI) Report

Spiroinolactone
(Spiroinolactone) C

Date:04/24/03ISR Number: 4102505-XReport Type:Expedited (15-DaCompany Report #2003016621

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Consumer	Dilantin Suspension	PS		ORAL
ORAL		Alopecia		(Phenytoin Sodium)			
		Cerebrovascular Accident		Neurontin	SS		ORAL
ORAL		Convulsion		(Gabapentin)			
		Crying		Carbamazepine	SS		
		Difficulty In Walking		(Carbamazepine)			
		Drug Ineffective		All Other	SS		
		Drug Level Decreased		Therapeutic Products			
		Hypersomnia					
		Medication Error					
		Movement Disorder					
		Partial Seizures					

Date:04/25/03ISR Number: 4100819-0Report Type:Expedited (15-DaCompany Report #033-0945-990010

Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Foreign	Neurontin	PS		ORAL
600 MG		Toxic Skin Eruption	Study	(Gabapentin)			
(DAILY), ORAL			Health				
			Professional	Triatec (Ramipril)	C		
				Bromazepam	C		
				(Bromazepam)			
				Clonazepam	C		
				(Clonazepam)			
				Perindopril	C		
				(Perindopril)			
				Esberiven Forte	C		
				(Rutoside, Melilot)			
				Metformin Embonate			

(Metformin Embonate) C
Metformin
Hydrochloride
(Metformin
Hydrochloride) C

Date:04/25/03ISR Number: 4103106-XReport Type:Expedited (15-DaCompany Report #2003016415
Age:21 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Haoperidol (Haloperidol)	SS		ORAL
ORAL				Benzatropine Mesilate (Benzatropine Mesilate)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/25/03ISR Number: 4103108-3Report Type:Expedited (15-DaCompany Report #2003016399
Age:11 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia	Literature	Gabapentin			
Hospitalization - (ONCE), ORAL		Blood Ph Increased	Health	(Gabapentin)	PS		ORAL
Initial or Prolonged Other 6000 MG (ONCE), ORAL		Blood Potassium Decreased Blood Sodium Increased Cardiac Arrest Completed Suicide Convulsion Disseminated Intravascular Coagulation Hypoglycaemia Hypotension Intestinal Ischaemia Intestinal Perforation Loss Of Consciousness Oliguria Sepsis Tachycardia	Professional	Nortriptyline (Nortriptyline)	SS		ORAL

Date:04/25/03ISR Number: 4103109-5Report Type:Expedited (15-DaCompany Report #2003016396
Age:36 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Gabapentin			
Other ORAL		Completed Suicide	Health	(Gabapentin)	PS		ORAL
ORAL			Professional	Oxycodone (Oxycodone)	SS		ORAL
ORAL				Paroxetine (Paroxetine)	SS		ORAL

Date:04/25/03ISR Number: 4103110-1Report Type:Expedited (15-DaCompany Report #2003016417
Age:38 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Zolpidem (Zolpidem)	SS		ORAL
ORAL							

Date:04/25/03ISR Number: 4103119-8Report Type:Expedited (15-DaCompany Report #2003016395
Age:19 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Level Increased	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Valproic Acid (Valproic Acid)	SS		ORAL
ORAL				Olanzapine (Olanzapine)	SS		ORAL
ORAL				All Other Therapeutic Products	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/25/03ISR Number: 4103162-9Report Type:Expedited (15-DaCompany Report #2003016394

Age:54 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional				

Date:04/25/03ISR Number: 4103187-3Report Type:Expedited (15-DaCompany Report #2003016397

Age:54 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		ORAL
ORAL				Zolpidem (Zolpidem)	SS		ORAL
ORAL				All Other Therapeutic Products	SS		ORAL

Date:04/25/03ISR Number: 4103188-5Report Type:Expedited (15-DaCompany Report #2003016619

Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Alcohol Detoxification Arthralgia	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Dysstasia Myalgia		Propranolol (Propranolol)	C		
				Valproic Acid (Valproic Acid)	C		
				Chlordiazepoxide (Chlordiazepoxide)	C		

Clonidine	
(Clonidine)	C
Furosemide	
(Furosemide)	C
Paroxetine	
Hydrochloride	
(Paroxetine	
Hydrochloride)	C
Lansoprazole	
(Lansoprazole)	C
Potassium	
(Potassium)	C
Olanzapine	
(Olanzapine)	C
Lorazepam	
(Lorazepam)	C

Date:04/25/03ISR Number: 4103189-7Report Type:Expedited (15-DaCompany Report #2003017525
Age: Gender:Female I/FU:I

Outcome	PT
Disability	Bone Pain
Other	Difficulty In Walking
	Disease Recurrence
	Drug Ineffective

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Menopausal Symptoms Muscle Spasms Neuralgia Weight Decreased	Report Source	Product	Role	Manufacturer	Route
3600 (QID), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL				Celecoxib (Celecoxib)	SS		ORAL
				Metoprolol Succinate (Metoprolol Succinate)	C		
				Esomeprazole (Esomeprazole)	C		
				Oxycodone (Oxycodone)	C		
				Ibuprofen (Ibuprofen)	C		
				Vitamins	C		
				Echinacea Extract (Echinacea Extract)	C		
				Spirolactone (Spirolactone)	C		

Date:04/25/03ISR Number: 4103190-3Report Type:Expedited (15-DaCompany Report #2003016387
Age:52 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Completed Suicide	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Galenic /Paracetamol/Codeine / (Codeine, Paracetamol)	SS		ORAL
ORAL				Fosinopril (Fosinopril)	SS		ORAL
				All Other			

ORAL

Date:04/28/03ISR Number: 4101863-XReport Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 191736

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Drug Level Decreased		Gabapentin	PS		
Initial or Prolonged	Pyrexia		Phenytoin	SS		
	Rash Erythematous					
	Rash Maculo-Papular					

Date:04/28/03ISR Number: 4103398-7Report Type:Expedited (15-DaCompany Report #2002050043
 Age: Gender:Female I/FU:F

Outcome	PT
Other	Adrenal Adenoma
	Asthenia
	Blood Arsenic Increased
	Condition Aggravated
	Creatinine Renal
	Clearance Decreased
	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
(TWICE DAILY), ORAL		Dyspepsia Dyspnoea Feeling Abnormal	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Gastrointestinal Disorder Headache	Professional				
		Helicobacter Infection		Lamotrigine (Lamotrigine)	SS		ORAL
ORAL		Hunger Hypertension		Clonazepam (Clonazepam)	SS		
		Laboratory Test Abnormal Movement Disorder Muscle Spasms Nervousness Oesophageal Spasm Pollakiuria Tardive Dyskinesia Throat Tightness Weight Increased					

Date:04/28/03ISR Number: 4104455-1Report Type:Expedited (15-DaCompany Report #2003016762
Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (DAILY), ORAL		Cardiac Failure Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Weight Increased	Professional Company Representative	Furosemide (Furosemide)	C		

Date:04/28/03ISR Number: 4104456-3Report Type:Expedited (15-DaCompany Report #2003016992
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Hepatic Failure	Foreign Health Professional	Gabapentin (Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Bone Pain	Consumer	Celebrex			
Other		Difficulty In Walking		(Celecoxib)	PS		ORAL
ORAL							
		Drug Ineffective		Neurontin			
300 MG, QD;		Feeling Hot		(Gabapentin) Capsule	SS		ORAL
		Flushing					
900 MG, QID;							
		Hyperhidrosis					
900 MG, TID,							
ORAL		Menopause					
		Muscle Spasms		Toprol Xl			
		Sleep Disorder		(Metoprolol			
		Unevaluable Event		Succinate)	C		
		Weight Decreased		Nexium	C		
				Oxycodone			
				(Oxycodone)	C		
				Motrin	C		
				Vitamins	C		
				Echinacea Extract			
				(Echinacea Extract)	C		
				Spirolactone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/30/03ISR Number: 4104735-XReport Type:Expedited (15-DaCompany Report #2003017701

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Other 600 MG (BID), ORAL	Hepatitis	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
			Prednisolone (Prednisolone)	C		
			Didronel Pmo "Norwich Eaton" (Calcium Carbonate Etidronate Disodium)	C		
			Paramol-118 (Paracetamol, Dihydrocodeine Bitartrate)	C		
			Dosulepin (Dosulepin)	C		
			Domperidone (Domperidone)	C		
			Omeprazole (Omeprazole)	C		
			Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
			Iron (Iron)	C		
			Atorvastatin (Atorvastatin)	C		
			Beclometasone Dipropionate (Beclometasone Dipropionate)	C		
			Finasteride (Finasteride)	C		
			Azathioprine (Azathioprine)	C		
			Ramipril (Ramipril)	C		
			Cyclophosphamide (Cyclophosphamide)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Condition Aggravated	Foreign	Neurontin			
Other		Liver Disorder	Literature	(Gabapentin)	PS		ORAL
600 MG		Myasthenia Gravis	Health				
(DAILY), ORAL		Neoplasm	Professional	Immunosuppressive			
		Pneumonia		Agents	C		
		Pulmonary Embolism					

Outcome	PT	Report Source
Hospitalization -	Cholestasis	Foreign
Initial or Prolonged	Hepatitis	Health
	Hepatocellular Damage	Professional
		Company

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
800 MG (DAILY)		Neurontin (Gabapentin)	PS		
		Lansoprazole Lercanidipine Hydrochloride (Lercanidipine Hydrochloride)	C C		

Date:04/30/03ISR Number: 4105635-1Report Type:Expedited (15-DaCompany Report #2003017239
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 , THREE TIMES DAILY), ORAL				Alprazolam Atenolol Allopurinol Furosemide Minoxidil	C C C C C		

Date:04/30/03ISR Number: 4105637-5Report Type:Expedited (15-DaCompany Report #2003017238
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 Other , THREE TIMES		Anaemia Asthenia Blood Iron Decreased	Health Professional	Neurontin (Gabapentin)	PS		ORAL

DAILY) ORAL

Hyperhidrosis

Lymphadenopathy
Lymphoma
Nausea
Platelet Count Decreased
Pyrexia
Serum Ferritin Decreased
Vomiting
Weight Decreased
White Blood Cell Count
Decreased

Risperidone C
Lorazepam C

Date:04/30/03ISR Number: 4105655-7Report Type:Expedited (15-DaCompany Report #2003017240
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Drug Ineffective		Insulin Injection, Isophane (Insulin Injection, Isophane)	C		
				Insulin Lispro	C		
				Diclofenac (Diclofenac)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/30/03ISR Number: 4105705-8Report Type:Expedited (15-DaCompany Report #2003010331

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nail Disorder	Health	Neurontin			
		Treatment Noncompliance	Professional	(Gabapentin)	PS		
				All Other			
				Therapeutic Products	SS		

Date:05/01/03ISR Number: 4106560-2Report Type:Expedited (15-DaCompany Report #2003008068

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Activities Of Daily	Consumer	Neurontin			
Other		Living Impaired		(Gabapentin)	PS		
		Anaemia		Acetylsalicylic Acid			
		Arthralgia		(Acetylsalicylic			
		Balance Disorder		Acid)	SS		
		Bradyphrenia		Vitamins With			
		Burns Second Degree		Minerals	SS		
		Cerebrovascular Accident					
		Convulsion					
		Coordination Abnormal					
		Crying					
		Difficulty In Walking					
		Disturbance In Attention					
		Dizziness					
		Dry Mouth					
		Fall					
		Headache					
		Heart Rate Increased					
		Hemiparesis					
		Movement Disorder					
		Muscle Disorder					
		Muscle Spasms					
		Nerve Injury					
		Nervousness					
		Pain					
		Pain In Extremity					
		Panic Attack					
		Paralysis					
		Speech Disorder					

Outcome PT
Other Adrenal Adenoma
Asthenia
Blood Heavy Metal
Increased
Condition Aggravated
Dizziness
Dyspnoea
Feeling Abnormal
Gastrointestinal Disorder
Headache
Helicobacter Infection
Hunger
Hypertension

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Laboratory Test Abnormal Muscle Spasms Nausea				Professional				
Nervousness Oesophageal Spasm				Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Pharmaceutical Product				Professional				
Complaint Pollakiuria					Lamotrigine (Lamotrigine)	SS		ORAL
Tracheal Disorder Weight Increased					Clonazepam (Clonazepam) Nifedipine (Nifedipine)	SS C		

Date:05/01/03ISR Number: 4106654-1Report Type:Expedited (15-DaCompany Report #2003006078
Age:19 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged				Literature Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other				Health Professional	Lamotrigine (Lamotrigine)	SS		ORAL
25 GRAM (ONCE), ORAL								
Hypertension Suicide Attempt Tachycardia								

Date:05/01/03ISR Number: 4106965-XReport Type:Periodic Company Report #DSA_80010_2002
Age:42 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Drug Effect Decreased				Health	Xyrem	PS		ORAL
Drug Interaction				Professional				
Inhibition					Neurontin	SS		

Pain

Date:05/02/03ISR Number: 4107150-8Report Type:Expedited (15-DaCompany Report #2002066682
Age:45 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Carpal Tunnel Syndrome	Consumer	Neurontin			
Initial or Prolonged	Cognitive Deterioration	Health	(Gabapentin)	PS		ORAL
400 MG						
Other	Condition Aggravated	Professional				
(DAILY), ORAL						
	Confusional State		Anxiolytics	SS		
	Disturbance In Attention		Tizanidine			
	Drug Interaction		Hydrochloride			
	Encephalopathy		(Tizanidine			
			Hydrochloride)	SS		
			Tramadol			
			Hydrochloride			
			(Tramadol			
			Hydrochloride)	SS		
			Alprazolam			
			(Alpazolam)	C		
			Oxycodone			
			Hydrochloride			
			(Oxycodone			
			Hydrochloride)	C		
			Hypnotics And			
			Sedatives	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/02/03ISR Number: 4107163-6Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 400 MG (BID), Disability ORAL Other	Brain Neoplasm Carotid Artery Disease Carotid Artery Stenosis Convulsion Dizziness Dysphemia Ependymoma Gastritis Gastrooesophageal Reflux Disease Memory Impairment Pain Somnolence Tremor Weight Decreased	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin) Buspirone Hydrochloride (Buspirone Hydrochloride)	PS C C C C		ORAL

Date:05/02/03ISR Number: 4107164-8Report Type:Expedited (15-DaCompany Report #2003011961

Age:57 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG (900 MG, BID), ORAL	Bronchiolitis Interstitial Lung Disease Pulmonary Fibrosis	Health Professional	Neurontin (Gabapentin) Bupropion Hydrochloride (Bupropion Hydrochloride) Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate) Fluticasone	PS C C		ORAL

Propionate
(Fluticasone
Propionate) C
Salbutamol
(Salbutamol) C

Date:05/02/03ISR Number: 4107166-1Report Type:Expedited (15-DaCompany Report #2003017389
Age:46 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 3600 MG Other (1200, TID), ORAL	Infection Lung Disorder Post Procedural Complication Pulmonary Oedema Surgical Procedure Repeated	Consumer	Neurontin (Gabapentin) Quetiapine Fumarate (Quetiapine Fumarate) Clonazepam (Clonazepam) Paroxetine Hydrochloride (Paroxetine	PS C C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C

Date:05/05/03ISR Number: 4107670-6Report Type:Expedited (15-DaCompany Report #2003150492US
 Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 200 MG, BID, Other ORAL	Accident Arthritis Eye Disorder	Consumer	Celebrex(Celecoxib) apsule	PS		ORAL
200 MG, BID, ORAL	Fibromyalgia Herpes Zoster		Neurontin(Gabapentin)	SS		ORAL
175 MG, ORAL	Intervertebral Disc Degeneration		Topamax(Topiramate)	SS		ORAL
EPIDURAL	EPIDURAL Intervertebral Disc Protrusion Nerve Compression Nerve Injury Neuralgia Neuropathy Peripheral Oedema Peripheral Post Procedural Complication Tremor		Unspecified Drug () Luvox Xanax Skelaxin (Metaxalone) Prozac (Fluoxetine Hydrochloride)	SS C C C C		

Date:05/05/03ISR Number: 4107985-1Report Type:Expedited (15-DaCompany Report #2003017659
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 400 MG (200, Other TWICE DAILY), ORAL	Condition Aggravated Diabetes Mellitus Poor Peripheral Circulation Surgery	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Ranitidine			

Throat Irritation
Urinary Tract Disorder

Hydrochloride
(Ranitidine
Hydrochloride) SS
Metformin
(Metformin) C
Tamsulosin
Hydrochloride
(Tamsulosin
Hydrochloride) C
Spironolactone
(Spironolactone) C
Potassium Chloride
(Potassium Chloride) C
Atenolol (Atenolol) C
Lansoprazole
(Lansoprazole) C

Date:05/05/03ISR Number: 4108173-5Report Type:Expedited (15-DaCompany Report #2003017240
Age:53 YR Gender:Female I/FU:F

Outcome PT
Other Arthritis
Convulsion
Drug Ineffective
Drug Withdrawal Syndrome
Exostosis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Intervertebral Disc Protrusion Movement Disorder	Report Source				
ORAL		Multiple Sclerosis Muscle Twitching	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Insulin Injection, Isophane (Insulin Injection, Isophane)	C		
				Insulin Lispro (Insulin Lispro)	C		
				Diclofenac (Diclofenac)	C		

Date:05/05/03ISR Number: 4108374-6Report Type:Expedited (15-DaCompany Report #2003157655FR
Age:51 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Alkaline Phosphatase Decreased Blood Bilirubin Increased	Foreign Health Professional	Medrol (Methylprednisolone) Tablet	PS		ORAL
SEE IMAGE		Diabetes Mellitus Inadequate Control Genital Candidiasis	Other	Glucophage (Metformin Hydrochloride)	SS		ORAL
850 MG, TID, ORAL		Hepatic Steatosis					
SUBCUTANEOUS	18UI/MORN, 10UI/EVENING, SUBCUTANEOUS	Hepatitis Cholestatic Oral Candidiasis		Insulin (Insulin)	SS		
1 DF, TID, ORAL				Diffu K (Potassium Chloride)	SS		ORAL
400 MG, TID				Neurontin (Gabapentin)	SS		

Date:05/06/03ISR Number: 4103780-8Report Type:Expedited (15-DaCompany Report #FR-BRISTOL-MYERS SQUIBB COMPANY-12261863
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Diabetes Mellitus	Health	Glucophage Tabs 850			
Hospitalization -		Inadequate Control	Professional	Mg	PS	Bristol-Myers Squibb	
Initial or Prolonged		Genital Candidiasis				Company	ORAL
		Glioblastoma		Mixtard	SS		
		Hepatic Steatosis		Diffu-K	SS		ORAL
		Hepatitis		Medrol	SS		ORAL
		Oral Candidiasis		Neurontin	SS		ORAL

Date:05/06/03ISR Number: 4108484-3Report Type:Expedited (15-DaCompany Report #2003005556
 Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		C-Reactive Protein	Foreign	Neurontin			
Initial or Prolonged		Increased	Health	(Gabapentin)	PS		ORAL
1200 MG							
(TID), ORAL		Dermatitis Allergic	Professional				
		Erythema Multiforme	Company	Morphine (Morphine)	C		
		Oedema Peripheral	Representative	Diclofenac			
		Pain		(Diclofenac)	C		
				Amitriptyline			
				(Amitriptyline)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/06/03ISR Number: 4108946-9Report Type:Expedited (15-DaCompany Report #2003018309

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Blood Urine Joint Swelling					
THREE TIMES DAILY, ORAL		Mental Status Changes					
		Penile Haemorrhage Pyrexia		Valaciclovir Hydrochloride Diuretics Warfarin Sodium	SS C C		

Date:05/06/03ISR Number: 4109051-8Report Type:Expedited (15-DaCompany Report #2003018024

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional	Neurontin (Gabapentin)	PS		
Other		Drug Withdrawal Syndrome Headache					
UNKNOWN	2400 MG	Treatment Noncompliance					
UNKNOWN							

Date:05/07/03ISR Number: 4107571-3Report Type:Direct Company Report #CTU 192468

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Hydromorphone Gabapentin	PS SS		
		Mental Status Changes					

Date:05/07/03ISR Number: 4109100-7Report Type:Expedited (15-DaCompany Report #055-0945-M0100042

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Asthenia	Foreign	Gabapentin			

Other	Back Pain	Consumer	(Gabapentin)	PS	ORAL
2000 MG (400	Blood Cholesterol				
MG, 5X/DAY),	Increased				
ORAL	Blood Triglycerides		Amitriptyline		
	Increased		Hydrochloride		
	Bone Neoplasm		(Amitriptyline		
	Bone Pain		Hydrochloride)	C	
	Chest Pain		Diazepam (Diazepam)	C	
	Condition Aggravated		Sertraline		
	Constipation		(Sertraline)	C	
	Depression		Venlafaxine		
	Dizziness		(Venlafaxine)	C	
	Fall		Alprazolam		
	Fibromyalgia		(Alprazolam)	C	
	Insomnia		Acetylsalicylic Acid		
	Neck Pain		(Acetylsalicylic		
	Neuropathic Pain		Acid)	C	
	Oedema				
	Pain In Extremity				
	Paraesthesia				
	Peripheral Coldness				
	Suicide Attempt				
	Treatment Noncompliance				
	Weight Increased				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/07/03ISR Number: 4109112-3Report Type:Expedited (15-DaCompany Report #2003018132
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Condition Aggravated	Foreign	Neurontin			
Hospitalization -		Cushingoid	Health	(Gabapentin)	PS		ORAL
1200 MG							
Initial or Prolonged		Diabetes Mellitus	Professional				
(TID), ORAL							
		Inadequate Control		Initard (Insulin,			
		Difficulty In Walking		Insulin Injection,			
		Genital Candidiasis		Isophane)	SS		
SUBCUTANEOUS	28 I.U.						
(TWICE		Hepatic Steatosis					
DAILY),		Oral Candidiasis					
SUBCUTANEOUS				Potassium Chloride			
(TID), ORAL				(Potassium Chloride)	SS		ORAL
(FIVE TIMES				Methylprednisolone			
PER DAY),				(Methylprednisolone)	SS		ORAL
ORAL							
				Metformin			
				Hydrochloride			
				(Metformin			
				Hydrochloride)	C		ORAL
2550 MG							
(TID), ORAL							

Date:05/07/03ISR Number: 4109135-4Report Type:Expedited (15-DaCompany Report #20020685598
 Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Asthma	Foreign	Gabapentin (Tablets)			

Initial or Prolonged 1200 MG	Bronchitis Acute	Study	(Gabapentin)	PS	ORAL
(THREE TIMES DAILY), ORAL	Condition Aggravated	Health			
	Cough	Professional			
	Drug Interaction		Carbamazepine		
	Dyspnoea		(Carbamazepine)	C	
	Inflammation		Clobazam (Clobazam)	C	
	Nasopharyngitis		Theophylline		
	Pain		(Theophylline)	C	
	Pyrexia		Pranlukast		
			(Pranlukast)	C	
			Salbutamol		
			(Salbutamol)	C	
			Budesonide		
			(Budesonide)	C	
			Tulobuterol		
			Hydrochloride		
			(Tulobuterol		
			Hydrochloride)	C	

Date:05/07/03ISR Number: 4109381-XReport Type:Expedited (15-DaCompany Report #2003018128
Age: Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Blood Urine
Initial or Prolonged	Condition Aggravated
Other	Coronary Artery
	Reocclusion
	Coronary Artery Surgery
	Drug Effect Decreased
	Lymphatic Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Complaint	Report Source	Product	Role	Manufacturer	Route
5400 MG		Lymphoedema Malabsorption Meningioma Muscle Disorder Pharmaceutical Product	Consumer	Neurontin (Gabapentin)	PS		ORAL
(TID), ORAL		Renal Cell Carcinoma Stage Unspecified Surgical Procedure Repeated Transitional Cell Carcinoma		Candesartan Cilexetil (Candesartan Cilexetil) Furosemide (Furosemide) Rofecoxib (Rofecoxib) Potassium Chloride (Potassium Chloride) Combivent (Ipratropium Bromide, Salbutamol Sulfate) Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate) Beclometasone Dipropionate (Beclometasone Dipropionate) Meclozine (Meclozine) Esomeprazole (Esomeprazole) Alprazolam (Alprazolam)	C C C C C C C		

Date:05/07/03ISR Number: 4109395-XReport Type:Expedited (15-DaCompany Report #2003018584
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Alopecia	Consumer	Neurontin			

300 MG (BID),	Balance Disorder	(Gabapentin)	PS	ORAL
ORAL	Condition Aggravated			
	Episcleritis	Clonazepam		
	Herpes Zoster	(Clonazepam)	SS	
	Muscle Twitching	Alprazolam		
	Myalgia	(Alprazolam)	SS	
	Paraesthesia	Prednisolone Acetate		
	Somnolence	(Prednisolone		
	Vision Blurred	Acetate)	SS	
		Ascorbic Acid		
		(Ascorbic Acid)	C	
		Esomeprazole		
		(Esomeprazole)	C	
		Montelukast Sodium		
		(Montelukast Sodium)	C	
		Fluticasone		
		Propionate		
		(Fluticasone		
		Propionate)	C	

Freedom Of Information (FOI) Report

Salmeterol Xinafoate
 (Salmeterol
 Xinafoate) C
 Salbutamol
 (Salbutamol) C
 Progesterone
 (Progesterone) C
 Hydrocortisone
 (Hydrocortisone) C
 All Other
 Therapeutic Products C

Date:05/07/03ISR Number: 4109552-2Report Type:Expedited (15-DaCompany Report #2003018310
 Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (DAILY), ORAL		Adhesion Facial Nerve Disorder	Foreign Consumer	Gabapentin (Tablets) (Gabapentin)	PS		ORAL
				All Other Therapeutic Products	C		

Date:05/08/03ISR Number: 4109172-XReport Type:Expedited (15-DaCompany Report #2003157842US
 Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 20 MG, QD, ORAL		Asthenia Difficulty In Walking	Health Professional	Bextra (Valdecoxib)	PS		ORAL
300 MG, QD, ORAL		Muscular Weakness Oliguria		Neurontin (Gabapentin)	SS		ORAL
		Tremor					
		Urine Output Decreased		Lortab (Paracetamol, Hydrocodone			

ORAL

Bitartrate) Tablet	SS
Lexapro	C
Norvasc (Amlodipine Besilate)	C
Aspirine	C
Vitamins	C

ORAL

Date:05/09/03ISR Number: 4110679-XReport Type:Expedited (15-DaCompany Report #2003009356

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Constipation	Consumer	Neurontin			
800 MG (400		Diarrhoea	Health	(Gabapentin)	PS		ORAL
MG, BID);		Fibroadenoma Of Breast	Professional				
ORAL (SEE		Flatulence					
IMAGE)		Gastrointestinal Disorder					
		Insomnia		Fluoxetine			
		Pharyngolaryngeal Pain		Hydrochloride	C		
		Stress					
		Suicidal Ideation					
		Weight Increased					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/09/03ISR Number: 4110689-2Report Type:Expedited (15-DaCompany Report #2003018325
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Infertility Male	Health Professional	Neurontin (Gabapentin)	PS		ORAL
UNKNOWN							

(TID); ORAL

Rofecoxib	C
Zolpidem Tartrate	C
Vicodin (Paracetamol, Hydrocodone Bitartrate)	C
Omeprazole	C
Cyclobenzaprine Hydrochloride	C
Salbutamol	C

Date:05/12/03ISR Number: 4111108-2Report Type:Expedited (15-DaCompany Report #2003018755
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthralgia	Consumer	Neurontin (Gabapentin)	PS		
Other		Brain Damage					
2400 MG (600, FOUR TIMES A DAY)		Cerebral Artery Occlusion					
		Cerebrovascular Accident					
		Convulsion		Anovlar (Norethisterone Acetate, Ethinylestradiol)	SS		
		Difficulty In Walking					
		Emotional Distress					
		Eye Pain					
		Hemiparesis					
		Migraine					
		Oral Pain					
		Pain					
		Peroneal Nerve Palsy					

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Anaemia	Consumer	Neurontin			
600 MG (BID)	Asthma		(Gabapentin)	PS		ORAL
Other	Condition Aggravated					
ORAL	Cystitis		Prednisone			
	Diabetes Mellitus		(Prednisone)	SS		
	Diabetic Neuropathy		Acetylsalicylic Acid	C		
	Essential Tremor		Insulin Lispro	C		
	Fall		All Other			
	Gilbert'S Syndrome		Therapeutic Products	C		
	Haemolytic Anaemia		Simvastatin	C		
	Headache		Seretide Mite			
	Infection		(Fluticasone			
	Kidney Infection		Propionate,			
	Malaise		Salmeterol			
	Pneumonia		Xinafoate)	C		
	Shoulder Operation		Furosemide	C		
	Urinary Tract Infection		Combivent			
	Weight Decreased		(Ipratropium			
			Bromide, Salbutamol			
			Sulfate)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Propacet
 (Paracetamol,
 Dextropropoxyphene
 Napsilate) C
 Digoxin C

Date:05/12/03ISR Number: 4111394-9Report Type:Expedited (15-DaCompany Report #2003005197
 Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Activities Of Daily	Consumer	Neurontin			
Other		Living Impaired		(Gabapentin)	PS		ORAL
900 MG (300		Arthralgia					
TID) ORAL		Back Pain		Norvasc (Amlodipine)	SS		ORAL
ORAL		Cholelithiasis		Meloxicam	C		
		Diarrhoea		Atenolol	C		
		Difficulty In Walking		Vitamins	C		
		Drug Ineffective					
		Dysstasia					
		Joint Lock					
		Joint Swelling					
		Musculoskeletal Stiffness					
		Sciatica					
		Skin Discolouration					
		Sleep Attacks					

Date:05/12/03ISR Number: 4111407-4Report Type:Expedited (15-DaCompany Report #2003018320
 Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Agranulocytosis	Foreign	Neurontin (Tablets)			
Initial or Prolonged		Bone Marrow Depression	Health	(Gabapentin)	PS		ORAL
900 MG		Staphylococcal Infection	Professional				
Other				Acetylsalicylic Acid			
(DAILY), ORAL				(Acetylsalicylic			
				Acid)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Deafness Neurosensory	Foreign	Neurontin			
Other		Tremor	Consumer	(Gabapentin)	PS		ORAL
1800 MG (600		Vaginal Discharge	Health				
, TID), ORAL		Vaginal Haemorrhage	Professional	Calcium Carbonate (Calcium Carbdate)	C		
				Prednisolone (Prednisolone)	C		
				Verapamil Hydrochloride (Verapamil			
				Hydrochloride)	C		
				Acetylsalicylic Acid (Acetylsalicylic			
				Acid)	C		
				Triazolam (Triazolam)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/12/03ISR Number: 4111420-7Report Type:Expedited (15-DaCompany Report #2003018752
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depressed Level Of Consciousness Hepatocellular Damage	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:05/12/03ISR Number: 4111908-9Report Type:Direct Company Report #USP 51747
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Neurontin	PS	Pd	

Date:05/12/03ISR Number: 4112203-4Report Type:Direct Company Report #USP 51667
 Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Neurontin	PS	Parke Davis	

Date:05/12/03ISR Number: 4112697-4Report Type:Direct Company Report #CTU 192772
 Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Speech Disorder		Gabapentin	PS		

Date:05/13/03ISR Number: 4108379-5Report Type:Expedited (15-DaCompany Report #IT-GLAXOSMITHKLINE-B0273503A
 Age:64 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening 1TAB Twice		Acute Myocardial	Health	Combivir	PS	Glaxosmithkline	ORAL

Hospitalization - per day	Infarction	Professional			
Initial or Prolonged 600MG Per day	Cardiac Arrest		Efavirenz	SS	ORAL
			Salmeterol	SS	Glaxosmithkline
			Gabapentin	SS	
			Amiodarone	C	ORAL
			Carvedilol	C	Glaxosmithkline
			Ramipril	C	
			Furosemide	C	Glaxosmithkline

Date:05/13/03ISR Number: 4111936-3Report Type:Expedited (15-DaCompany Report #ERD2003A00073
Age:70 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Anaemia
Initial or Prolonged	Blood Immunoglobulin G
	Increased
	C-Reactive Protein
	Increased
	Cardiomegaly
	Epstein-Barr Virus
	Antibody Positive
	Hepatotoxicity
	Liver Function Test
	Abnormal

Freedom Of Information (FOI) Report

Ventricular Dysfunction

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
45 MG ORAL	368 DAY	Study Health Professional	Pioglitazone Hydrochloride (Placebo) (Pioglitazone Hydrochloride)	PS		ORAL
257 DAY			Gabapentin (Gabapentin)	SS		
3 YR			Simvastatin (Simvastatin)	SS		
5 YR			Allopurinol (Allopurinol)	SS		
			Metformin (Metformin)	C		
			Insuline (Insulin)	C		
			Thiamazol (Thiamazole)	C		
			Furosemid (Furosemide)	C		
			Nitro Glycerine (Glyceryl Trinitrate)	C		
			Clopidogrel (Clopidogrel)	C		
			Fosinopril (Fosinopril)	C		
			Folic Acid (Folic Acid)	C		
			Alpha-Lipoic Acid (Thioctic Acid)	C		

Date:05/13/03ISR Number: 4111982-XReport Type:Expedited (15-DaCompany Report #2003010680
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Condition Aggravated	Foreign	Neurontin			

Other 900 MG (DAILY), ORAL	Hyperkalaemia Hypokalaemia Sympathetic Nerve Destruction	Health Professional	(Gabapentin)	PS	ORAL
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Date:05/14/03ISR Number: 4112399-4Report Type:Expedited (15-DaCompany Report #2003019418
Age: Gender: I/FU:I

Outcome Dose Duration Hospitalization - Initial or Prolonged Other THREE TIMES DAILY), ORAL	PT	Report Source	Product	Role	Manufacturer	Route
	Abdominal Pain Burning Sensation Cold Sweat Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
	Joint Swelling Parkinson'S Disease		Citalopram Hydrobromide (Citalopram Hydrobromide) Famotidine (Famotidine) Multivitamins (Ergocalciferol,	C C		

Freedom Of Information (FOI) Report

Ascorbic Acid, Folic
 Acid, Thiamine
 Hydrochloride, C
 Paracetamol
 (Paracetamol) C

Date:05/14/03ISR Number: 4112400-8Report Type:Expedited (15-DaCompany Report #2003019414
 Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Interaction Irritable Bowel Syndrome	Health Professional	Neurontin (Gabapentin)	PS		ORAL
2700 MG (TID), ORAL		Jaw Disorder					
		Muscle Spasms Muscle Twitching		Mirtazapine (Mirtazapine)	SS		ORAL
2700 MG (TID), ORAL		Speech Disorder					
				Hydroxychloroquine Phosphate (Hydroxychloroquine Phosphate)	C		
				Celecoxib	C		
				Prednisone (Prednisone)	C		
				Multivitamins And Iron	C		
				Calcium (Calcium)	C		
				Cilest (Ethinylestradiol, Norgestimate)	C		
				Zolpidem Tartrate (Zolpidem Tartrate)	C		
				Nifedipine (Nifedipine)	C		
				Glyceryl Trinitrate (Glyceryl Trinitrate)	C		
				Rabeprazole Sodium (Rabeprazole Sodium)	C		
				Etanercept (Etanercept)	C		
				Tramadol			

Hydrochloride	
(Tramadol	
Hydrochloride)	C
Fish Oil (Fish Oil)	C
Chondroitin/Glucosamine (Glucosamine, Chondroitin)	C
Promethazine	
(Promethazine)	C
Prochlorperazine Edisylate	
(Prochlorperazine Edisylate)	C
Alprazolam	
(Alprazolam)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/15/03ISR Number: 4112864-XReport Type:Expedited (15-DaCompany Report #2003019626

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 20 MG Other (DAILY), ORAL	Arteriosclerosis Blood Cholesterol Increased Blood Triglycerides Drug Effect Decreased Myocardial Infarction Pain In Extremity	Consumer	Lipitor (Atorvastatin)	PS		ORAL
			Gabapentin (Gabapentin)	SS		
			Fluvastatin Sodium(Fluvastatin Sodium)	SS		
			Ezetimibe (Ezetimibe)	SS		
			Benazepril Hydrochloride (Benazepril Hydrochloride)	C		
			Atenolol (Atenolol)	C		
			Hyzaar (Hydrochlorothiazide , Losartan Potassium)	C		
			Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
			Alprazolam(Alprazola m)	C		

Date:05/15/03ISR Number: 4112875-4Report Type:Expedited (15-DaCompany Report #2003011978

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL	Abnormal Behaviour Anger	Consumer	Neurontin (Gabapentin)	PS		ORAL
	Anxiety Dependence Euphoric Mood Irritability Thinking Abnormal		Bupropion Hydrochloride (Bupropion Hydrochloride)	C		
			Doxycycline			

(Doxycycline)

C

Date:05/16/03ISR Number: 4112441-0Report Type:Expedited (15-DaCompany Report #2003019581

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased	Health	Neurontin			
		Inguinal Hernia	Professional	(Gabapentin)	PS		ORAL
3600 MG							
(1200, THREE		Quadriplegia	Company				
TIMES DAILY)		Renal Disorder	Representative				
ORAL							

Baclofen(Baclofen)

C

Date:05/16/03ISR Number: 4112658-5Report Type:Expedited (15-DaCompany Report #2003019420

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser	Consumer	Neurontin			
		Unevaluable Event		(Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/16/03ISR Number: 4112712-8Report Type:Expedited (15-DaCompany Report #2003157842US

Age:71 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Constipation Difficulty In Walking	Health Professional	Bextra (Valdecoxib) Tablet	PS		ORAL
20 MG, QD, ORAL		Dizziness					
		Dysarthria Muscular Weakness		Neurontin (Gabapentin)	SS		ORAL
300 MG, QD, ORAL		Oliguria					
		Tremor		Lortab (Paracetamol, Hydrocodone Bitartrate) Tablet	SS		ORAL
ORAL				Lexapro Norvasc (Amlodipine Besilate) Aspirine Vitamins	C C C C		

Date:05/16/03ISR Number: 4113548-4Report Type:Expedited (15-DaCompany Report #2003019549

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Disease Recurrence Dyspepsia	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (BID), Other ORAL		Fatigue					
		Nausea Thinking Abnormal		Carbamazepine (Carbamazepine)	SS		ORAL
ORAL		Trigeminal Neuralgia Vomiting		Fentanyl (Fentanyl) All Other Therapeutic Products	C C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL Other	Cerebral Atrophy Cerebrovascular Accident Coma Confusional State Dyskinesia Extrapyramidal Disorder Loss Of Consciousness Restlessness Somnolence Urinary Tract Infection	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Olanzapine (Olanzapine) Tetrabenazine (Tetrabenazine) Levothyroxine Sodium (Levothyroxine Sodium) Torasemide (Torasemide) Insulin Human Injection, Isophane (Insulin Human Injection, Isophane) Lactulose (Lactulose)	PS C C C C C C		ORAL

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 - Initial or Prolonged	Mental Status Changes		Duragesic	PS		
			Vicodin	SS		
			Cardura	SS		
			Wellbutrin	SS		
			Neurontin	SS		
			Tequin	SS		

Date:05/19/03ISR Number: 4114349-3Report Type:Direct
Age:49 YR Gender:Male I/FU:I

Company Report #CTU 193373

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening 300MG PO BID	Haemodialysis		Neurontin	PS		ORAL
Hospitalization - Initial or Prolonged Required	Renal Failure Acute Sleep Disorder		Klonopin	C		
Intervention to Prevent Permanent Impairment/Damage			Effexor	C		
			Vicodin	C		
			Cocaine	C		

Date:05/19/03ISR Number: 4114926-XReport Type:Expedited (15-DaCompany Report #2003019777
Age:14 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Chest Pain Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL
6400 MG TID, ORAL	Hypoaesthesia Rickets		Paracetamol (Paracetamol)	C		

Date:05/19/03ISR Number: 4114928-3Report Type:Expedited (15-DaCompany Report #2003019790
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	600 MG BID,	Depression Dialysis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other ORAL		Dysuria Renal Failure Acute		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		

Date:05/19/03ISR Number: 4114932-5Report Type:Expedited (15-DaCompany Report #2003009221
Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG	Condition Aggravated Grand Mal Convulsion	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other (DAILY), ORAL		Visual Disturbance	Professional	Didanosine (Didanosine)	C		
				Stavudine (Stavudine)	C		
				Saquinavir (Saquinavir)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Kaletra (Ritonavir, Lopinavir)	C
Bactrim (Sulfamethoxazole, Trimethoprim)	C
Azithromycin (Azithromycin)	C
Fluconazole (Fluconazole)	C
Levothyroxine Sodium (Levothyroxine Sodium)	C
Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C
Diazepam (Diazepam)	C
Oxycodone (Oxycodone)	C
Dronabinol (Dronabinol)	C
Nystatin (Nystatin)	C
Benzonatate (Benzonatate)	C
Phenytoin Sodium (Phenytoin Sodium)	C
Cetirizine Hydrochloride (Cetirizine Hydrochloride)	C
Celecoxib (Celecoxib)	C

Date:05/19/03ISR Number: 4115295-1Report Type:Expedited (15-DaCompany Report #2003UW06025
Age:86 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 20 MG DAILY		Blood Creatinine	Foreign	Elavil	PS		ORAL
Initial or Prolonged PO		Increased	Health				
Required 1800 MG DAILY		Blood Pressure Decreased	Professional	Neurontin	SS		ORAL

Intervention to	Fatigue	Other			
PO					
Prevent Permanent 1200 MG DAILY Impairment/Damage PO	Haematuria		Geavir	SS	ORAL
	Pyrexia				
	Somnolence		Tramadol Hydrochloride	SS	ORAL
200 MG DAILY					
PO			Alvedon	C	
			Lanacrist	C	
			Trombyl	C	

Date:05/19/03ISR Number: 4115345-2Report Type:Expedited (15-DaCompany Report #2003019901
Age:69 YR Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Arthritis
Hospitalization -	Asthenia
Initial or Prolonged	Blood Creatine
Other	Phosphokinase Increased
	Blood Lactate

FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
800 MG (BID), ORAL		Dehydrogenase Increased Leukopenia Mucosal Inflammation Rash Erythematous	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Renal Failure Renal Impairment Swelling		Atorvastatin (Atorvastatin) Gliclazide (Gliclazide) Irbesartan (Irbesartan) Metformin Hydrochloride (Metformin Hydrochloride)	C C C C		

Date:05/19/03ISR Number: 4115349-XReport Type:Expedited (15-DaCompany Report #2003019548
Age: Gender:Female I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Fall Hypoaesthesia Sensory Loss	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:05/20/03ISR Number: 4114267-0Report Type:Expedited (15-DaCompany Report #2003019956
Age:74 YR Gender:Male I/FU:I

Outcome Dose Hospitalization - Initial or Prolonged ORAL Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Oedema Peripheral Venous Pressure Jugular	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Increased		Nitrazepam (Nitrezepam) Quinine (Quinine) Panadeine Co	C C		

(Codeine Phosphate,
Paracetamol) C

Date:05/20/03ISR Number: 4114351-1Report Type:Expedited (15-DaCompany Report #2003017145
Age:76 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Blood Creatinine	Foreign	Neurontin (Tablets)			
Initial or Prolonged	Blood Potassium Increased	Health	(Gabapentin)	PS		ORAL
800 MG (BID),						
ORAL	C-Reactive Protein	Professional				
	Increased	Company	Lansoprazole			
	Cholestasis	Representative	(Lansoprazole)	C		
	Haemoglobin Decreased		Lercanidipine			
	Hepatitis		Hydrochloride			
	Hepatocellular Damage		(Lercanidipine			
	Platelet Count		Hydrochloride)	C		
	Red Blood Cell					
	Sedimentation Rate					
	Increased					
	Urinary Tract Infection					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/20/03ISR Number: 4114565-0Report Type:Expedited (15-DaCompany Report #2003020204

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Colorectal Cancer	Foreign	Neurontin			
ORAL		Constipation	Consumer	(Gabapentin)	PS		ORAL
		Drug Ineffective		Morphine (Morphine)	SS		ORAL
ORAL		Hepatic Neoplasm		All Other			
		Malignant		Therapeutic Products	SS		
		Pain		Vitamins	C		
		Recurrent Cancer					

Date:05/21/03ISR Number: 4113016-XReport Type:Expedited (15-DaCompany Report #GB-GLAXOSMITHKLINE-B0299906A

Age:46 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Maculopathy		Lamotrigine	PS	Glaxosmithkline	ORAL
200MG Twice		Visual Acuity Reduced					
per day				Topiramate	SS		
UNKNOWN	100MG Twice						
per day				Gabapentin	SS		
UNKNOWN	900MG Twice						
per day				Paracetamol	C	Glaxosmithkline	
UNKNOWN							

Date:05/21/03ISR Number: 4115541-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Amnesia	Consumer	Neurontin			
Initial or Prolonged		Aspartate	Health	(Gabapentin)	PS		ORAL
400 MG (BID),							

Disability	Aminotransferase	Professional		
ORAL				
Other	Increased	Company	Valproate Sodium	
	Brain Neoplasm Benign	Representative	(Valproate Sodium)	SS
	Cardiac Arrest		Fluoxetine	
	Convulsion		Hydrochloride	
	Diplopia		(Fluoxetine	
	Fatigue		Hydrochloride)	C
	Gallbladder Disorder		Omeprazole	
	Headache		(Omeprazole)	C
	Incoherent		Capsaicin	
	Mood Swings		(Capsaicin)	C
	Muscle Twitching		Buspirone	
	Somnolence		Hydrochloride	
	Tardive Dyskinesia		(Buspirone	
	Tinnitus		Hydrochloride)	C

Date:05/21/03ISR Number: 4115806-6Report Type:Direct Company Report #CTU 87209
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medication Error		Noroxin 400mg Roberts Pharmaceuticals	PS	Roberts Pharmaceuticals	ORAL
400MG 3WKS PO				Neurontin 400mg Parke-Davis	SS	Parke-Davis	ORAL
400MG 3/WK PO							

[NEVER-ERROR]

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/21/03ISR Number: 4115881-9Report Type:Expedited (15-DaCompany Report #2003020214
Age:49 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL Other	Condition Aggravated Depression	Health Professional	Neurontin (Gabapentin)	PS		ORAL
	Dialysis Renal Failure Acute Renal Tubular Necrosis		Cocaine (Cocaine) Vicodin (Paracetamol, Hydrocodone Bitartrate)	C C		

Date:05/21/03ISR Number: 4116072-8Report Type:Expedited (15-DaCompany Report #2003019549
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG (BID), Other ORAL	Dental Necrosis Disease Recurrence	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
	Dry Mouth					
	Dyspepsia Fatigue		Carbamazepine (Carbamazepine)	SS		ORAL
	Mental Impairment Nausea Pharmaceutical Product Complaint Treatment Noncompliance Trigeminal Neuralgia Vomiting		Fentanyl (Fentanyl) All Other Therapeutic Products	C C		

Date:05/22/03ISR Number: 4116254-5Report Type:Expedited (15-DaCompany Report #2003020659
Age:45 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL	Anxiety Chills	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL

Condition Aggravated
Feeling Cold
Salivary Hypersecretion

Amitriptyline
(Amitriptyline) SS
Fluoxetine
Hydrochloride
(Fluoxetine
Hydrochloride) SS
Dihydrocodeine
(Dihydrocodeine) C
Morphine (Morphine) C

Date:05/22/03ISR Number: 4116491-XReport Type:Expedited (15-DaCompany Report #2003012704

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Effect Decreased Eating Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300, THREE TIMES DAILY) ORAL		Hypoaesthesia Hypotonia Medication Error Nerve Injury Oral Discomfort Oral Dysaesthesia Paraesthesia Road Traffic Accident		Acetylaslicylic Acid Ibuprofen Allegra-D (Pseudoephedrine Hydrochloride Fexofenadine)	C C C C		

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FDA - Adverse Event Reporting System (AERS)

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Mometasone Furoate C
 Cyclobenzaprine
 Hydrochloride C
 Esomeprazole C
 Iron C
 Vicodin
 (Paracetamol,
 Hydrocodone
 Bitartrate) C

Date:05/22/03ISR Number: 4116564-1Report Type:Expedited (15-DaCompany Report #2003020459
 Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Coagulation Time Prolonged	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (BID), ORAL		General Physical Health	Professional				
10 MG (DAILY), ORAL		Deterioration Hepatocellular Damage		Norvasc (Amlodipine)	C		ORAL
1200 MG (QID), ORAL		Jaundice Liver Function Test Abnormal		Carbamazepine(Carbamazepine)	C		ORAL
				Atenolol (Atenolol) Enalapril Maleate (Enalapril Maleate)	C C		

Date:05/22/03ISR Number: 4116628-2Report Type:Expedited (15-DaCompany Report #EMADSS2003004079
 Age:46 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Maculopathy Visual Acuity Reduced	Foreign Health	Topamax(Unspecified) (Topiramate)	PS		
100 MG, 2 IN 1 DAILY,			Professional				

900 MG, 2 IN	Gabapentin (Gabapentin)	SS	
1 DAILY			
400 MG.	Carbamazepine (Carbamazepine)	SS	ORAL
DAILY, ORAL			
200 MG, 2 IN	Lamotrigine (Lamotrigine)	SS	ORAL
1 DAILY, ORAL			
	Paracetamol (Paracetamol)	C	

Date:05/23/03ISR Number: 4116949-3Report Type:Expedited (15-DaCompany Report #2003016992
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Hepatic Failure	Foreign	Gabapentin			
Hospitalization -	Liver Transplant	Health	(Gabapentin)	PS		
Initial or Prolonged		Professional				
Other						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/23/03ISR Number: 4116957-2Report Type:Expedited (15-DaCompany Report #2003021076
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	900 MG (300MG TID),	Somnolence Weight Increased	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:05/23/03ISR Number: 4116997-3Report Type:Expedited (15-DaCompany Report #2003021210
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Disability		Dyskinesia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:05/23/03ISR Number: 4117133-XReport Type:Expedited (15-DaCompany Report #2003020570
 Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100 MG (DAILY), ORAL		Diplopia Eye Disorder Medication Error Swelling Vision Blurred	Consumer	Neurontin (Gabapentin) Diltiazem Hydrochloride (Diltiazem Hydrochloride) Celecoxib (Celecoxib) Cetirizine Hydrochloride	PS C C		ORAL

(Cetirizine
Hydrochloride) C
Clopidogrel Sulfate
(Clopidogrel
Sulfate) C

Date:05/23/03ISR Number: 4117136-5Report Type:Expedited (15-DaCompany Report #2003020574
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased Intestinal Operation Malabsorption Pharmaceutical Product Complaint	Consumer	Neurontin (Gabapentin) All Other Therapeutic Products	PS C		

Date:05/23/03ISR Number: 4117139-0Report Type:Expedited (15-DaCompany Report #2003020630
Age: Gender:Male I/FU:I

Outcome	PT
Other	Brain Damage Cataract Condition Aggravated

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Freedom Of Information (FOI) Report

Dose	Duration	Neuropathic Pain Retinopathy Transient Ischaemic Attack Treatment Noncompliance Vascular Occlusion	Report Source	Product	Role	Manufacturer	Route
600 MG (300, BID), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Amlodipine Besilate (Amlodipine Besilate)	C		
				Insulin Human (Insulin Human)	C		
				Simvastatin (Simvastatin)	C		
				Asasantin (Acetylsalicylic Acid, Dipyridamole)	C		
				Potassium (Potassium)	C		
				Furosemide (Furosemide)	C		
				Folic Acid (Folic Acid)	C		
				Pramipexole (Pramipexole)	C		
				Alprazolam (Alprazolam)	C		
				Cyanocobalamin (Cyanocobalamin)	C		
				Iron (Iron)	C		
				Insulin Human Injection, Isophane (Insulin Human Injection, Isophane)	C		

Date:05/23/03ISR Number: 4117157-2Report Type:Expedited (15-DaCompany Report #2003020736

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Abdominal Pain Upper Anxiety	Consumer	Neurontin (Gabapentin)	PS		ORAL

Dysstasia
Exostosis
Fatigue
Malaise
Nausea
Pain
Visual Disturbance

Date:05/23/03ISR Number: 4117162-6Report Type:Expedited (15-DaCompany Report #2003017659

Age: Gender:Male I/FU:F

Outcome PT
Hospitalization - Condition Aggravated
Initial or Prolonged Diabetes Mellitus
Other Double Vessel Bypass
Graft
Micturition Disorder
Poor Peripheral

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Circulation
Throat Irritation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
400 MG (200, TWICE DAILY), ORAL		Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Ranitidine Hydrochloride (Ranitidine Hydrochloride)	SS		
			Metformin (Metformin)	C		
			Tamsulosin Hydrochloride (Tamsulosin Hydrochloride)	C		
			Spiroinolactone (Spiroinolactone)	C		
			Potassium Chloride (Potassium Chloride)	C		
			Atenolol (Atenolol)	C		
			Lansoprazole (Lansoprazole)	C		

Date:05/23/03ISR Number: 4117166-3Report Type:Expedited (15-DaCompany Report #2003015764
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG (600, Other QID)		Atrioventricular Block Intestinal Obstruction Treatment Noncompliance	Consumer Health Professional	Neurontin (Gabapentin)	PS		
				All Other Therapeutic Products	C		

Outcome	PT
Death	Alopecia
Hospitalization -	Amnesia
Initial or Prolonged	Angiopathy
	Anxiety
	Arterial Occlusive
	Disease
	Arthralgia
	Asthma
	Atherosclerosis
	Back Pain
	Carpal Tunnel Syndrome
	Chest Pain
	Coma
	Condition Aggravated
	Depression
	Disturbance In Attention
	Drug Ineffective
	Emphysema
	Fatigue
	Femoral Bruit

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Finger Deformity Flushing Hilar Lymphadenopathy				
		Hypoaesthesia Influenza Influenza Like Illness Insomnia	Consumer Health Professional Other			
10 MG,			Oxycontin Tablets 10 Mg(Oxycodone Hydrochloride) Cr Tablet			PS
		Irritability Libido Decreased Memory Impairment Myalgia				
20 MG,	1008 DAY		Oxycontin Tablets 20 Mg(Oxycodone Hydrochloride) Cr Tablet			SS
		Nausea Nervousness Night Sweats Overdose Pain	Hydrocodone Bitartrate (Similar To Ind 59,175) (Hydrocodone Bitartrate) Unknown			SS
1008 DAY						
		Pain In Extremity Paraesthesia	Alprazolam (Alprazolam)			SS
1008 DAY						
		Peripheral Occlusive Disease	Ephedrine (Ephedrine)			SS
1008 DAY						
		Pneumonia Pulmonary Oedema	Pseudoephedrine (Pseudoephedrine)			SS
1008 DAY						
		Pulse Absent Rash Pruritic	Effexor(Venlafaxine Hydrochloride)			SS
75 MG, DAILY	1008 DAY					
		Shock	Codeine (Codeine)			SS
1008 DAY						
		Somnolence Toxicologic Test Abnormal	Neurontin(Gabapentin)			SS
100 MG, TID,	1008 DAY					
		Upper Respiratory Tract Infection	Metoclopramide(Metoc lopramide)			SS
1008 DAY						
		Urinary Incontinence Weight Increased	Quetiapine(Quetiapin e)			SS
1008 DAY						
			Claritin (Loratadine) Vitamin C (Ascorbic Acid)			C C

Celebrex (Celecoxib)	C
Prilosec	
(Omeprazole)	C
Daypro (Oxaprozin)	C
Trandate (Labetalol	
Hydrochloride)	C
Axid (Nizatidine)	C
Paxil (Paroxetine	
Hydrochloride)	C
Medrol	
(Methylprednisolone)	C
Macro Antioxidant	
(Ascorbic Acid,	
Cystine, Tocopherol,	
Calcium Ascorbate,	
Betacarotene,	C
Zocor (Simvastatin)	C
Risperdal	
(Risperidone)	C
Fioricet	
(Butalbital)	C
Flonase (Fluticasone	
Propionate)	C
Phenobarbital	
(Phenobarbital)	C
Donnatal (Atropine	

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Sulfate, Hyoscine
 Hydrobromide,
 Hyoscyamine Sulfate,
 Phenobarbital) C
 Bellergal-S
 (Belladonna
 Alkaloids,
 Ergotamine Tartrate,
 Phenobarbital) C
 Weight Loss
 Supplement (Does Not
 Code) C
 Ibuprofen C
 Ambien (Zolpidem
 Tartrate) C
 Baclofen (Baclofen) C
 Trandate (Labetatol
 Hydrochloride) C
 Celexa (Citalopram
 Hydrbromide) C
 Cortisone
 (Cortisone) C
 Atarax (Hydroxyzine
 Hydrochloride) C
 Proventil Tablet
 (Salbutamol Sulfate) C
 Amitriptyline
 (Amitriptyline) C
 Cyclobenzaprine
 (Cyclobenzaprine) C
 Augmentin
 (Amoxicillin
 Trihydrate,
 Clavulanate
 Potassium) C
 Ergobel
 (Nicergoline) C

Date:05/27/03ISR Number: 4118061-6Report Type:Expedited (15-DaCompany Report #001-0945-950345

Age: Gender:Male I/FU:F

Outcome PT
 Hospitalization - Abdominal Discomfort
 Initial or Prolonged Anticonvulsant Drug Level
 Other Increased
 Asthenia

Cervical Vertebral
Fracture
Chest Discomfort
Convulsion
Deafness
Deafness Neurosensory
Dental Plaque
Difficulty In Walking
Dizziness
Dysstasia
Feeling Abnormal
Hypoaesthesia
Injury
Joint Stiffness
Muscle Spasms

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Paralysis Road Traffic Accident Somnolence	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		Spinal Cord Injury Tinnitus	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
(UNKNOWN),		Tooth Discolouration	Professional				
ORAL		Tooth Disorder					
UNKNOWN		Visual Acuity Reduced		Dilantin Kapseals (Phenytoin Sodium)	SS		ORAL
(UNKNOWN),							
ORAL				Zoloft	SS		ORAL
UNKNOWN							
(UNKNOWN),							
ORAL				Lamotrigine (Lamotrigine)	SS		
UNKNOWN	UNKNOWN						
(UNKNOWN),							
UNKNOWN							
UNKNOWN				Topiramate (Topiramate)	SS		
(UNKNOWN),	UNKNOWN						
UNKNOWN							
UNKNOWN				Tizanidine Hydrochloride (Tizanidine Hydrochloride)	SS		ORAL
(UNKNOWN),							
ORAL							

Cefuroxime Axetil	C
Fluoxetine	
Hydrochloride	C
Citalopram	
Hydrobromide	C
Levetiracetam	C
Lactulose	C
Lorazepam	C

Date:05/27/03ISR Number: 4118339-6Report Type:Expedited (15-DaCompany Report #2003021558

Age:55 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Chest Pain	Consumer	Neurontin			
Initial or Prolonged	Dizziness		(Gabapentin)	PS		
Other	Dyspnoea		Cisapride			
20 MG	Feeling Abnormal		(Cisapride)	SS		
	Oedema Peripheral		Salmeterol Xinafoate			
	Swelling Face		(Salmeterol			
			Xinafoate)	C		
			Clonazepam			
			(Clonazepam)	C		
			Aciclovir			
			(Aciclovir)	C		
			Lovastatin			
			(Lovastatin)	C		
			Famotidine			
			(Famotidine)	C		
			Dyazide			
			(Hydrochlorothiazide			
			, Triamterene)	C		
			Verapamil			

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Freedom Of Information (FOI) Report

(Verapamil) C

Date:05/28/03ISR Number: 4117884-7Report Type:Direct
Age:61 YR Gender:Male I/FU:I

Company Report #CTU 194088

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 5MG Q12H PRN Initial or Prolonged ORAL	Accidental Overdose		Oxycodone 5mg	PS		ORAL
1400MG QD ORAL			Gabapentin	SS		ORAL
			Tacrolimus	C		
			Cotrimoxazole	C		
			Lansoprazole	C		
			Ketoconazole	C		
			Simvastatin	C		
			Paroxetine	C		
			Warfarin	C		
			Prednisone	C		
			Azathioprine	C		

Date:05/28/03ISR Number: 4117952-XReport Type:Direct
Age:69 YR Gender:Male I/FU:I

Company Report #CTU 194139

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Angioneurotic Oedema		Citalopram	PS		
			Gabapentin	SS		
			Lisinopril	SS		

Date:05/28/03ISR Number: 4118670-4Report Type:Expedited (15-DaCompany Report #200321716
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	No Adverse Drug Effect	Consumer	Neurontin (Gabapentin)	PS		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Antineutrophil	Foreign	Gabapentin			
Initial or Prolonged	Cytoplasmic Antibody	Literature	(Gabapentin)	PS		
	Positive	Health	Tolbutamide			
	Antinuclear Antibody	Professional	(Tolbutamide)	C		
	Positive		Acetylsalicylic Acid			
	Biopsy Skin Abnormal		(Acetylsalicylic			
	Blister		Acid)	C		
	Leukocytoclastic					
	Vasculitis					
	Pain					
	Purpura					
	Pyrexia					
	Rash					
	Swelling					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/28/03ISR Number: 4119693-1Report Type:Expedited (15-DaCompany Report #2003021322
Age:46 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	1800 MG (900, BID)	Maculopathy	Foreign Health Professional	Gabapentin (Gabapentin)	PS		
	200 MG (100, BID),			Topiramate (Topiramate)	SS		
	400 MG (DAILY), ORAL			Carbamazepine (Carbamazepine)	SS		ORAL
	400 MG (200, BID), ORAL			Lamotrigine (Lamotrigine)	SS		ORAL
				Paracetamol (Paracetamol)	C		

Date:05/29/03ISR Number: 4119724-9Report Type:Expedited (15-DaCompany Report #2003021321
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Status Epilepticus	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:05/29/03ISR Number: 4120357-9Report Type:Expedited (15-DaCompany Report #2003001026
Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Cardiac Failure	Foreign	Zoloft (Sertraline)	PS		ORAL
50 MG, ORAL							
Hospitalization -		Cognitive Disorder	Health	Neurontin			
Initial or Prolonged		Condition Aggravated	Professional	(Gabapentin)	SS		ORAL
300 MG							
Other		Depressed Mood					
(DAILY), ORAL							
		Diabetes Mellitus		Fentanyl (Fentanyl)	SS		
TOPICAL	TOPICAL						
		Diaphragmatic Paralysis		Prednisone			
		Difficulty In Walking		(Prednisone)	SS		ORAL
50 MG							
		Erythema					
(DAILY), ORAL							
		Hemiparesis		Carbamazepine			
		Leukoencephalomyelitis		(Carbamazepine)	C		
		Oedema Peripheral		Metamizole Sodium			
		Oral Intake Reduced		(Metamizole Sodium)	C		
		Pain		Baclofen (Baclofen)	C		
		Somnolence		Metoprolol Succinate			
		Stupor		(Metoprolol			
		Swelling Face		Succinate)	C		
		Trigeminal Neuralgia		Valsartan			
				(Valsartan)	C		
				Furosemide			
				(Furosemide)	C		
				Thiamazole			
				(Thiamazole)	C		
				Valoron N (Naloxone			
				Hydrochloride,			
				Tilidine			
				Hydrochloride)	C		
				Amitriptyline			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Amitriptyline) C

Date:05/30/03ISR Number: 4120111-8Report Type:Expedited (15-DaCompany Report #B0299997A
 Age:40 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Paxil (Formulation Unknown) (Paroxetine Hydrochloride)	PS		ORAL
ORAL				Tricyclic Antidepressant (Formulation Unknown) (Tricyclic Antidepressant)	SS		ORAL
ORAL				Gabapentin (Formulation Unknown) (Gabapentin)	SS		ORAL

Date:05/30/03ISR Number: 4120632-8Report Type:Expedited (15-DaCompany Report #USA-2002-0001677
 Age:41 YR Gender:Female I/FU:F

Outcome	PT
Death	Alopecia
Hospitalization - Initial or Prolonged	Amnesia Anxiety Apnoea Arthralgia Arthritis Atherosclerosis Back Pain Cardiac Arrest Cardiac Disorder Chest Pain Coma Condition Aggravated Depression Disturbance In Attention Drug Ineffective Emphysema

Fatigue
Finger Deformity
Flushing
Hilar Lymphadenopathy
Hyperventilation
Hypoaesthesia
Increased Tendency To
Bruise
Influenza
Initial Insomnia
Insomnia
Irritability
Libido Decreased
Memory Impairment
Muscular Weakness
Nausea
Nervousness
Night Sweats
Overdose
Pain

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pain In Extremity Paraesthesia Peripheral Occlusive Disease Pneumonia Portal Triaditis Pruritus	Report Source	Product	Role	Manufacturer	Route
10 MG			Consumer Health Professional Other	Oxycontin Tablets 10 Mg (Oxycodone Hydrochloride) Cr Tablet	PS		
20 MG	1008 DAY	Pulmonary Oedema Rash Pruritic Rash Scaly Shock		Oxycontin Tablets 20 Mg (Oxycodone Hydrochloride) Cr Tablet	SS		
1008 DAY		Somnolence Upper Respiratory Tract Infection Urinary Incontinence Weight Increased		Hydrocodone Bitartrate (Similar To Ind 59,175) (Hydrocodone Bitartrate)	SS		
1008 DAY				Alprazolam (Alprazolam)	SS		
1008 DAY				Ephedrine (Ephedrine)	SS		
1008 DAY				Pseudoephedrine (Pseudoephedrine)	SS		
75 MG, DAILY	1008 DAY			Effexor (Venlafaxine Hydrochloride)	SS		
1008 DAY				Codeine (Codeine)	SS		
100 MG, TID	1008 DAY			Neurontin (Gabapentin)	SS		
1008 DAY				Metoclopramide (Metoclopramide)	SS		
1008 DAY				Quetiapine (Quetiapine)	SS		
				Claritin (Loratadine) Vitamin C (Ascorbic Acid)	C C		

Celebrex (Celecoxib)	C
Prilosec	
(Omeprazole)	C
Daypro (Oxaprozin)	C
Trandate (Labetalol	
Hydrochloride)	C
Axid (Nizatidine)	C
Paxil (Paroxetine	
Hydrochloride)	C
Medrol	
(Methylprednisolone)	C
Macro Antioxidant	
(Ascorbic Acid,	
Cystine, Tocopherol,	
Calcium Ascorbate,	
Betacarotene,	C
Zocor (Simvastatin)	C
Risperdal	
(Risperidone)	C
Fioricet	
(Butalbital)	C
Flonase (Fluticasone	
Propionate)	C
Phenobarbital	
(Phenobarbital)	C
Donnatal (Atropine	

Freedom Of Information (FOI) Report

Sulfate, Hyoscine
 Hydrobromide,
 Hyoscyamine Sulfate,
 Phenobarbital) C
 Bellergal-S
 (Belladonna
 Alkaloids,
 Ergotamine Tartrate,
 Phenobarbital) C
 Weight Loss
 Supplement (Does Not
 Code) C
 Ibuprofen C
 Ambien (Zolpidem
 Tartrate) C
 Baclofen (Baclofen) C
 Trandate (Labetalol
 Hydrochloride) C
 Celexa (Citalopram
 Hydrobromide) C
 Cortisone
 (Cortisone) C
 Atarax (Hydroxyzine
 Hydrochloride) C
 Proventil Tablet
 (Salbutamol Sulfate) C
 Amitriptyline
 (Amitriptyline) C
 Cyclobenzaprine
 (Cyclobenzaprine) C
 Augmentin
 (Amoxicillin
 Trihydrate,
 Clavulanate
 Potassium) C
 Ergobel
 (Nicergoline) C

Date:05/30/03ISR Number: 4121488-XReport Type:Expedited (15-DaCompany Report #2003022137

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 400 MG		Anticonvulsant Drug Level Decreased	Consumer	Neurontin (Gabapentin)	PS		

Condition Aggravated
Convulsion
Pharmaceutical Product
Complaint

Dilantin Suspension
(Phenytoin Sodium) SS

Date:06/02/03ISR Number: 4120645-6Report Type:Direct
Age:39 YR Gender:Female I/FU:I

Company Report #CTU 194457

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG BID Required PO		Stevens-Johnson Syndrome		Gabapentin (Parke Davis)	PS	Parke Davis	ORAL
Intervention to Prevent Permanent Impairment/Damage				Ranitidine Metformin Tylenol/Codine	C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/02/03ISR Number: 4121759-7Report Type:Expedited (15-DaCompany Report #2003021783

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health	Neurontin			
1200 MG		Drug Toxicity	Professional	(Gabapentin)	PS		ORAL
(ONCE), ORAL		Medication Error					
				Lamotrigine			
(ONCE) , ORAL				(Lamotrigine)	SS		ORAL
				Carbamazepine			
(ONCE), ORAL				(Carbamazepine)	SS		ORAL
				Furosemide			
				(Furosemide)	C		
				Potassium			
				(Potassium)	C		
				Insulin (Insulin)	C		
				Oxybutynin			
				(Oxybutynin)	C		
				Sertraline			
				Hydrochloride			
				(Sertraline			
				Hydrochloride)	C		
				Leuprorelin Acetate			
				(Leuprorelin			
				Acetate)	C		
				Risperidone			
				(Risperidone)	C		

Date:06/02/03ISR Number: 4121903-1Report Type:Expedited (15-DaCompany Report #PHFR2003GB02057

Age:46 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Macular Degeneration	Foreign	Carbamazepine			
400 MG/DAY,		Maculopathy	Health	(Carbamazepine)	PS		ORAL
ORAL		Retinal Deposits	Professional				
		Visual Acuity Reduced	Other	Topiramate			

100 MG, BID	(Topiramate)	SS	
	Gabapentin (Gabapentin)	SS	
900 MG, BID	Lamotrigine (Lamotrigine)	SS	ORAL
200 MG, BID, ORAL	Paracetamol	C	

Date:06/02/03ISR Number: 4121995-XReport Type:Expedited (15-DaCompany Report #2003013223
Age:1 DY Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly	Agitation Neonatal	Foreign	Neurontin			
	Blood Glucose Decreased	Health	(Gabapentin)	PS		
	Blood Glucose Increased	Professional	Citalopram			
	Caesarean Section		(Citalopram)	C		
	Complications Of Maternal		Methyldopa			
	Exposure To Therapeutic		(Methyldopa)	C		
	Drugs					
	Fallot'S Tetralogy					
	Irritability					
	Laboratory Test Abnormal					
	Maternal Drugs Affecting					
	Foetus					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/02/03ISR Number: 4121996-1Report Type:Expedited (15-DaCompany Report #2003010615

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 400 MG (DAILY)		Bone Marrow Depression	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:06/02/03ISR Number: 4121997-3Report Type:Expedited (15-DaCompany Report #2003019008

Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1600 MG (QID) ORAL		Dyskinesia Gamma-Glutamyltransferase Increased Hyperhidrosis Hyperkinesia Pyrexia Urinary Tract Infection	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Furosemide (Furosemide) Prednisolone (Prednisolone) Esomeprazole (Esomeprazole) Acetylsalicylic Acid (Acetylsalicylic Acid) Mirtazapine (Mirtazapine) Valoron N (Naloxone Hydrochloride, Tilidine Hydrochloride) Zopiclone (Zopiclone) Amitriptyline Hydrochloride (Amitriptyline Hydrochloride) Enalapril	PS C C C C C C C C C		ORAL

(Enalapril)

C

Date:06/02/03ISR Number: 4122713-1Report Type:Expedited (15-DaCompany Report #2003021781

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Dysphagia	Foreign	Neurontin			
Initial or Prolonged	Local Swelling	Consumer	(Gabapentin)	PS		ORAL
ORAL	Pharyngolaryngeal Pain		Levothyroxine Sodium (Levothyroxine Sodium)	C		
			All Other Therapeutic Products	C		

Date:06/03/03ISR Number: 4123196-8Report Type:Expedited (15-DaCompany Report #2003022132

Age: 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

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG DAILY		Somnolence	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL				Furosemide	C		
				Potassium Chloride	C		
				Losartan Potassium	C		
				Pravastatin Sodium	C		
				Warfarin Sodium	C		
				Digoxin	C		
				Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)	C		
				Glipizide	C		
				Entex (Phenylephrine Hydrochloride, Guaifenesin, Phenylpropanolamine Hydrochloride)	C		
				Metoprolol Succinate	C		
				Pantoprazole	C		
				Combivent (Ipratropium Bromide, Salbutamol Sulfate)	C		
				All Other Therapeutic Products	C		

Date:06/03/03ISR Number: 4123200-7Report Type:Expedited (15-DaCompany Report #2002056208

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cholecystitis	Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Drug Hypersensitivity	Professional				
		Drug Ineffective		Atenolol	C		
		Hepatitis Cholestatic		Acetylsalicylic Acid	C		

Murphy'S Sign Positive
Pyrexia
Splenomegaly

Ranitidine
Hydrochloride C
All Other
Therapeutic Products C

Date:06/03/03ISR Number: 4123493-6Report Type:Expedited (15-DaCompany Report #2003022526

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Facial Palsy	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
1500 MG (300			Professional				
, FIVE TIMES							
DAILY) ORAL							

Amitripytline C
Dosulepin C
Acetylsalicylic Acid C
Fenofibrate C
Mirtazapine C
Furosemide C

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Freedom Of Information (FOI) Report

Gaviscon (Sodium
Bicarbonate, Sodium
Alginate) C
Senna C
Lactulose C

Date:06/04/03ISR Number: 4123564-4Report Type:Expedited (15-DaCompany Report #2003022993
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abnormal Behaviour	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:06/04/03ISR Number: 4123570-XReport Type:Expedited (15-DaCompany Report #2003022681
Age:86 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG Other (TID), ORAL		Blood Creatinine Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
20 MG (DAILY), ORAL		Blood Pressure Decreased	Professional				
		Cachexia Chills Dehydration Fatigue		Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		ORAL
1200 MG (DAILY), ORAL		Haematuria					
		Loss Of Consciousness Medication Error		Aciclovir (Aciclovir)	SS		ORAL
		Pyrexia					
		Somnolence Treatment Noncompliance		Tramadol Hydrochloride (Tramadol			

200 MG

Hydrochloride) SS ORAL

(DAILY), ORAL

Digoxin
(Digoxin) C
Paracetamol
(Paracetamol) C
Acetylsalicylic Acid C

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) Gabapentin (Formulation Unknown) (Gabapentin)	SS SS		

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Date:06/04/03ISR Number: 4124357-4Report Type:Expedited (15-DaCompany Report #2003014853

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 2400 MG (800, Initial or Prolonged TID) Disability	Cardiac Failure Congestive Congestive Cardiomyopathy	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:06/04/03ISR Number: 4124707-9Report Type:Expedited (15-DaCompany Report #2003022646

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL Initial or Prolonged Other	Abortion Spontaneous Chest Pain Complications Of Maternal Exposure To Therapeutic Drugs Condition Aggravated Constipation	Consumer	Zoloft (Sertraline) Neurontin (Gabapentin) Zyrtec-D 12 Hour (Cetirizine/Pseudoep hedrine)	PS SS SS		ORAL ORAL ORAL
ORAL ORAL	Disease Recurrence Fatigue		Zyrtec (Cetirizine Hydrochloride)	SS		ORAL
	Fibroma Hearing Impaired Hypoaesthesia Hypotonia Ileus Incision Site Complication Maternal Drugs Affecting Foetus Ovarian Cyst Ovarian Hyperstimulation Syndrome Panic Attack Postoperative Wound Complication Seasonal Allergy		All Other Therapeutic Products Ketamine (Ketamine) Hydroxyzine Embonate (Hydroxyzine Embonate) Alprazolam (Alprazolam)	SS SS C C		

Sinus Congestion
Somnolence
Urinary Retention
Vertigo

Date:06/05/03ISR Number: 4123419-5Report Type:Expedited (15-DaCompany Report #WAES 0306FRA00005

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	26 DAY	Drug Eruption		Noroxin	PS	Merck & Co., Inc	ORAL
Initial or Prolonged	56 DAY	Eosinophilia		Carbamazepine	SS		ORAL
	16 DAY	Hepatocellular Damage		Gabapentin	SS		ORAL
		Hypersensitivity					
		Lymph Node Palpable					
		Oedema Peripheral					
		Pyrexia					

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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			
UNKNOWN		Maternal Drugs Affecting	Professional	(Gabapentin)	PS		ORAL
(UNKNOWN),		Foetus					
ORAL		Pregnancy					
UNKNOWN	UNKNOWN			Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		
(UNKNOWN),							
UNKNOWN				Rizatriptan Benzoate (Rizatriptan Benzoate)	SS		
UNKNOWN	UNKNOWN						
(UNKNOWN),							
UNKNOWN							

Date:06/05/03ISR Number: 4124779-1Report Type:Expedited (15-DaCompany Report #2003023124

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dental Caries	Consumer	Neurontin			
Initial or Prolonged		Diarrhoea		(Gabapentin)	PS		ORAL
2700 MG		Gallbladder Disorder					
(TID), ORAL		Hiatus Hernia		Ultracet	C		
		Liver Function Test		Tramadol	C		
		Abnormal		Oxycocet	C		
		Tooth Loss		Propacet	C		
		Weight Increased		Vicodin	C		

Methadone C
Clonidine C
Amitriptyline
Hydrochloride C

Date:06/05/03ISR Number: 4125341-7Report Type:Expedited (15-DaCompany Report #2003001026
Age:74 YR Gender:Female I/FU:F

Outcome PT
Life-Threatening Cardiac Failure
Hospitalization - Depressed Level Of
Initial or Prolonged Consciousness
Other Diaphragmatic Paralysis
Difficulty In Walking
Disturbance In Attention
Drug Interaction
Dry Mouth
Encephalitis
Fall
Femoral Neck Fracture
Fluid Intake Reduced
General Physical Health
Deterioration
Multiple Sclerosis
Oral Intake Reduced
Overdose
Paresis
Somnolence

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Dose	Duration	Therapeutic Agent	Report Source	Product	Role	Manufacturer	Route
50 MG, ORAL		Stupor Swelling Face Toxicity	Foreign	Zoloft (Sertraline)	PS		ORAL
300 MG (DAILY), ORAL			Health Professional	Neurontin (Gabapentin)	SS		ORAL
TOPICAL	TOPICAL			Fentanyl (Fentanyl)	SS		
50 MG (DAILY) ORAL				Prednisone (Prednisone)	SS		ORAL
				Carbamazepine (Carbamazepine)	C		
				Metamizole Sodium (Metamizole Sodium)	C		
				Baclofen (Baclofen)	C		
				Metoprolol Succinate (Metoprolol Succinate)	C		
				Valsartan (Valsartan)	C		
				Furosemide (Furosemide)	C		
				Thiamazole (Thiamazole)	C		
				Valoron N (Naloxone Hydrochloride, Tilidine Hydrochloride)	C		
				Amitriptyline (Amitriptyline)	C		
				Metoclopramide Hydrochloride (Netoclopramide Hydrochloride)	C		
				Diazepam (Diazepam)	C		

Age:43 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG Other (DAILY), ORAL	Colitis Coma Cytomegalovirus Infection Hepatitis Leukopenia Neutropenia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
UNKNOWN (UNKNOWN), ORAL			Nevirapine (Nevirapine)	SS		ORAL
UNKNOWN (BID), ORAL			Kaletra (Ritonavir, Lopinavir)	SS		ORAL
UNKNOWN (BID), ORAL			Tenofovir Disoproxil Fumarate (Tenofovir Disoproxil Fumarate)	SS		ORAL
UNKNOWN (DAILY), ORAL			Valganciclovir (Valcanciclovir)	SS		ORAL

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(BID), ORAL

Paracetamol (Paracetamol)	C
Loperamide (Loperamide)	C
Rifampicin (Rifampicin)	C
Ethambutol Dihydrochloride (Ethambutol Dihydrochloride)	C
Clarithromycin (Clarithromycin)	C
Bactrim (Sulfamethoxazole, Trimethoprim)	C
Calcium Folate (Calcium Folate)	C
Filgrastim (Filgrastim)	C

Date:06/09/03ISR Number: 4126228-6Report Type:Expedited (15-DaCompany Report #2003015349

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (DAILY), ORAL		Anticonvulsant Drug Level Above Therapeutic Apallic Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
750 MG (DAILY), ORAL		Cognitive Disorder Crying Drug Toxicity Nightmare		Valproate Semisodium (Valproate Semisodium)	SS		ORAL
		Oral Intake Reduced Respiratory Arrest Screaming Speech Disorder Weight Gain Poor		Lamotrigine (Lamotrigine) Topiramate(Topiramate)	SS SS		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 50 MG, ORAL	Duration Blood Glucose Increased	Foreign	Zoloft (Sertraline)	PS		ORAL
Hospitalization - Initial or Prolonged 300 MG	Cardiac Failure Drug Toxicity	Health Professional	Neurontin (Gabapentin)	SS		ORAL
Other (DAILY), ORAL	Encephalitis					
TOPICAL	Hemiparesis TOPICAL		Fentanyl (Fentanyl)	SS		
50 MG	Iatrogenic Injury Oedema Peripheral		Prednisone (Prednisone)	SS		ORAL
(DAILY), ORAL	Oral Intake Reduced					
	Pain		Carbamazepine (Carbamazepine)	C		
	Stupor		Metamizole Sodium (Metamizole Sodium)	C		
	Swelling		Baclofen (Baclofen)	C		
	Trigeminal Neuralgia		Metoprolol Succinate (Metoprolol Succinate)	C		
			Valsartan			

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(Valsartan)	C
Furosemide	
(Furosemide)	C
Thiamazole	
(Thiamazole)	C
Valoron N (Naloxone Hydrochloride, Tilidine Hydrochloride)	C
Amitriptyline (Amitriptyline)	C
Metoclopramide Hydrochloride (Metoclopramide Hydrochloride)	C
Diazepam (Diazepam)	C

Date:06/09/03ISR Number: 4126330-9Report Type:Expedited (15-DaCompany Report #2003012656
 Age:69 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	100 MG ONCE,	Transient Ischaemic Attack	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL			Company Representative	Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Diltiazem (Diltiazem)	C		
				Dipyridamole (Dipyridamole)	C		
				Estrogens Conjugated (Estrogens Conjugated)	C		
				Naproxen (Naproxen)	C		
				Omeprazole (Omeprazole)	C		
				Tramadol (Tramadol)	C		

Date:06/09/03ISR Number: 4126331-0Report Type:Expedited (15-DaCompany Report #2003022679
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG, ORAL		Fall Hypotension Mydriasis Somnolence	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Loprazolam (Loprazolam) All Other Therapeutic Products	PS C C		ORAL

Date:06/10/03ISR Number: 4126743-5Report Type:Expedited (15-DaCompany Report #2003023115
Age:53 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Blood Gases Abnormal Blood Potassium Decreased Headache Loss Of Consciousness Somnolence

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
900 MG (300, TID) , ORAL		Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Carisoprodol (Carisoprodol)	C		
			Paracetamol (Paracetamol)	C		
			Panadeine Co (Codeine Phosphate, Paracetamol)	C		
			Diazepam (Diazepam)	C		

Date:06/10/03ISR Number: 4127552-3Report Type:Expedited (15-DaCompany Report #2003023632
Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	1200 MG (TID); ORAL	Neutrophil Count Platelet Count White Blood Cell Count	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Clonazepam	C		
				Heparin-Fraction, Sodium Salt	C		

Date:06/10/03ISR Number: 4127839-4Report Type:Expedited (15-DaCompany Report #2003023513
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	400 MG (DAILY), ORAL	Balance Disorder Chest Pain Confusional State Disorientation	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Venlafaxine			

Feeling Abnormal	Hydrochloride	
Joint Dislocation	(Venlafaxine	
Muscular Dystrophy	Hydrochloride)	C
Nervous System Disorder	Nefazodone	
Nightmare	Hydrochloride	
Tremor	(Nefazodone	
	Hydrochloride)	C
	Ropinirole	
	Hydrochloride	
	(Ropinirole	
	Hydrochloride)	C
	Cyclobenzapine	
	Hydrochloride	
	(Cyclobenzaprine	
	Hydrochloride)	C
	Obetrol	
	(Dexamfetamine	
	Sulfate, Amfetamine	
	Sulfate,	
	Dexamfetamine	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/11/03ISR Number: 4127076-3Report Type:Direct
 Age:62 YR Gender:Male I/FU:I

Company Report #CTU 195605

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Wall Pain		Gabapentin			
		Diarrhoea		(Neurontin)	PS		ORAL
300 MG PO TID		Dyspepsia		Beclomethasone Nasal			
				Inhaler	C		
				Clonidine Patch	C		
				Glyburide	C		
				Lisinopril	C		
				Metformin	C		
				Nifedipine	C		
				Ntg	C		
				Prazosin	C		
				Rabeprazole	C		

Date:06/11/03ISR Number: 4127078-7Report Type:Direct
 Age:44 YR Gender:Male I/FU:I

Company Report #CTU 195601

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache		Gabapentin			
		Treatment Noncompliance		(Neurontin)	PS		
300 MG PO QD							
TITRATE UP TO							
TID OVER 3							
WEEKS							
				Albuterol	C		
				Diazepam	C		
				Vicodin	C		
				Paroxetine	C		

Date:06/11/03ISR Number: 4128727-XReport Type:Expedited (15-DaCompany Report #2003006332
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged 1200 MG (QID), ORAL	Back Pain Bipolar Disorder Fall Fatigue Joint Swelling Myalgia Oedema Peripheral Tendonitis Weight Increased	Consumer	Neurontin (Gabapentin)	PS	ORAL
			Buspirone Hydrochloride (Buspirone Hydrochloride)	C	
			Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C	
			Levothyroxine Sodium (Levothyroxine Sodium)	C	
			Hypnotics And Sedatives	C	
			Hydroxyzine Embonate (Hydroxyzine Embonate)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/11/03ISR Number: 4128747-5Report Type:Expedited (15-DaCompany Report #2003020214
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (BID), Other ORAL		Depression Dialysis Drug Screen Positive Renal Failure Acute Renal Tubular Necrosis	Health Professional	Neurontin (Gabepentin) Cocaine (Cocaine) Vicodin (Paracetamol, Hydrocodone Bitartrate)	PS C C		ORAL

Date:06/11/03ISR Number: 4128827-4Report Type:Expedited (15-DaCompany Report #2003012199
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG (DAILY), ORAL		Condition Aggravated Cystitis Noninfective Eye Swelling Fluid Retention Rash Macular Swelling Face	Consumer Health Professional	Neurontin (Gabapentin) Pentosan Polysulfate Sodium (Pentosan Polysulfate Sodium) Amitriptyline Hydrochloride (Amitriptyline Hydrochloride) Clonazepam (Clonazepam) Rofecoxib (Rofecoxib) Simvastatin (Simvastatin) Estradiol (Estradiol) Oxycocet (Paracetamol, Oxycodone Hydrochloride)	PS C C C C C C		

Bupropion
Hydrochloride
(Bupropion
Hydrochloride) C

Date:06/11/03ISR Number: 4129266-2Report Type:Expedited (15-DaCompany Report #B0299996A

Age:28 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Paxil (Paroxetine Hydrochloride)	PS		ORAL
ORAL				Gabapentin (Gabapentin)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/12/03ISR Number: 4128872-9Report Type:Expedited (15-DaCompany Report #2002052306
 Age:83 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (THREE TIMES DAILY), ORAL	Arrhythmia Cardiac Failure Drug Toxicity Dyspnoea	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
			Amitriptyline (Amitriptyline)	C		
			Ginkgo Biloba (Ginkgo Biloba)	C		
			Omeprazole (Omeprazole)	C		
			Verapamil (Verapamil)	C		
			Spironolactone (Spironolactone)	C		
			Methadone Hydrochloride (Methadone Hydrochloride)	C		
			Meloxicam (Meloxicam)	C		
			Homeopathic Drug	C		

Date:06/13/03ISR Number: 4128423-9Report Type:Direct Company Report #CTU 195712
 Age:71 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Required 300 MG PO TID Intervention to Prevent Permanent Impairment/Damage	Abdominal Pain Asthenia Back Pain Hyperhidrosis		Gabapentin	PS		ORAL

Date:06/13/03ISR Number: 4129661-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
 Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Alanine Aminotransferase
Initial or Prolonged	Increased
Disability	Amnesia
Other	Anhedonia
	Anticonvulsant Drug Level
	Below Therapeutic
	Back Pain
	Blood Cholesterol
	Increased
	Blood Glucose Increased
	Brain Neoplasm
	Cardio-Respiratory Arrest
	Cognitive Disorder
	Confusional State
	Decreased Appetite
	Depression
	Dizziness
	Dysphoria
	Eosinophil Count
	Increased
	Ependymoma

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (BID), ORAL		Fall Fatigue Feeling Abnormal Grand Mal Convulsion Hyperlipidaemia Hypertension Hypertriglyceridaemia Irritability Joint Sprain Libido Decreased Mean Platelet Volume Decreased Memory Impairment Nervous System Disorder Osteoarthritis Pain In Extremity Paraesthesia Rash Red Blood Cell Sedimentation Rate Increased Sleep Disorder Social Avoidant Behaviour Somnolence Speech Disorder Tinnitus Tremor Vomiting	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin)	PS SS C C C C		ORAL

Date:06/13/03ISR Number: 4129836-1Report Type:Expedited (15-DaCompany Report #2003021210
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG (DAILY), ORAL		Coordination Abnormal Drug Ineffective Dyskinesia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Asthma	Consumer	Neurontin			
Initial or Prolonged	Condition Aggravated	Health	(Gabapentin)	PS		ORAL
600 MG (BID),						
Other	Cystitis	Professional				
ORAL						
	Fall		Prednisone			
	Gilbert'S Syndrome		(Prednisone)	SS		
	Haemolytic Anaemia		Acetylsalicyclic			
	Headache		Acid			
	Kidney Infection		(Acetylsalicylic			
	Pneumonia		Acid)	C		
	Pyrexia		Insulin Lispro			
	Tremor		(Insulin Lispro)	C		
	Urinary Tract Infection		All Other			
	Weight Decreased		Therapeutic Products	C		
			Simvastatin			
			(Simvastatin)	C		
			Seretide Mite			

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Freedom Of Information (FOI) Report

(Fluticasone
 Propionate,
 Salmeterol
 Xinafoate) C
 Furosemide
 (Furosemide) C
 Combivent
 (Ipratropium
 Bromide, Salbutamol
 Sulfate) C
 Propacet
 (Paracetamol,
 Dextropropoxyphene
 Napsilate) C
 Digoxin (Digoxin) C
 Epoetin Alfa
 (Epoetin Alfa) C

Date:06/16/03ISR Number: 4129589-7Report Type:Direct
 Age:45 YR Gender:Female I/FU:I

Company Report #CTU 195891

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination Nausea		Gabapentin (Neurontin)	PS		

Date:06/16/03ISR Number: 4130303-XReport Type:Expedited (15-DaCompany Report #2003020630
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract Neuropathic Pain	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (300, BID), ORAL		Retinopathy	Professional				
		Transient Ischaemic Attack		Amlodipine Besilate (Amlodipine Besilate)	C		
				Insulin Human (Insulin Human)	C		
				Simvastatin (Simvastatin)	C		
				Asasantin			

(Acetylsalicylic Acid, Dipyrindamole)	C
Potassium (Potassium)	C
Furosemide (Furosemide)	C
Folic Acid (Folic Acid)	C
Pramipexole (Pramipexole)	C
Alprazolam (Alprazolam)	C
Cyanocobalamin (Cyanocobalamin)	C
Iron (Iron)	C
Insulin Human Injection, Isophane (Insulin Human Injection, Isophane)	C

Freedom Of Information (FOI) Report

Date:06/16/03ISR Number: 4130307-7Report Type:Expedited (15-DaCompany Report #2003015172
 Age:82 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG Other (DAILY), ORAL	Abdominal Distension Anuria Balance Disorder Blood Potassium Increased Blood Pressure Decreased Dizziness Dyspnoea Fatigue Gait Disturbance Haemoptysis Heart Rate Decreased Heart Rate Irregular Joint Swelling Movement Disorder Muscular Weakness Po2 Decreased Renal Failure Acute Somnolence Treatment Noncompliance	Consumer	Neurontin (Gabapentin) Lisinopril (Lisinopril) Antihypertensives Verapamil (Verapamil) Estrogens Conjugated (Estrogens Conjugated)	PS C C C C		ORAL

Date:06/16/03ISR Number: 4130332-6Report Type:Expedited (15-DaCompany Report #002#2#2002-00109 (0)
 Age:43 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability	Akathisia Arthropathy Blepharospasm Blood Pressure Increased Chest Discomfort Condition Aggravated Conversion Disorder Depression Diarrhoea Disturbance In Attention Drug Withdrawal Syndrome Dry Mouth Dysphagia Dysphemia

Dyspnoea
Dystonia
Eye Pain
Feeling Abnormal
Halo Vision
Headache
Hyperhidrosis
Impaired Driving Ability
Malaise
Migraine
Muscle Spasms
Nausea
Nervousness
Nightmare
Obsessive-Compulsive
Disorder
Palpitations
Paraesthesia
Photosensitivity Reaction

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
10MG, 4 IN 1 D, ORAL		Posture Abnormal Restlessness Sleep Apnoea Syndrome Speech Disorder Strabismus Syncope Tardive Dyskinesia Tic Tremor Visual Acuity Reduced	Consumer	Reglan-10mg-Tablet (Metoclopramide Hcl)	PS		ORAL
				Prochlorperazine-Edi sylate	SS		
				Haloperidol	SS		
				Gabapentin	SS		
				Olanzapine	SS		
				Lisinopril	C		
				Diazepam	C		
				Mylanta	C		
				Metoprolol	C		
				Heparin	C		
				Clopidogrel	C		

Date:06/16/03ISR Number: 4130664-1Report Type:Expedited (15-DaCompany Report #2003024541

Age: Gender:Female I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
300 MG (UNKNOWN), ORAL		PT Asthenia Brain Oedema Fatigue	Foreign Health Professional Company Representative	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Morphine	C		

Date:06/16/03ISR Number: 4130667-7Report Type:Expedited (15-DaCompany Report #2003018752

Age:74 YR Gender:Male I/FU:F

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
ORAL		Cytolytic Hepatitis Depressed Level Of	Foreign Health	Neurontin (Gabapentin)	PS		ORAL

Consciousness	Professional	Paroxetine	
Disturbance In Attention	Company	(Paroxetine)	C
	Representative	Hydromorphone	
		(Hydromorphone)	C
		Mianserin	
		(Mianserin)	C
		Morphine Sulfate	
		(Morphine Sulfate)	C

Date:06/16/03ISR Number: 4130682-3Report Type:Expedited (15-DaCompany Report #2003024245
Age:44 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Angiopathy
Initial or Prolonged	Cholestasis
Other	Cytolytic Hepatitis
	Drug Rash With
	Eosinophilia And Systemic
	Symptoms
	Hemiplegia
	Lymphadenopathy
	Oedema Peripheral
	Rash Erythematous
	Rash Papular

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Freedom Of Information (FOI) Report

Dose	Duration	Rash Pruritic White Blood Cell Count	Report Source	Product	Role	Manufacturer	Route
1800 MG, ORAL			Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
400 MG, ORAL				Carbamazepine (Carbamazepine)	SS		ORAL
400 MG, ORAL				Norfloxacin (Norfloxacin)	SS		ORAL
				Ketoprofen (Ketoprofen)	C		
				Bromazepam (Bromazepam)	C		
				Enoxaparin (Heparin)	C		

Date:06/17/03ISR Number: 4131360-7Report Type:Expedited (15-DaCompany Report #2003024542
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Intolerance	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:06/17/03ISR Number: 4131364-4Report Type:Expedited (15-DaCompany Report #2003024623
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (BID), Other ORAL		Blister Drug Hypersensitivity	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Drug Ineffective					
		Dyspnoea Fibrocystic Breast Disease		Benadryl Injection (Diphenhydramine)	SS		
INTRAVENOUS	INTRAVENOUS			Ciprofloxacin			

Infection
 Menorrhagia
 Ovarian Cyst
 Swollen Tongue
 Tongue Blistering

(Ciprofloxacin) SS
 Paracetamol
 (Paracetamol) C

Date:06/17/03ISR Number: 4131465-0Report Type:Expedited (15-DaCompany Report #2003024696
 Age:58 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG Other (BID), GASTROSTOMY TUBE	Agitation Balance Disorder Drug Withdrawal Syndrome Dyskinesia Hallucination Hyperhidrosis Mental Disorder Tremor	Foreign Health Professional	Neurontin (Gabapentin) Lansoprazole (Lansoprazole) Nulytely (Sodium Bicarbonate, Potassium Chloride, Sodium Chloride, Macrogol) Morphine Sulfate (Morphine Sulfate)	PS C C C		

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Paroxetine
 Hydrochloride
 (Paroxetine
 Hydrochloride) C
 Fentanyl (Fentanyl) C
 Baclofen (Baclofen) C

Date:06/17/03ISR Number: 4131558-8Report Type:Expedited (15-DaCompany Report #2003001285
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Gamma-Glutamyltransferase Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
(THREE TIMES DAILY), ORAL		Hepatic Enzyme Increased	Professional				
		Hepatic Steatosis		Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
				Clonazepam (Clonazepam)	C		

Date:06/17/03ISR Number: 4131704-6Report Type:Expedited (15-DaCompany Report #2003017240
 Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Convulsion					
		Drug Ineffective		Insulin Injection, Isophane (Insulin Injection, Isophane)	C		
		Drug Withdrawal Syndrome		Insulin Lispro (Insulin Lispro)	C		
		Dystonia		Diclofenac (Diclofenac)	C		
		Exostosis					
		Intervertebral Disc Protrusion					
		Multiple Sclerosis					
		Muscle Twitching					
		Treatment Noncompliance					

Date:06/17/03ISR Number: 4131903-3Report Type:Expedited (15-DaCompany Report #2003024694
Age:15 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Foreign	Neurontin			
		Loss Of Consciousness	Consumer	(Gabapentin)	PS		
UNKNOWN	400 MG						

(DAILY),

UNKNOWN (SEE
IMAGE)

Date:06/18/03ISR Number: 4132401-3Report Type:Expedited (15-DaCompany Report #2003024807
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Convulsion	Consumer	Neurontin (Tablets)			
Other		Dementia Alzheimer'S Type		(Gabapentin)	PS		ORAL
1800 MG (600,		Disease Progression					
TID), ORAL							

5 MG (DAILY),

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				Norvasc (Amlodipine Besilate) (Amlodipine)	SS		ORAL
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ORAL

Rivastigmine	
Tartrate	
(Rivastigmine	
Tartrate)	C
Olanzapine	
(Olanzapine)	C
Acetylsalicylic Acid	
(Acetylsalicylic	
Acid)	C
Tocopherol	
(Tocopherol)	C
Retinol (Retinol)	C

Date:06/18/03ISR Number: 4132405-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
 Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Adjustment Disorder
Initial or Prolonged	Albumin Globulin Ratio
Disability	Decreased
Other	Anxiety
	Back Pain
	Brain Neoplasm
	Cardiac Arrest
	Chest Pain
	Cognitive Disorder
	Confusional State
	Convulsion
	Depression
	Diplopia
	Disturbance In Attention
	Dysarthria
	Dysphemia
	Emotional Disorder
	Enchondromatosis
	Ependymoma
	Fall
	Fatigue
	Feeling Abnormal
	Glucose Tolerance
	Impaired
	Headache
	Hearing Impaired
	Memory Impairment

Monocyte Count Decreased
Mood Altered
Osteoarthritis
Pain In Extremity
Patellofemoral Pain
Syndrome
Rash
Sick Sinus Syndrome
Somnolence
Speech Disorder
Stress
Surgical Procedure
Repeated
Synovitis
Tension
Tremor

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Freedom Of Information (FOI) Report

Dose	Duration	Urine Ketone Body Present Visual Disturbance	Report Source	Product	Role	Manufacturer	Route
400 MG (BID), ORAL			Consumer Health Professional Company Representative	Neurontin (Gabapentin) Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin)	PS SS C C C C		ORAL

Date:06/18/03ISR Number: 4132438-4Report Type:Expedited (15-DaCompany Report #2003024808
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Body Temperature Decreased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Catatonia Coma	Company Representative	Alprazolam (Alprazolam)	SS		ORAL
		Intentional Misuse Pupil Fixed Pyrexia		Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride)	SS		

Date:06/19/03ISR Number: 4132868-0Report Type:Expedited (15-DaCompany Report #2003024695
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour Anorexia Cerebral Ischaemia	Foreign Health Professional	Atarax (Tablet) (Hydroxyzine Hydrochloride)	PS		ORAL
100 MG DAILY, ORAL		Cerebrovascular Accident					
1200 MG DAILY, ORAL		Coma Depression Somnolence		Neurontin (Gabapentin)	SS		ORAL
100 MG (DAILY), ORAL		Urinary Incontinence		Cyamemazine (Cyamemazine)	SS		ORAL
ORAL				Aporex (Paracetmol, Dextropropoxyphene Hydrochloride)	SS		ORAL
30 MG DAILY, ORAL				Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		ORAL
				Clorazepate Dipotassium (Clorazepate			

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25 MG DAILY,
ORAL

Dipotassium) SS ORAL

Zopiclone
(Zopiclone) C

Date:06/19/03ISR Number: 4132884-9Report Type:Expedited (15-DaCompany Report #2003018779
Age:90 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Pressure Abnormal	Foreign	Neurontin (Tablets)			
Other		Cerebral Haemorrhage	Health	(Gabapentin)	PS		ORAL
1200 MG BID		Coma	Professional				
ORAL			Company Representative				

Date:06/19/03ISR Number: 4132948-XReport Type:Expedited (15-DaCompany Report #2003025422
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Fatigue	Consumer	Tikosyn (Dofetilide)	PS		ORAL
1000 MCG		Hypersomnia					
Initial or Prolonged		Somnolence		Neurontin			
(BID), ORAL		Weight Decreased		(Gabapentin)	SS		ORAL
ORAL				Tacrolimus			
				(Tacrolimus)	SS		
				Citalopram			
				Hydrobromide			
				(Citalopram			
				Hydrobromide)	SS		ORAL
ORAL				Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	SS		
				Mycophenolate			

1 GRAM (BID),

ORAL

Mofetil (Mycophenolate Mofetil)	SS	ORAL
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Goserelin (Goserelin)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C
Warfarin (Warfarin)	C

Date:06/19/03ISR Number: 4137835-9Report Type:Periodic
Age:60 YR Gender:Female I/FU:I

Company Report #325112

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Toxicity	Literature	Clonazepam			
Hospitalization - ORAL			Health	(Clonazepam)	PS		ORAL
Initial or Prolonged			Professional	Acetaminophen/ Hydrocodone (Acetaminophen/ Hydrocodone Bitartrate)	SS		ORAL
ORAL				Alprazolam (Alprazolam)	SS		ORAL
ORAL				Paroxetine			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL			(Paroxetine)	SS		ORAL
ORAL			Pentazocine (Pentazocine)	SS		ORAL
ORAL			Gabapentin (Gabapentin)	SS		ORAL
ORAL			Venlafaxine (Venlafaxine Hydrochloride)	SS		ORAL
ORAL			Morphine (Morphine Sulfate)	SS		ORAL

Date:06/19/03ISR Number: 4188922-0Report Type:Periodic Company Report #325146
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health	Klonopin (Clonazepam)	PS		ORAL
ORAL			Professional	Elavil (Amitriptyline Hydrochloride)	SS		ORAL
ORAL				Adderall (Amphetamine Aspartate/Amphetamin e Sulfated/Dextroamphe	SS		ORAL
ORAL				Vioxx (Rofecoxib)	SS		ORAL
ORAL				Topamax(Topiramate)	SS		ORAL
ORAL				Geodon (Ziprasidone Hydrochloride)	SS		ORAL
ORAL				Remeron (Mirtazapine)	SS		ORAL
ORAL				Methadone (Methadone			

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL				Hydrochloride)	SS		ORAL
				Neurontin (Gabapentin)	SS		ORAL
ORAL							
Date:06/20/03ISR Number: 4133916-4Report Type:Expedited (15-DaCompany Report #03P-056-0221579-00							
Age:64 YR Gender:Female I/FU:I							
Hospitalization - Initial or Prolonged		Cerebellar Syndrome Dysarthria Nervous System Disorder	Foreign Health Professional	Kaletra Soft Gelatin Capsules (Kaletra) (Lopinavir/Ritonavir)	PS		ORAL
2 DOSAGE							
FORMS, 1 IN 1							
D, PER ORAL				Enalapril	SS		ORAL
2.5 MG, 1 IN							
1 D, PER ORAL				Fluoxetine	SS		ORAL
20 MG, 1 IN 1							
D, PER ORAL				Gabapentin	SS		ORAL
300 MG, 2 IN							
1 D, PER ORAL				Valganciclovir	SS		ORAL
450 MG,2 IN 1							
D, PER ORAL				Tenofovir	SS		ORAL
245 MG, 2 IN							
1 D, PER ORAL				Aspirin	SS		ORAL
PER ORAL				Levothyroxine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/23/03ISR Number: 4134463-6Report Type:Expedited (15-DaCompany Report #2003025018

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Confusional State	Consumer	Neurontin (Gabapentin)	PS		
Other		Convulsion		Dilantin Suspension (Phenytoin Sodium)	SS		ORAL
		Drug Level Decreased					ORAL

Date:06/23/03ISR Number: 4134464-8Report Type:Expedited (15-DaCompany Report #2003025605

Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dental Caries	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300, TID), ORAL		Drug Effect Decreased					
		Face Injury					
2400 MG (800, TID), ORAL		Nervous System Disorder		Ibuprofen (Ibuprofen)	SS		ORAL
		Toothache					
		Ulcer					
		White Blood Cell Count Decreased		Estrogens Conjugated (Estrogens Conjugated)	C		
				Lorazepam (Lorazepam)	C		

Date:06/23/03ISR Number: 4134467-3Report Type:Expedited (15-DaCompany Report #2003025233

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Crying	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG (600, Other QID), ORAL		Disorientation					
		Fear					
		Tinnitus		Vistrail (Caps)			

(PRN), ORAL

(Hydroxyzien Pamote) SS

ORAL

Amitriptyline
Hydrochloride
(Amitriptyline
Hydrochloride) C
Trazodone
(Trazodone) C
Zaleplon (Zaleplon) C

Date:06/23/03ISR Number: 4134497-1Report Type:Expedited (15-DaCompany Report #2003015172

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

Outcome	PT
Hospitalization -	Abdominal Distension
Initial or Prolonged	Anuria
Other	Blood Pressure Decreased
	Difficulty In Walking
	Dizziness
	Drug Level Above
	Therapeutic
	Dyspnoea
	Fatigue
	Gait Disturbance
	Haemoptysis
	Heart Rate Decreased
	Heart Rate Irregular

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Movement Disorder Muscular Weakness Renal Failure Acute	Report Source	Product	Role	Manufacturer	Route
300 MG		Sedation	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL			Professional	Lisinopril (Lisinopril) Antihypertensives	C		
				Verapamil (Verapamil)	C		
				Estrogens Conjugated (Estrogens Conjugated)	C		

Date:06/23/03ISR Number: 4134499-5Report Type:Expedited (15-DaCompany Report #2003018309
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Pressure Increased Blood Urine	Consumer	Neurontin (Gabapentin)	PS		ORAL
THREE TIMES DAILY, ORAL		Faeces Discoloured					
		Gingival Bleeding International Normalised Ratio Increased Joint Swelling Mental Status Changes Penile Haemorrhage Prothrombin Time Prolonged Pyrexia Tooth Disorder Vascular Fragility		Valaciclovir Hydrochloride (Valaciclovir Hydrochloride) Warfarin Sodium (Warfarin Sodium) Diuretics Lidocaine Hydrochloride (Lidocaine Hydrochloride)	SS SS C C		

Date:06/23/03ISR Number: 4134615-5Report Type:Expedited (15-DaCompany Report #2003025601
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Pancytopenia	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
300 MG			Professional				
(DAILY), ORAL				Methotrexate (Methotrexate)	C		

Date:06/24/03ISR Number: 4134692-1Report Type:Direct Company Report #CTU 196524
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Angioneurotic Oedema		Neurontin	PS		
200MG TID (2		Oedema Peripheral					
Intervention to							
DOSES)							
Prevent Permanent							
Impairment/Damage							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/24/03ISR Number: 4135421-8Report Type:Expedited (15-DaCompany Report #2003019548

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fall	Foreign	Neurontin			
900 MG (300,		Hypoaesthesia	Health	(Gabapentin)	PS		ORAL
TID) ORAL		Respiratory Disorder	Professional				
			Company				
			Representative				

Date:06/24/03ISR Number: 4135425-5Report Type:Expedited (15-DaCompany Report #2003025778

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Optic Neuritis	Foreign	Neurontin			
3000 MG		Vision Blurred	Health	(Gabapentin)	PS		
(DAILY),			Professional				
				Amitriptyline			
				(Amitriptyline)	C		
				Clonidine			
				(Clonidine)	C		
				Metoprolol			
				(Metoprolol)	C		
				Promethazine			
				Hydrochloride			
				(Promethazine			
				Hydrochloride)	C		

Date:06/24/03ISR Number: 4135728-4Report Type:Expedited (15-DaCompany Report #2003025618

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Disorientation	Consumer	Neurontin			
Initial or Prolonged		Feeling Abnormal		(Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG QD PO		Cerebellar Syndrome	Foreign	Tenofovir Df	PS		ORAL
Initial or Prolonged 2.5 MG QD PO		Cerebral Atrophy	Health	Renitec	SS		ORAL
20 MG QD PO		Coordination Abnormal	Professional	Prozac	SS		ORAL
300 MG BID PO		Dysarthria	Other	Neurontin	SS		ORAL
2 BID PO		Pneumocystis Jiroveci		Kaletra	SS		ORAL
450 MG BID PO		Pneumonia		Valcyte	SS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Torsade De Pointes	Foreign Health Professional	Neurontin (Gabapentin) Enalapril Maleate (Enalapril Maleate) Atorvastatin Calcium (Atorvastatin Calcium)	PS C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/26/03ISR Number: 4137007-8Report Type:Expedited (15-DaCompany Report #2003025957

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fluid Retention	Foreign	Neurontin			
1800 MG		Glossitis	Health	(Gabapentin)	PS		
(DAILY)		Lichen Planus	Professional				
		Oral Mucosal Disorder		Fluoxetine			
		Paraesthesia		(Fluoxetine)	C		
		Rash		Levothyroxine			
				(Levothyroxine)	C		
				Multivitamins			
				(Ergocalciferol,			
				Ascorbic Acid, Folic			
				Acid, Thiamine			
				Hydrochloride,	C		
				Insulin Zinc			
				Suspension (Insulin			
				Zinc Suspension)	C		
				Insulin Spart			
				(Insulin Aspart)	C		

Date:06/26/03ISR Number: 4137048-0Report Type:Expedited (15-DaCompany Report #2003026090

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Respiratory Failure	Foreign	Neurontin			
Other			Health	(Gabapentin)	PS		ORAL
ORAL			Professional				
			Company				
			Representative				

Date:06/26/03ISR Number: 4137054-6Report Type:Expedited (15-DaCompany Report #2003026124

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Convulsion	Consumer	Neurontin (Tablets)			

Other Depressed Level Of (Gabapentin) PS
Consciousness

Date:06/26/03ISR Number: 4143396-0Report Type:Expedited (15-DaCompany Report #US-SHR-03-009149
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Pressure Abnormal Dehydration	Consumer	Betaseron (Interferon Beta-1b)	PS		
SUBCUTANEOUS	8 MIU, EVERY 2D,	Depressed Level Of Consciousness					
SUBCUTANEOUS		Somnolence		Lasix (Furosemide) Neurontin (Gabapentin)	SS SS		
SEE IMAGE				Provigil (Modafinil)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/27/03ISR Number: 4139326-8Report Type:Expedited (15-DaCompany Report #2003026291

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 TID), ORAL		Bacterial Infection Cardiac Flutter	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Sumatriptan Succinate (Sumatriptan Succinate)	C		
				Amitriptyline (Amitriptyline)	C		
				Nitrofurantoin (Nitrofurantoin)	C		
				Diazepam (Diazepam)	C		
				Oxaprozin (Oxaprozin)	C		
				Biselect (Hydrochlothiazide, Bisoprolol Fumarate)	C		

Date:06/30/03ISR Number: 4139490-0Report Type:Expedited (15-DaCompany Report #GB-JNJFOC-20030602504

Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1 IN 5 HOUR 1200 MG, 2 IN 1 DAY, ORAL		Back Pain Convulsion Medication Error Tic	Foreign Health Professional	Zydol (Tramadol Hydrochloride) Neurontin (Gabapentin)	PS SS		ORAL
				Piroxicam (Piroxicam)	C		
				Clonidine (Clonidine)	C		

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300, Other THREE TIMES DAILY), ORAL	Abdominal Pain Burning Sensation Cold Sweat Condition Aggravated Dizziness Joint Swelling Parkinson'S Disease	Consumer Health Professional	Neurontin (Gabapentin) Citalopram Hydrobromide (Citalopram Hydrobromide) Famotidine (Famotidine) Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Paracetamol (Paracetamol) Sinemet (Levodopa, Carbidopa)	PS C C C C C		ORAL

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Freedom Of Information (FOI) Report

Date:06/30/03ISR Number: 4140310-9Report Type:Expedited (15-DaCompany Report #2003017389
Age:46 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 3600 MG Other (1200, TID), ORAL	Infection Lung Disorder Pulmonary Oedema	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Quetiapine Fumarate (Quetiapine Fumarate)	C		
			Clonazepam (Clonazepam)	C		
			Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		

Date:06/30/03ISR Number: 4140311-0Report Type:Expedited (15-DaCompany Report #2003014454
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Facial Pain Trigeminal Neuralgia	Consumer	Neurontin (Gabapentin)	PS		

Date:07/01/03ISR Number: 4138753-2Report Type:Expedited (15-DaCompany Report #US-ROCHE-315823
Age:71 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Creatinine Renal Clearance Decreased Dry Mouth Dysphonia Tardive Dyskinesia Urine Arsenic Increased Urine Mercury Abnormal Weight Increased		Klonopin Neurontin Lamictal Diavan Vitamins	PS SS SS C C	Roche	ORAL ORAL ORAL ORAL ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depressed Level Of Consciousness	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (300, TID), ORAL		Dysphagia					
		Eye Disorder		Potassium (Potassium)	C		
		Lethargy		Propacet (Paracetamol, Dextropropoxyphene Napsilate)	C		
		Muscular Weakness		Captopril (Captopril)	C		
		Urinary Incontinence		Doxazosin (Doxazosin)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/01/03ISR Number: 4140597-2Report Type:Expedited (15-DaCompany Report #2003019626

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 20 MG Other (DAILY), ORAL	Atherosclerosis Blood Cholesterol Increased Blood Pressure Inadequately Controlled Blood Triglycerides Drug Ineffective Myocardial Infarction Pain In Extremity	Consumer	Lipitor (Atorvastatin)	PS		ORAL
			Neurontin (Gabapentin)			
			(Gabapentin)	SS		
			Fluvastatin Sodium (Fluvastatin Sodium)	SS		
			Ezetimibe (Ezetimibe)	SS		
			Benazepril Hydrochloride (Benazepril Hydrochloride)	C		
			Atenolol (Atenolol)	C		
			Hyzaar (Hydrochlorothiazide , Losartan Potassium)	C		
			Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
			Alprazolam (Alprazolam)	C		

Date:07/01/03ISR Number: 4140600-XReport Type:Expedited (15-DaCompany Report #2003026754

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Dyslexia Speech Disorder	Health Professional	Neurontin (Gabapentin)	PS		
			Oxycocet (Paracetamol, Oxycodone Hydrochloride)	SS		
			Alprazolam (Alprazolam)	SS		

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Condition Aggravated	Health	Neurontin			
Initial or Prolonged	Depression	Professional	(Gabapentin)	PS		ORAL
600 MG (BID),						
Other	Dialysis					
ORAL	Renal Failure Acute		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/02/03ISR Number: 4141483-4Report Type:Expedited (15-DaCompany Report #A-US2003-02718

Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia	Health	Tracleer (Bosentan)			
Hospitalization -		Coma	Professional	Tablet	PS		ORAL
62.5 MG, BID, Initial or Prolonged ORAL; 125 MG, BID		Confusional State	Distributor				
		Dysphagia					
		Haematocrit		Flolan (Epoprostenol			
		Hyperhidrosis		Sodium)	SS		
INTRAVENOUS	UNK, UNK,	Hypoglycaemia					
INTRAVENOUS		Hyponatraemia		Vioxx (Rofecoxib)	SS		
		Pneumonia Aspiration		Cardizem (Diltiazem			
		Pyrexia		Hydrochloride)	SS		
		Rash Erythematous		Neurontin			
		Rash Generalised		(Gabapentin)	SS		
		Respiratory Arrest		Prevacid			
		Thrombocytopenia		(Lansoprazole)	C		
		Weight Decreased		Aldactone			
				(Spironolactone)	C		
				Elavil			
				(Amitriptyline			
				Hydrochloride)	C		
				Celexa (Citalopram			
				Hydrobromide)	C		
				Oxycontin (Oxycodone			
				Hydrochloride)	C		
				Allegra			
				(Fexofenadine			
				Hydrochloride)	C		
				Lasix (Furosemide)	C		
				Coumadin (Warfarin			
				Sodium)	C		

Date:07/02/03ISR Number: 4141786-3Report Type:Expedited (15-DaCompany Report #2003027160

Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Death Other 2100 MG (DAILY), ORAL	Bronchopneumonia Sputum Purulent	Foreign Health Professional	Neurontin (Gabapentin)	PS	ORAL
120 MG (DAILY), ORAL			Ketogin (Ketobromide Hydrochloride, Dimethyl-3, 3-Diphenyl-1-Methyla llylamine Hcl)	SS	ORAL
225 MG (DAILY), ORAL			Ketobemidone (Ketobemidone)	SS	ORAL
			Glyceryl Trinitrate (Glyceryl Trinitrate)	C	
			Levothyroxine Sodium (Levothyroxine Sodium)	C	
			Propranolol Hydrochloride (Propranolol Hydrochloride)	C	
			Estrogens Conjugated		

Freedom Of Information (FOI) Report

(Estrogens
 Conjugated) C
 Flunitrazepam
 (Flunitrazepam) C
 Omeprazole Magnesium
 (Omeprazole
 Magnesium) C

Date:07/02/03ISR Number: 4141788-7Report Type:Expedited (15-DaCompany Report #2003027159
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcoholism Cardiomegaly	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
800 MG (DAILY), ORAL		Drug Abuser Toxicologic Test Abnormal	Professional				
150 MG (DAILY), PLACENTAL				Methadone Hydrochloride (Methadone Hydrochloride)	SS		
				Mepenzolate Bromide (Mepenzolate Bromide)	C		
				Paracetamol (Paracetamol)	C		
				Propiomazine Maleate (Propiomazine Maleate)	C		
				Levomepromazine (Levomepromazine)	C		
				Carisoprodol (Carisoprodol)	C		
				Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		

Outcome	PT
Other	Abnormal Behaviour
	Alopecia
	Aphasia
	Autism
	Cognitive Disorder
	Communication Disorder
	Crying
	Decreased Appetite
	Developmental Delay
	Drug Toxicity
	Encephalopathy
	Insomnia
	Nightmare
	Oral Intake Reduced
	Pseudodementia
	Psychomotor Hyperactivity
	Rett'S Disorder
	Screaming

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Speech Disorder Vomiting Weight Gain Poor	Report Source	Product	Role	Manufacturer	Route
SEE IMAGE			Consumer Health	Neurontin (Gabapentin)	PS		ORAL
750 MG			Professional	Depakote (Valproate Semisodium)	SS		
(DAILY),				Lamotrigine (Lamotrigine)	SS		
				Topiramate (Topiramate)	SS		
				Hydroxyzine Hydrochloride (Hydroxyzine Hydrochloride)	C		
				Phenobarbital (Phenobarbital)	C		
				Lactulose (Lactulose)	C		
				Senokot (Senna Fruit)	C		
				Ryna-12s (Chlorphenamine, Pseudoephedrine Hydrochloride)	C		

Date:07/02/03ISR Number: 4141791-7Report Type:Expedited (15-DaCompany Report #2003027172

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Musculoskeletal Stiffness Nervous System Disorder	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (BID),		Pain In Extremity					
ORAL		Paraesthesia Paralysis					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SUBCUTANEOUS	8 MIU, EVERY	Apnoea Bradycardia Coronary Artery Occlusion	Consumer Health Professional	Betaseron(Interferon Beta - 1b) Injection	PS		
2D, SUBCUTANEOUS		Oedema Peripheral		Klonopin(Clonazepam)	SS		ORAL
0.5 MG, IN THE MORNING, ORAL; SEE IMAGE		Somnolence		Neurontin (Gabapentin)	SS		ORAL
300 MG, IN THE MORNING, ORAL; SEE IMAGE				Efexor - Slow Release (Venlafaxine Hydrochloride) Levothyroxine (Levothyroxine)	C C		

Freedom Of Information (FOI) Report

Potassium Chloride	C
Zocor "Msd" (Simvastatin)	C
Evista (Raloxifene Hydrochloride)	C
Baclofen	C
Morphine (Morphine)	C
Nexium (Esomeprazole)	C
Aminopyridine (Fampridine)	C
Nasacort Aq (Triamcinolone Acetonide)	C

Date:07/02/03ISR Number: 4175585-3Report Type:Periodic
Age:69 YR Gender:Male I/FU:I

Company Report #EMADSS2003002954

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged TRANSDERMAL	100 MCG/H, 1	Confusional State Pneumonia	Foreign Health Professional	Durogesic (100 Mcg/Hr Patch) (Fentanyl)	PS		
IN 72 HOUR(S), TRANSD				Finlepsin (Carbamazepine)	SS		ORAL
600 MG, 2 IN 1 DAILY, ORAL				Neurontin (Gabapentin)	SS		ORAL
300 MG, 3 IN 1 DAILY, ORAL				Saroten (Amitriptyline Hydrochloride)	SS		ORAL
10 MG, 3 IN 1							

DAILY, ORAL

Lasix (Furosemide) C
Novalgin (Metamizole Sodium) C
Rytmonorm (Propafenone Hydrochloride) C

Date:07/02/03ISR Number: 4181027-4Report Type:Periodic
Age:59 YR Gender:Male I/FU:I

Company Report #NSADSS2002039704

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged TRANSDERMAL TRANSD	Confusional State	Health Professional	Duragesic (Patch) (Fentanyl) Neurontin (Gabapentin) Soma (Carisoprodol) Doxepin Lidoderm Patch Testosterone Zocor (Simvastatin) Bextra Ultram (Tramadol Hydrochloride) Roxanol (Morphine	PS SS C C C C C C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Sulfate) C
 Risperdal
 (Risperidone) C

Date:07/02/03ISR Number: 4181648-9Report Type:Periodic
 Age:77 YR Gender:Male I/FU:I

Company Report #EMADSS2001007696

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	25 MCG/H, 1	Pneumonitis Somnolence	Foreign Health	Durogesic (25 Mcg/Hr Patch) (Fentanyl)	PS		
TRANSDERMAL Required IN 3 DAY(S), Intervention to TRANSD Prevent Permanent Impairment/Damage		Therapeutic Response Increased	Professional	Neurontin (Gabapentin) Celebrex (Celecoxib)	SS C		

Date:07/02/03ISR Number: 4181791-4Report Type:Periodic
 Age:35 YR Gender:Male I/FU:I

Company Report #EMADSS2003000044

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	100 MCG,	Coma Hypoaesthesia Paraesthesia	Foreign Health Professional	Duragesic (100 Mcg/Hr Patch) (Fentanyl)	PS		
TRANSDERMAL TRANSD 2700 MG, DAILY, ORAL		Therapeutic Response Increased		Gabapentin (Gabapentin)	SS		ORAL
				Trandolapril (Trandolapril) Diazepam (Diazepam) Etodolac (Etodolac) Beclomethasone (Beclometasone) Salbutamol (Salbutamol)	C C C C C C		

Zydol (Tablet)
(Tramadol
Hydrochloride) C
Pethidine
(Pethidine) C

Date:07/02/03ISR Number: 4182719-3Report Type:Periodic Company Report #NSADSS2002024247
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Confusional State Somnolence	Foreign Health	Duragesic (Patch) (Fentanyl)	PS		
TRANSDERMAL	225 MCG/H, 1	Professional				

IN 72

HOUR(S),

TRANSD

50 MG, 1 IN 1

NIGHT(S),

ORAL

Elavil
(Amitriptyline
Hydrochloride) SS ORAL

75 MG, 1 IN 1

NIGHT(S),

Elavil
(Amitriptyline
Hydrochloride) SS

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

OTHER

1800; 2800;

900, DAILY,

ORAL

Neurontin
(Gabapentin) SS ORAL

Metoclopramide
(Metoclopramide) C
Dilaudid
(Hydromorphone
Hydrochloride) C
Vioxx (Rofecoxib) C
Pantoloc
(Pantoprazole
Sodium) C

Date:07/03/03ISR Number: 4141452-4Report Type:Direct
Age:73 YR Gender:Male I/FU:I

Company Report #CTU 197224

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Neurontin 100 Mg	PS		ORAL
Other		Ear Discomfort					
300 MG DAY		Ear Disorder					
ORAL		Ear Infection					
		Ear Pain					
		Hearing Impaired					
		Otitis Externa					

Date:07/03/03ISR Number: 4142690-7Report Type:Expedited (15-DaCompany Report #2003027161
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Dysgeusia					
ORAL		Headache		Diazepam	C		
		Increased Appetite		Oxycocet			

Stomach Discomfort
Stress
Urticaria
Weight Increased

(Paracetamol,
Oxycodone
Hydrochloride) C

Date:07/03/03ISR Number: 4142692-0Report Type:Expedited (15-DaCompany Report #2003027063
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased Body Height Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG		Compression Fracture					
(DAILY) ORAL		Weight Increased		Celecoxib (Celecoxib)	SS		ORAL
200 MG							
(DAILY) ORAL				Levothyroxine Sodium	C		
				Alendronate Sodium	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/07/03ISR Number: 4142191-6Report Type:Expedited (15-DaCompany Report #FR-ROCHE-341229

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Cerebellar Syndrome	Consumer	Valcyte	PS	Roche	ORAL
Initial or Prolonged		Dysarthria		Viread	SS		ORAL
		Nuclear Magnetic		Kaletra	SS		ORAL
		Resonance Imaging		Neurontin	SS		ORAL
		Abnormal		Prozac	SS		ORAL
				Renitec	SS		ORAL
				Aspegic	SS		ORAL
UNKNOWN				Levothyrox	C		

Date:07/07/03ISR Number: 4144070-7Report Type:Expedited (15-DaCompany Report #2003027449

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia	Consumer	Neurontin			
Other		Dizziness		(Gabapentin)	PS		
		Dysarthria		Methadone			
(TID)		Hyperventilation		(Methadone)	SS		
		Somnolence		Clonazepam			
				(Clonazepam)	C		
				Paroxetine			
				Hydrochloride			
				(Paroxetine			
				Hydrochloride)	C		
				Digoxin (Digoxin)	C		
				Metoprolol Tartrate			
				(Metoprolol			
				Tartrate)	C		
				Rabeprazole			
				(Rabeprazole)	C		
				Temazepam			
				(Temazepam)	C		
				Lansoprazole			
				(Lansoprazole)	C		
				Simvastatin			
				(Simvastatin)	C		
				All Other			
				Therapeutic Products	C		

Salbutamol
(Salbutamol)

C

Date:07/07/03ISR Number: 4144102-6Report Type:Expedited (15-DaCompany Report #2003015585
Age:43 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 2700 MG, TID, Initial or Prolonged ORAL Disability Other	Cardiomyopathy Infectious Mononucleosis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Desipramine	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/07/03ISR Number: 4144157-9Report Type:Expedited (15-DaCompany Report #2002050043

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Distension	Consumer	Neurontin			
(TWICE		Adrenal Adenoma	Health	(Gabapentin)	PS		ORAL
DAILY), ORAL		Asthenia	Professional				
ORAL		Blood Arsenic Increased		Lamotrigine			
		Blood Mercury Abnormal		(Lamotrigine)	SS		ORAL
		Creatinine Renal		Clonazepam			
		Clearance Decreased		(Clonazepam)	SS		
		Dizziness		Nifedipine			
		Dry Mouth		(Nifedipine)	C		
		Dyspepsia					
		Dyspnoea					
		Headache					
		Helicobacter Infection					
		Hunger					
		Hypertension					
		Malaise					
		Muscle Spasms					
		Nausea					
		Nervousness					
		Oesophageal Spasm					
		Pharmaceutical Product					
		Complaint					
		Pollakiuria					
		Respiratory Disorder					
		Sensation Of Heaviness					
		Tardive Dyskinesia					
		Throat Tightness					
		Tracheal Obstruction					
		Weight Increased					

Date:07/07/03ISR Number: 4144202-0Report Type:Expedited (15-DaCompany Report #2003027534

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser	Health	Neurontin (Tablets)			
			Professional	(Gabapentin)	PS		

Date:07/07/03ISR Number: 4144204-4Report Type:Expedited (15-DaCompany Report #2003027533

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		

Date:07/07/03ISR Number: 4144207-XReport Type:Expedited (15-DaCompany Report #2003027532

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/07/03ISR Number: 4144881-8Report Type:Expedited (15-DaCompany Report #2003027568

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG Other (DAILY),		Dry Mouth Dyspnoea Myalgia Neuralgia	Foreign Consumer	Neurontin (Gabapentin) Oral Antidiabetics	PS C		

Date:07/08/03ISR Number: 4144987-3Report Type:Expedited (15-DaCompany Report #2003027915

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Anticonvulsant Toxicity Autoimmune Disorder	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Epistaxis Hypersensitivity Muscle Disorder		Phenytoin Suspension (Phenytoin) (Phenytoin Sodium)	SS		ORAL

Date:07/08/03ISR Number: 4144993-9Report Type:Expedited (15-DaCompany Report #2003018752

Age:74 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Alanine Aminotransferase Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Aspartate Aminotransferase Increased Cytolytic Hepatitis Depressed Level Of Consciousness	Professional Company Representative	Paroxetine (Paroxetine) Hydromorphone (Hydromorphone) Mianserin (Mianserin) Morphine Sulfate (Morphine Sulfate)	C C C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcoholism Cardiomegaly	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
800 MG (DAILY), ORAL		Drug Abuser	Professional				
				Methadone Hydrochloride (Methadone Hydrochloride)	SS		ORAL
150 MG (DAILY), ORAL							
				Mepenzolate Bromide (Mepenzolate Bromide)	C		
				Paracetamol (Paracetamol)	C		
				Propiomazine Maleate (Propiomazine Maleate)	C		
				Levomepromazine (Levomepromazine)	C		
				Carisoprodol (Carisoprodol)	C		
				Venlafaxine			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride
(Venlafaxine
Hydrochloride) C

Date:07/09/03ISR Number: 4144419-5Report Type:Direct
Age:88 YR Gender:Female I/FU:I

Company Report #CTU 197507

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300MG PO QID		Catatonia Muscle Rigidity Mydriasis Tremor		Neurontin 300 Mg Po Qid	PS		ORAL
				Metoprolol	C		
				Novolin	C		
				Plavix	C		
				Protonix	C		
				Celexa	C		
				Compazine	C		
				Seroquel	C		

Date:07/09/03ISR Number: 4145991-1Report Type:Expedited (15-DaCompany Report #2003027782
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 3600 MG (DAILY), ORAL		Conversion Disorder Drug Tolerance Economic Problem Pain Paraesthesia Post Procedural Pain	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Fentanyl (Fentanyl)	C		
				Ramipril (Ramipril)	C		
				Allopurinol (Allopurinol)	C		
				Acetazolamide (Acetazolamide)	C		
				Sodium Bicarbonate (Sodium Bicarbonate)	C		
				Colchicine (Colchicine)	C		
				Tolterodine L-Tartrate (Tolterodine L-Tartrate)	C		

Calcium (Calcium)	C
Hydrochlorothiazide	
(Hydrochlorothiazide	
)	C
Pethidine	
Hydrochloride	
(Pethidine	
Hydrochloride)	C
Montelukast Sodium	
(Montelukast Sodium)	C
Ranitidine	
Hydrochloride	
(Ranitidine	
Hydrochloride)	C
Cetirizine	
Hydrochloride	
(Cetirizine	
Hydrochloride)	C
Diphenhydramine	
Hydrochloride	
(Diphenhydramine	

Freedom Of Information (FOI) Report

Hydrochloride) C
 Salmeterol Xinafoate
 (Salmeterol
 Xinafoate) C
 Fluticasone
 Propionate
 (Fluticasone
 Propionate) C
 Salbutamol
 (Salbutamol) C
 Epinephrine
 (Epinephrine) C
 Prednisone
 (Prednisone) C
 Vicodin
 (Paracetamol,
 Hydrococone
 Bitartrate) C

Date:07/09/03ISR Number: 4146420-4Report Type:Expedited (15-DaCompany Report #2003024137

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Asthenia	Consumer	Neurontin (Gabapentin)	PS		
Other		Balance Disorder Hypoaesthesia		Ropinirole Hydrochloride (Ropinirole Hydrochloride)	SS		ORAL
	(DAILY), ORAL			Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
				Hyoscyamine (Hyoscyamine)	C		
				Tizanidine Hydrochloride (Tizanidine Hydrochloride)	C		
				Diclofenac Sodium (Diclofenac Sodium)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Anxiety	Consumer	Neurontin			
Initial or Prolonged	Circulatory Collapse		(Gabapentin)	PS		
Disability	Dysgeusia					
Other	Emotional Distress					
	General Physical Health					
	Deterioration					
	Impaired Self-Care					
	Pneumonia					
	Syncope					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/10/03ISR Number: 4147070-6Report Type:Expedited (15-DaCompany Report #2003027916
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Foreign	Neurontin (Tablets)			
2400 MG		Muscle Spasms	Health	(Gabapentin)	PS		ORAL
(QID), ORAL			Professional				

Date:07/11/03ISR Number: 4147770-8Report Type:Expedited (15-DaCompany Report #2003021716
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Economic Problem	Consumer	Neurontin			
UNKNOWN		No Adverse Drug Effect	Health	(Gapaentin)	PS		
(UNKNOWN),			Professional				
UNKNOWN							

Date:07/14/03ISR Number: 4146494-0Report Type:Direct Company Report #CTU 197818
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Aggression		Neurontin 300mg			
Hospitalization -		Injury		Parke-Davis	PS	Parke-Davis	ORAL
SEE IMAGE							
Initial or Prolonged		Intentional Self-Injury					
Other		Psychiatric Symptom					
		Suicide Attempt					

Date:07/14/03ISR Number: 4146504-0Report Type:Direct Company Report #CTU 197830
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 800MG T.I.D. Required ORAL	Tooth Loss	Neurontin 800mg	PS	ORAL
Intervention to Prevent Permanent Impairment/Damage		Methadone Plaquenil Prednisone Xanax Imitrex	C C C C C	

Date:07/14/03ISR Number: 4148241-5Report Type:Expedited (15-DaCompany Report #2003025618
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour Disorientation	Consumer	Neurontin (Gabapentin)	PS		

Date:07/14/03ISR Number: 4148435-9Report Type:Expedited (15-DaCompany Report #2003028825
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Blood Cholesterol Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Blood Triglycerides Euphoric Mood Foot Fracture Headache Weight Increased		Pravastatin Sodium Tamoxifen Pantoprazole Fluoxetine Hydrochloride	C C C C C		

Freedom Of Information (FOI) Report

Fioricet W/Codeine
 (Codeine Phosphate,
 Caffeine,
 Butalbital,
 Paracetamol) C

Date:07/14/03ISR Number: 4148438-4Report Type:Expedited (15-DaCompany Report #2003022137
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurotin (Gabapentin)	PS		
400 MG		Drug Level Decreased					
		Medication Error		Dilantin Suspension (Phenytoin Sodium)	SS		
		Pharmaceutical Product Complaint					

Date:07/14/03ISR Number: 4148439-6Report Type:Expedited (15-DaCompany Report #2003028324
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged BID, ORAL		Adrenal Disorder Anaphylactic Reaction	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Blindness		Calcitriol	C		
		Diplopia		Simvastatin	C		
		Hypothalamo-Pituitary Disorders		Trimethoprim	C		
		Intracranial Pressure Increased		Bupropion Hydrochloride	C		
		Migraine		Levothyroxine Sodium	C		
		Optic Disc Disorder		Oxaprozin	C		
		Pain		Alendronate Sodium	C		
		Thyroid Disorder		Metaxalone	C		
		Vision Blurred		Sibutramine	C		
		Visual Disturbance		Hydrochloride	C		
				Potassium	C		
				Nortriptyline	C		
				Lansoprazole	C		

Date:07/14/03ISR Number: 4149274-5Report Type:Expedited (15-DaCompany Report #2003028852

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Stevens-Johnson Syndrome Toxic Epidermal Necrolysis	Foreign Health Professional	Gabapentin (Gabapentin) Valproate Sodium (Valproate Sodium) Clonazepam (Clonazepam)	PS C C		ORAL

Date:07/14/03ISR Number: 4149278-2Report Type:Expedited (15-DaCompany Report #2003026277

Age:71 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Coordination Abnormal Drug Level Increased Dyskinesia Myoclonus Restlessness

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Freedom Of Information (FOI) Report

Tongue Disorder

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
600 MG	(DAILY), ORAL	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Simvastatin (Simvastatin)	C		
			Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
			Metoprolol Succinate (Metoprolol Succinate)	C		
			Isosorbide Dinitrate (Isosorbide Dinitrate)	C		
			Molsidomine (Molsidomine)	C		
			Calcium Acetate (Calcium Acetate)	C		
			Sevelamer Hydrochloride (Sevelamer Hydrochloride)	C		
			Calcitriol (Calcitriol)	C		
			Colecalciferol (Colecalciferol)	C		
			Human Mixtard (Insulin Human, Insulin Human Injection, Isophane)	C		
			Glyceryl Trinitrate (Glyceryl Trinitrate)	C		
			Metamizole Sodium (Metamizole Sodium)	C		
			Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
			Sodium Bicarbonate (Sodium Bicarbonate)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Abortion Spontaneous	Foreign	Neurontin			
Initial or Prolonged	Complications Of Maternal	Health	(Gabapentin)	PS		
1800 MG (600						
TID)	Exposure To Therapeutic	Professional				
	Drugs		Folic Acid (Folic			
	Maternal Drugs Affecting		Acid)	C		
	Foetus		Phenobarbital			
			(Phenobarbital)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/14/03ISR Number: 4149336-2Report Type:Expedited (15-DaCompany Report #2003028326

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coma	Foreign	Neurontin			
TID		Pulse Abnormal	Health	(Gabapentin)	PS		
			Professional	Capozide			
			Company	(Hydrochlorothiazide			
			Representative	, Captopril)	C		
				Di-Gesic			
				(Paracetamol,			
				Dextropropoxyphene)	C		
				Caffeine	C		

Date:07/15/03ISR Number: 4148529-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -							
Initial or Prolonged		Brain Neoplasm	Consumer	Neurontin			
400 MG (BID),		Cardiac Arrest	Health	(Gabapentin)	PS		ORAL
Disability		Conversion Disorder	Professional				
ORAL			Company	Valproate Sodium			
Other		Dizziness	Representative	(Valproate Sodium)	SS		
		Dysphemia		Buspirone			
		Fall		Hydrochloride			
		Implant Site Reaction		(Buspirone			
		Pain In Extremity		Hydrochloride)	C		
		Scar		Fluoxetine			
		Tenderness		Hydrochloride			
		Tremor		(Fluoxetine			
				Hydrochloride)	C		
				Omeprazole			
				(Omeprazole)	C		
				Capsaicin			
				(Capsaicin)	C		

Date:07/15/03ISR Number: 4149260-5Report Type:Expedited (15-DaCompany Report #2003018325

Age:46 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Infertility Male	Health Professional	Neurontin (Gabapentin)	PS		ORAL
(TID), ORAL				Rofecoxib (Rofecoxib)	C		
				Zolpidem Tartrate (Zolpidem Tartrate)	C		
				Vicodin (Paracetamol, Hydrocodone Bitartrate)	C		
				Omeprazole (Omeprazole)	C		
				Cyclobenzaprine (Cyclobenzaprine)	C		
				Salbutamil (Salbutamol)	C		
				Amitriptyline (Amitriptyline)	C		
				Carisoprodol (Carisoprodol)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/15/03ISR Number: 4150123-XReport Type:Expedited (15-DaCompany Report #2002056560

Age:80 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Arrhythmia Atrial Thrombosis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL	Drug Toxicity	Professional	Clobazam (Clobazam)	SS		ORAL
	Encephalopathy Overdose Rash Maculo-Papular Somnolence		Valsartan Hydrochlorothiazide (Hydrochlorothiazide , Valsartan) Fluindione (Fluindione)	C C		

Date:07/15/03ISR Number: 4150124-1Report Type:Expedited (15-DaCompany Report #2003028905

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Bone Marrow Depression Bone Marrow Transplant	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:07/16/03ISR Number: 4150575-5Report Type:Expedited (15-DaCompany Report #2003029269

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Drug Ineffective Drug Level Below Therapeutic Malignant Neoplasm Of Spinal Cord	Consumer	Neurontin (Gabapentin)	PS		

Date:07/16/03ISR Number: 4151002-4Report Type:Expedited (15-DaCompany Report #2003025422

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1000 MCG		Bradycardia	Consumer	Tikosyn (Dofetilide)	PS		ORAL
Initial or Prolonged (BID), ORAL		Fatigue	Health				
Other		Somnolence	Professional	Neurontin (Gabapentin)	SS		ORAL
ORAL				Tacrolimus (Tacrolimus)	SS		ORAL
ORAL				Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		ORAL
ORAL				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		
1 GRAM (BID),				Mycophenolate Mofetil (Mycophenolate Mofetil)	SS		ORAL
ORAL				Goserelin (Goserelin)	C		
				Acetylsalicylic Acid (Acetylsalicylic			

Freedom Of Information (FOI) Report

Acid) C
 Warfarin (Warfarin) C
 Goserelin
 (Goserelin) C
 Oxycocet
 (Paracetamol,
 Oxycodone
 Hydrochloride) C

Date:07/16/03ISR Number: 4151027-9Report Type:Expedited (15-DaCompany Report #2003027172
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (BID), ORAL		Feeling Abnormal Joint Stiffness Pain In Extremity Paraesthesia Paralysis	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:07/16/03ISR Number: 4151287-4Report Type:Expedited (15-DaCompany Report #US-SHR-03-009149
 Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - SUBCUTANEOUS Initial or Prolonged 2D, SUBCUTANEOUS 40 MG 1X/DAY 400 MG, 3XDAY	8 MIU, EVERY	Blood Pressure Decreased Dehydration Feeling Cold Loss Of Consciousness Somnolence Sudden Onset Of Sleep Treatment Noncompliance	Consumer	Betaseron Lasix (Furosemide) Neurontin (Gabapentin) Provigil (Modafinil) Macrochantin (Nitrofurantoin) Mirapex	PS SS SS C C		

"Pharmacia-Upjohn"		
(Pramipexole		
Hydrochloride)	C	Pharmacia-Upjohn
Bextra (Bucindolol		
Hydrochloride)	C	
Wellbutrin		
(Amfebutamone		
Hydrochloride)	C	
Zanaflex (Tizanidine		
Hydrochloride)	C	
Sinemet (Carbidopa,		
Levodopa)	C	
Potassium		
(Potassium)	C	
Baclofen	C	
Percocet (Oxycodone		
Hydrochloride)	C	

Date:07/17/03ISR Number: 4150464-6Report Type:Expedited (15-DaCompany Report #PHNU2003DE01380
Age:70 YR Gender:Male I/FU:F

Outcome PT
Hospitalization - Drug Interaction
Initial or Prolonged Fatigue

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pain In Extremity Weight Increased	Report Source	Product	Role	Manufacturer	Route
50 MG/DAY, ORAL			Foreign Health Professional	Lioresal (Baclofen) Tablet, 10 Mg	PS		ORAL
1800 MG/DAY,			Other	Neurontin (Gabapentin)	SS		
				Saroten "Bayer Vital" (Amitriptyline Hydrochloride)	SS	Bayer Vital	
				L-Thyroxin "Henning Berlin"	C		
				Candesartan (Candesartan)	C		
				Oxybutynin (Oxybutynin)	C		

Date:07/17/03ISR Number: 4151500-3Report Type:Expedited (15-DaCompany Report #2003167945US
Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Life-Threatening ORAL Other ORAL		Myocardial Infarction	Study Consumer	Celebrex (Celecoxib) Capsule	PS		ORAL
				Neurontin(Gabapentin)	SS		ORAL
				Procardia(Nifedipine)	SS		

Date:07/18/03ISR Number: 4151670-7Report Type:Expedited (15-DaCompany Report #2003029347
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dyskinesia		Gabapentin			

Other Left Ventricular Failure (Gabapentin) PS ORAL
 1500 MG (FIVE
 TIMES DAILY),
 ORAL

Amitriptyline
 (Amitriptyline) SS
 Perindopril
 (Perindopril) C
 Spironolactone
 (Spironolactone) C
 Furosemide
 (Furosemide) C
 Paracetamol
 (Paracetamol) C

Date:07/18/03ISR Number: 4151671-9Report Type:Expedited (15-DaCompany Report #2003029157
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State Fall Vertigo	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/18/03ISR Number: 4151677-XReport Type:Expedited (15-DaCompany Report #2003030105
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hepatic Enzyme Increased	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:07/18/03ISR Number: 4151784-1Report Type:Expedited (15-DaCompany Report #2003028998
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (TID), ORAL		Circulatory Collapse Epilepsy	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Insomnia		Lamotrigine (Lamotrigine)	C		
		Irritability					

Date:07/18/03ISR Number: 4151790-7Report Type:Expedited (15-DaCompany Report #2003029349
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Abortion Spontaneous Complications Of Maternal	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
		Exposure To Therapeutic Drugs Maternal Drugs Affecting Foetus					

Date:07/18/03ISR Number: 4152539-4Report Type:Expedited (15-DaCompany Report #2003030062
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amputation Diabetes Mellitus Surgery	Consumer	Lipitor (Atorvastatin) Neurontin (Gabapentin) (Gabapentin) Cilostazol (Cilostazol)	PS SS C		

Date:07/18/03ISR Number: 4152626-0Report Type:Expedited (15-DaCompany Report #2003029575
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death							
Hospitalization - Initial or Prolonged		Gastrointestinal Tube Insertion Haematocrit Decreased Pneumonia Aspiration Pulmonary Hypertension Thrombocytopenia	Consumer	Neurontin (Gabapentin) Diltiazem Hydrochloride (Diltiazem Hydrochloride) Rofecoxib (Rofecoxib) Lansoprazole (Lansoprazole) Spironolactone	PS SS SS C		

Freedom Of Information (FOI) Report

(Spironolactone)	C
Amitriptyline	
(Amitriptyline)	C
Citalopram	
Hydrobromide	
(Citalopram	
Hydrobromide)	C
Oxycodone	
Hydrochloride	
(Oxycodone	
Hydrochloride)	C
Fexofenadine	
Hydrochloride	
(Fexofenadine	
Hydrochloride)	C
Furosemide	
(Furosemide)	C
Warfarin (Warfarin)	C

Date:07/18/03ISR Number: 4154062-XReport Type:Expedited (15-DaCompany Report #2003025875

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abnormal Behaviour
Initial or Prolonged	Aggression
Disability	Angina Unstable
Other	Blood Thyroid Stimulating
	Hormone Decreased
	Carotid Artery Atheroma
	Carotid Bruit
	Cerebral Haematoma
	Colonic Polyp
	Coma
	Confusional State
	Coronary Artery Stenosis
	Cough
	Depression
	Diastolic Dysfunction
	Difficulty In Walking
	Dilatation Atrial
	Dizziness
	Encephalomalacia
	Fall
	Folate Deficiency
	Gait Disturbance
	Haemorrhage

Haemorrhage Intracranial
Haemorrhoidal Haemorrhage
Head Injury
Heart Rate Increased
Hypothyroidism
International Normalised
Ratio Increased
Lung Infiltration
Medication Error
Mental Status Changes
Mitral Valve Incompetence
Nuclear Magnetic
Resonance Imaging Brain
Abnormal
Obesity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
	Oedema Peripheral Orthostatic Hypotension Overdose					
(DAILY), ORAL	Pain Pneumonia	Consumer Health	Lipitor (Atorvastatin)	PS		ORAL
	Prothrombin Time Prolonged	Professional	Gabapentin (Gabapentin)	SS		
	Pulmonary Granuloma		Fentanyl (Fentanyl)	SS		
	Pyrexia		Warfarin Sodium (Warfarin Sodium)	C		
	Respiratory Rate Increased		Lansoprazole (Lansoprazole)	C		
	Road Traffic Accident		Venlafaxine Hydrochloride			
	Sedation		(Venlafaxine Hydrochloride)	C		
	Sleep Apnoea Syndrome		Potassium Chloride (Potassium Chloride)	C		
	Subdural Haematoma		All Other			
	Tricuspid Valve Incompetence		Therapeutic Products	C		
	Venous Insufficiency		Montelukast Sodium (Montelukast Sodium)	C		
	Ventricular Hypertrophy		Vicodin (Paracetamol, Hydrocodone Bitartrate)	C		
	Visual Disturbance		Lorazepam (Lorazepam)	C		
	Vitamin B12 Deficiency		Novothyral (Levothyroxine Sodium, Liothyronine Sodium)	C		
	Weight Increased		Tocopherol (Tocopherol)	C		
			Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		
			Cyclobenzaprine Hydrochloride			

(Cyclobenzaprine
Hydrochloride) C
Vicoprofen
(Hydrocodone
Bitartrate,
Ibuprofen) C

Date:07/21/03ISR Number: 4150970-4Report Type:Expedited (15-DaCompany Report #WAES 0307USA01625
Age:57 YR Gender:Male I/FU:I

Outcome PT
Death Coma
Hospitalization - Confusional State
Initial or Prolonged Dyskinesia Oesophageal
Other Hypoglycaemia

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Disability	Condition Aggravated	Consumer	Neurontin	
Other	Fatigue		(Gabapentin)	PS
	Headache		Primidone	
	Tremor		(Primidone)	C
			Nadolol (Nadolol)	C
			All Other	
			Therapeutic Products	C

Date:07/21/03ISR Number: 4152892-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain Lower
Initial or Prolonged	Abdominal Pain Upper
Disability	Abdominal Tenderness
Other	Acrochordon
	Albumin Globulin Ratio
	Anticonvulsant Drug Level
	Above Therapeutic
	Anxiety
	Blood Cholesterol
	Increased

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Pain Of Skin
Scar
Speech Disorder
Syncope
Tension
Toe Deformity
Tremor

Date:07/22/03ISR Number: 4153884-9Report Type:Expedited (15-DaCompany Report #2003030105
Age:51 YR Gender:Male I/FU:F

Outcome	PT	Report Source
Death	Hepatic Enzyme Increased	Foreign Health Professional Company

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Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
		Neurontin (Gabapentin)	PS		
		Metformin Hydrochloride (Metformin Hydrochloride)	C		
		All Other Therapeutic Products	C		

Date:07/22/03ISR Number: 4153887-4Report Type:Expedited (15-DaCompany Report #2003024694
Age:15 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Tremor	Foreign Consumer	Neurontin (Gabapentin)	PS		
400 MG			Health				
(DAILY)			Professional				

Date:07/22/03ISR Number: 4153969-7Report Type:Expedited (15-DaCompany Report #2003018310
Age:80 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Adhesion Facial Nerve Disorder	Foreign Consumer	Gabapentin (Tablets) (Gabapentin)	PS		ORAL
300 MG							
(DAILY), ORAL				All Other Therapeutic Products	C		

Date:07/22/03ISR Number: 4153971-5Report Type:Expedited (15-DaCompany Report #2003029580
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		General Physical Health	Foreign	Gabapentin			
Hospitalization -		Deterioration	Consumer	(Gabapentin)	PS		ORAL
ORAL							
Initial or Prolonged		Lymphoma					

Date:07/23/03ISR Number: 4152310-3Report Type:Direct Company Report #CTU 198469
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Movement Disorder		Neurontin 100 Mg			
1-4 CAPSULES		Muscle Twitching		Capsules	PS	Parke Davi	
3T , A DAY							
AND EVERY							
BEDTIME - AS							
NEEDED							

Date:07/23/03ISR Number: 4155238-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
 Age: Gender:Male I/FU:F

Outcome
 Hospitalization -
 Initial or Prolonged

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Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Difficulty In Walking	Foreign	Neurontin			
400 MG DAILY		Foot Fracture	Consumer	(Gabapentin)	PS		
		Muscular Weakness		Pravastatin Sodium			
		Thrombosis		(Pravastatin Sodium)	SS		
				Cortisone	C		
				Warfarin	C		

Age:63 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Anaemia
Initial or Prolonged	Dermatitis Exfoliative
	Fungal Infection
	Nausea
	Oral Intake Reduced

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Photosensitivity Reaction Thrombocytopenia	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Foreign Health Professional	Zyvoxid(Linezolid) Tablet	PS		ORAL
600 MG, BID,							
ORAL			Other	Neurontin (Gabapentin)	SS		ORAL
300 MG, BID,							
ORAL				Doliprane(Paracetamol)	SS		ORAL
500 MG, ORAL							
1 DF, BID,				Clobazam (Clobazam)	SS		ORAL
ORAL							
400 MG, BID,				Peflacine(Pefloxacin Mesilate)	SS		ORAL
ORAL							
				Domperidone (Domperidone)	C		
				Cetirizine (Cetirizine)	C		
				Amphotericin B	C		

Date:07/23/03ISR Number: 4155700-8Report Type:Expedited (15-DaCompany Report #2003029760
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Drug Effect Decreased Pharmaceutical Product					
2400 MG ORAL		Complaint		Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
				Valsartan			

(Valsartan) C
 Loratadine C
 (Loratadine) C
 Levothyroxine Sodium C
 (Levothyroxine Sodium) C
 Calcium (Calcium) C
 Glucosamine C
 (Glucosamine) C
 Vitamins C
 Acetylsalicylic Acid C
 (Acetylsalicylic Acid) C

Date:07/23/03ISR Number: 4155702-1Report Type:Expedited (15-DaCompany Report #2003029798
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Eye Haemorrhage Optic Ischaemic Neuropathy	Health Professional	Neurontin (Gabapentin)	PS		ORAL
2000 MG BID, ORAL				Trazodone (Trazodone) Methylphenidate (Methylphenidate) Topiramate	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Topiramate) C

Date:07/24/03ISR Number: 4156108-1Report Type:Expedited (15-DaCompany Report #2003030526

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Aggression Bipolar Disorder Condition Aggravated Weight Decreased	Health Professional	Neurontin (Gabapentin)	PS		

Date:07/24/03ISR Number: 4156435-8Report Type:Expedited (15-DaCompany Report #2003019414

Age:21 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 2700 MG (900, TID), ORAL	Drug Interaction Irritable Bowel Syndrome Jaw Disorder	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
50 MG (QD), ORAL	Muscle Spasms Muscle Twitching Speech Disorder		Mirtazapine (Mirtazapine)	SS		ORAL
150, Q5H ORAL	Tremor		Tramadol (Tramadol)	SS		ORAL
			Hydroxychloroquine Phosphate (Hydroxychloroquine Phosphate) Celecoxib Prednisone (Prednisone) Multivitamins And Iron Calcium (Calcium) Cilest (Ethinylestradiol , Norgestimate) Zolpidem Tartrate (Zolpidem Tartrate)	C C C C C C C		

Nifedipine	
(Nifedipine)	C
Glyceryl Trinitrate	
(Glyceryl	
Trinitrate)	C
Rabeprazole Sodium	
(Rabeprazole Sodium)	C
Etanercept	
(Etanercept)	C
Fish Oil (Fish Oil)	C
Chondroitin/Glucosamine (Glucosamine,	
Chondroitin)	C
Promethazine	
(Promethazine)	C
Prochlorperazine	
Edisylate	
(Prochlorperazine	
Edisylate)	C
Alprazolam	
(Alprazolam)	C
Acetylsalicylic Acid	
(Acetylsalicylic	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Acid) C

Date:07/24/03ISR Number: 4156503-0Report Type:Expedited (15-DaCompany Report #2003003636

Age:6 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Neurontin			
Other		Convulsion	Health	(Gabapentin)	PS		ORAL
900 MG (TID),		Crying	Professional				
ORAL		Dreamy State		Dilantin Suspension			
		Drug Level Below		(Phenytoin Sodium)	SS		ORAL
ORAL		Therapeutic		Prednisone			
		Electroencephalogram		(Prednisone)	SS		ORAL
ORAL		Abnormal		Sulfasalazine			
		Gingival Disorder		(Sulfasalazine)	C		
		Hair Growth Abnormal		Fish Oil (Fish Oil)	C		
		Increased Appetite		Folic Acid (Folic			
		Thinking Abnormal		Acid)	C		
				Mesalazine			
				(Mesalazine)	C		
				All Other			
				Therapeutic Products	C		

Date:07/24/03ISR Number: 4156514-5Report Type:Expedited (15-DaCompany Report #DSA_23002_2003

Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma	Health	Cardizem	PS		
Hospitalization -		Confusional State	Professional	Tracleer	SS		
125 MG BID							
Initial or Prolonged		Haematocrit Decreased		Tracleer	SS		ORAL
62.5 MG BID		Hypoglycaemia					
PO		Pneumonia		Flolan	SS		
		Pneumonia Aspiration		Vioxx	SS		
		Respiratory Arrest		Neurontin	SS		
		Thrombocytopenia		Prevacid	C		

White Blood Cell Count
Decreased

Aldactone C
Elavil C
Celexa C
Oxycontin C
Allegra C
Lasix C
Warfarin Sodium C

Date:07/24/03ISR Number: 4156596-0Report Type:Expedited (15-DaCompany Report #2002069713

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 100 MG (DAILY), ORAL	Constipation Dizziness Feeling Abnormal Nervousness Oral Intake Reduced Tinnitus	Consumer	Neurontin (Gabapentin)	PS		ORAL

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Date:07/24/03ISR Number: 4156787-9Report Type:Expedited (15-DaCompany Report #2003030344
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	600 MG (BID), ORAL	Cerebrovascular Accident Hypertension	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
				Glimepiride (Glimepiride)	C		
				Enalapril Maleate (Enalapril Maleate)	C		
				Digoxin (Digoxin)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Polyvitamin (Vitamins Nos)	C		

Date:07/24/03ISR Number: 4156790-9Report Type:Expedited (15-DaCompany Report #2003030521
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	600 MG (BID), ORAL	Visual Acuity Reduced	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
	1 DAY		Company Representative	All Other Therapeutic Products	C		

Date:07/25/03ISR Number: 4153944-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0417428A
Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia		Flolan	PS	Glaxosmithkline	
Hospitalization - 62.5MG Twice		Coma		Tracleer	SS		ORAL

Initial or Prolonged	Confusional State			
per day	53 DAY			
Other	Dysphagia	Vioxx	SS	
	Gastrointestinal Motility Disorder	Cardizem	SS	Glaxosmithkline
	Haematocrit Decreased	Neurontin	SS	
	Hyperhidrosis	Prevacid	C	
	Hypoglycaemia	Aldactone	C	
	Lung Infiltration	Elavil	C	Glaxosmithkline
	Pneumonia Aspiration	Celexa	C	
	Pyrexia	Oxycontin	C	
	Rash Erythematous	Allegra	C	
	Rash Generalised	Lasix	C	Glaxosmithkline
	Respiratory Arrest	Coumadin	C	Glaxosmithkline
	Sepsis			
	Thrombocytopenia			
	Weight Decreased			
	White Blood Cell Count Decreased			

Date:07/25/03ISR Number: 4156849-6Report Type:Expedited (15-DaCompany Report #2003030526
Age: Gender:Female I/FU:F

Outcome
Hospitalization -
Initial or Prolonged

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Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4800 MG		Aggression Bipolar Disorder	Consumer Health	Neurontin (Gabapentin)	PS		
(DAILY),		Psychotic Disorder	Professional				
		Weight Decreased					

Date:07/25/03ISR Number: 4156852-6Report Type:Expedited (15-DaCompany Report #2003030108
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Pressure Decreased Dizziness	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Prostate Cancer		Propranolol Hydrochloride (Propranolol Hydrochloride)	SS		ORAL
120 MG, ORAL				Valproate Semisodium	C		

Date:07/25/03ISR Number: 4157180-5Report Type:Expedited (15-DaCompany Report #2003017894
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 10 MG DAILY, Initial or Prolonged ORAL		Aphonia	Consumer	Norvasc (Amlodipine)	PS		ORAL
Other		Blood Urine	Health				
ORAL		Chromaturia Memory Impairment	Professional	Neurontin (Gabapentin)	SS		ORAL
		Myocardial Infarction		Acetylsalicylic Acid (Acetylsalicylic Acid) Multivitamins And Iron	C C		

Losartan Potassium	
(Losartan Potassium)	C
Glipizide	
(Glipizide)	C
Bupropion	
Hydrochloride	
(Bupropion	
Hydrochloride)	C
Metoprolol Tartrate	
(Metoprolol	
Tartrate)	C
Olanzapine	
(Olanzapine)	C
Iron (Iron)	C
Isosorbide	
Mononitrate	
(Isosorbide	
Mononitrate)	C
Mirtazapine	
(Mirtazapine)	C
Glyceryl Trinitrate	
(Glyceryl	
Trinitrate)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/25/03ISR Number: 4157602-XReport Type:Expedited (15-DaCompany Report #2003022646

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Abdominal Pain Lower	Consumer	Zoloft (Sertraline)	PS		ORAL
ORAL						
Initial or Prolonged	Abortion Spontaneous		Neurontin			
Other	Agoraphobia		(Gabapentin)	SS		
	Alopecia		Zyrtec-D12 Hour			
	Anaesthetic Complication		(Cetirizine			
ORAL	Apallic Syndrome		/Pseudoephedrine)	SS		ORAL
	Chest Pain		Zyrtec			
	Constipation		(Cetirizine			
ORAL	Death Of Sibling		Hydrochloride)	SS		ORAL
	Dyspnoea		All Other			
	Fungal Infection		Therapeutic Product	SS		
	Heart Rate Abnormal		Ketamine			
	Hypotonia		Hydrochloride			
	Ileus		(Ketamine)	SS		
	Incision Site		Hydroxyzine Embonate	C		
	Complication		Alprazolam	C		
	Nasal Congestion					
	Neoplasm					
	Ovarian Cyst					
	Panic Attack					
	Respiratory Disorder					
	Seasonal Allergy					
	Urinary Retention					

Date:07/25/03ISR Number: 4157610-9Report Type:Expedited (15-DaCompany Report #2003030466

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Heat Stroke	Health Professional	Neurontin			
			(Gabapentin)	PS		

Date:07/28/03ISR Number: 4157764-4Report Type:Expedited (15-DaCompany Report #2003030361

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis	Consumer	Neurontin			
		Bipolar Disorder		(Gabapentin)	PS		ORAL
ORAL		Drug Hypersensitivity		Venlafaxine			
		Nerve Injury		Hydrochloride			
		Overweight		(Venlafaxine			
				Hydrochloride)	SS		
				Anti-Diabetics	C		
				Estrogens	C		
				Thyroid (Thyroid)	C		

Date:07/28/03ISR Number: 4157781-4Report Type:Expedited (15-DaCompany Report #2003030520
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Testosterone	Health	Neurontin			
		Decreased	Professional	(Gabapentin)	PS		
		Erythema		Testosterone			
		Hypothalamo-Pituitary		(Testosterone)	SS		
		Disorders					
		Oedema Peripheral					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/28/03ISR Number: 4157992-8Report Type:Expedited (15-DaCompany Report #2003030604
 Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Culture Positive	Consumer	Neurontin			
Hospitalization -		Coma		(Gabapentin)	PS		ORAL
ORAL							
Initial or Prolonged		Convulsion		All Other			
Other		Dysphemia		Therapeutic Products	C		
		Pruritus					
		Somnolence					
		Tremor					

Date:07/28/03ISR Number: 4158202-8Report Type:Expedited (15-DaCompany Report #2003030174
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blindness	Consumer	Neurontin			
				(Gabapentin)	PS		

Date:07/28/03ISR Number: 4158211-9Report Type:Expedited (15-DaCompany Report #001-0945-950345
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abasia	Consumer	Neurontin			
Initial or Prolonged		Abdominal Discomfort	Health	(Gabapentin)	PS		ORAL
ORAL							
Other		Asthenia	Professional	Dilantin Kapseals			
		Chest Discomfort		(Phenytoin Sodium)	SS		ORAL
ORAL							
		Convulsion		Zoloft (Sertraline)	SS		ORAL
ORAL							
		Deafness		Lamotrigine			
		Dental Plaque		(Lamotrigine)	SS		
		Difficulty In Walking		Topiramate			
		Dizziness		(Topiramate)	SS		
		Hypoaesthesia		Tizanidine			
		Muscle Spasms		Hydrochloride			
		Paralysis		(Tizanidine)			

ORAL	Road Traffic Accident	Hydrochloride)	SS	ORAL
	Somnolence	Cefuroxime Axetil		
	Spinal Disorder	(Cefuroxime Axetil)	C	
	Tinnitus	Fluoxetine		
	Tooth Discolouration	Hydrochloride		
	Victim Of Abuse	(Fluoxetine		
	Visual Acuity Reduced	Hydrochloride)	C	
		Citalopram		
		Hydrobromide		
		(Citalopram		
		Hydrobromide)	C	
		Levetiracetam		
		(Levetiracetam)	C	
		Lactulose		
		(Lactulose)	C	
		Lorazepam		
		(Lorazepam)	C	

Date:07/29/03ISR Number: 4157541-4Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 198885

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Gabapentin 100mg Cap	PS		
2 QHS							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/03ISR Number: 4159634-4Report Type:Expedited (15-DaCompany Report #2003030653

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence	Health	Neurontin			
ORAL		Epilepsy	Professional	(Gabapentin)	PS		ORAL
		Medication Error		Fentanyl (Fentanyl)	C		

Date:07/29/03ISR Number: 4159635-6Report Type:Expedited (15-DaCompany Report #2003031058

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Burning Sensation	Consumer	Neurontin			
Initial or Prolonged		Cystitis		(Gabapentin)	PS		ORAL
300 MG							
Other		Dysgeusia					
(DAILY), ORAL		Hepatic Cyst		Levetiracetam			
		Hypoaesthesia		(Levetiracetam)	SS		
		Multiple Allergies		Sertraline			
		Muscle Spasms		Hydrochloride			
		Pain		(Sertraline			
		Swelling		Hydrochloride)	C		
				Entex (Phenylephrine			
				Hydrochloride,			
				Guaifenesin,			
				Phenylpropanolamine			
				Hydrochloride)	C		
				Cyclobenzaprine			
				(Cyclobenzaprine)	C		
				Meloxicam			
				(Meloxicam)	C		
				Lansoprazole			
				(Lansoprazole)	C		
				Sucralfate			
				(Sucralfate)	C		
				Salbutamol			
				(Salbutamol)	C		
				Montelukast Sodium			
				(Montelukast Sodium)	C		
				Anovlar			
				(Norethisterone			

Acetate,
 Ethinylestradiol) C
 Alendronate Sodium C
 (Alendronate Sodium) C
 Budesonide
 (Budesonide) C

Date:07/29/03ISR Number: 4159636-8Report Type:Expedited (15-DaCompany Report #2003030655

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other 3000 MG (DAILY), ORAL		Drug Ineffective Emotional Disorder Muscle Spasms Pain Paralysis	Consumer	Neurontin (Gabapentin) Vicodin (Paracetamol, Hydrocodone Bitartrate) Metformin Hydrochloride	PS C		ORAL

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Freedom Of Information (FOI) Report

(Metformin
Hydrochloride) C
Pioglitazone
(Pioglitazone) C
Insulin (Insulin) C
Temazepam (Temazepam) C
Fluoxetine
Hydrochloride
(Fluoxetine
Hydrochloride) C

Date:07/30/03ISR Number: 4160367-9Report Type:Expedited (15-DaCompany Report #S03-FRA-03138-01
Age:55 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 20 MG QD PO	Cyanosis Dermatitis Bullous	Foreign Health	Seropram (Citalopram)	PS		ORAL
80 MG QD PO	Lymphoedema Pruritus	Professional Other	Skenen (Morphine Sulfate)	SS		ORAL
60 MG QD PO	Rash Papular		Skenan (Morphine Sulfate)	SS		ORAL
300 MG QD PO			Neurontin (Gabapentin)	SS		ORAL
			Lactulose	SS		

Date:07/30/03ISR Number: 4160787-2Report Type:Expedited (15-DaCompany Report #GBR-2003-0000658
Age:68 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2 UNIT, QID	Confusional State Constipation Hallucination Mobility Decreased	Foreign Health Professional Other	Oxycodone Hcl Cr Tablets(Oxycodone Hydrochloride) Cr Tablet	PS		
300 MG, TID,	Sedation Speech Disorder		Gabapentin(Gabapenti n)	SS		ORAL

ORAL

Celecoxib(Celecoxib) SS

ORAL

100 MG, BID,

ORAL

Acetylsalicylic Acid C
 Hydroxocobalamin C
 Diclofenac C
 Co-Danthrusate C
 Lisinopril C
 Quinine Sulphate (Quinine Sulfate) C
 Iron Sulfate (Ferrous Sulfate) C

Date:07/30/03ISR Number: 4160885-3Report Type:Expedited (15-DaCompany Report #2003169331GB

Age:68 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 100 MG, BID,	Confusional State Constipation	Foreign Health	Celebrex(Celecoxib) apsule	C PS		ORAL
ORAL 2 DF, QID	Hallucination Mobility Decreased	Professional	Oxycodone(Oxycodone)	SS		
300 MG, TID,	Sedation Speech Disorder	Other	Gabapentin(Gabapenti n)	SS		ORAL

ORAL

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Aspirine	C
Hydroxocobalamin	C
Diclofenac	C
Co-Danthrusate (Docusate Sodium, Dantron)	C
Lisinopril	C
Quinine Sulphate	C
Iron Sulfate	C

Date:07/30/03ISR Number: 4160973-1Report Type:Expedited (15-DaCompany Report #2003028207

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Decreased	Health	Geodon (Ziprasidone)	PS		
80 MG (BID)		Dizziness	Professional	Neurontin (Tablets)			
		Dry Mouth	Company	(Gabapentin)	SS		
2400 MG (TID)		Electrocardiogram Qt	Representative	Sertraline			
		Corrected Interval		Hydrochloride			
		Prolonged		(Sertraline			
		Heart Rate Decreased		Hydrochloride)	C		
		Heart Rate Irregular					
		Pruritus					
		Somnolence					

Date:07/30/03ISR Number: 4161253-0Report Type:Expedited (15-DaCompany Report #2003030759

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaesthetic Complication	Consumer	Neurontin			
		Anxiety		(Gabapentin)	PS		
		Bronchial Infection		All Other			
		Extrasystoles		Therapeutic Products	SS		
		Injury		Fluoxetine			
		Micturition Disorder		Hydrochloride			
		Panic Disorder		(Fluoxetine			
		Post-Traumatic Stress		Hydrochloride)	C		
		Disorder		Alprazolam			
		Suicide Attempt		(Alprazolam)	C		

Urethral Disorder

Risperidone
(Risperidone)

C

Date:07/30/03ISR Number: 4161256-6Report Type:Expedited (15-DaCompany Report #2003027160
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchopneumonia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other	2100 MG DAILY		Professional				ORAL
				Ketogen (Ketobemidone Hydrochloride, Dimethyl-3,3-Diphenyl-1-1-Methylallylamine	SS		ORAL
	120 MG DAILY						ORAL
				Ketobemidone (Ketobemidone)	SS		ORAL
	225 MG DAILY						ORAL

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Freedom Of Information (FOI) Report

Glyceryl Trinitrate C
 Levothyroxine Sodium C
 Propranolol
 Hydrochloride C
 Estrogens Conjugated C
 Flunitrazepam C
 Omeprazole Magnesium C

Date:07/30/03ISR Number: 4161378-XReport Type:Expedited (15-DaCompany Report #2003030994
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dizziness	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:07/30/03ISR Number: 4161415-2Report Type:Expedited (15-DaCompany Report #2003030659
 Age:88 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (300, QID), ORAL		Catatonia Muscle Rigidity Mydriasis Tremor	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Metoprolol (Metoprolol)	C		
				Human Mixtard (Insulin Human, Insulin Human Injection, Isophane)	C		
				Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
				Pantoprazole (Pantoprazole)	C		
				Citalopram Hydrobromide (Citalopram Hydrobromide)	C		
				Prochlorperazine			

Edisylate
 (Prochlorperazine
 Edisylate) C
 All Other
 Therapeutic Products C

Date:07/31/03ISR Number: 4161959-3Report Type:Expedited (15-DaCompany Report #2003031277
 Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300, TID), ORAL		Confusional State Constipation Hallucination	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
200 MG (100, BID), ORAL		Mobility Decreased Sedation Speech Disorder		Celecoxib (Celecoxib)	SS		ORAL
(FOUR TIMES)				Oxycodone (Oxycodone)	SS		
				Acetylsalicyclic Acid			

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(Acetylsalicylic Acid)	C
Hydroxocobalamin (Hydroxocobalamin)	C
Diclofenac (Diclofenac)	C
Coloxyl With Danthron (Dantron, Docusate Sodium)	C
Lisinopril (Lisinopril)	C
Quinine Sulfate (Quinine Sulfate)	C

Date:07/31/03ISR Number: 4161985-4Report Type:Expedited (15-DaCompany Report #2003030996
 Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dermatitis Allergic	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
2400 MG, ORAL			Professional	Esomeprazole (Esomeprazole)	C		
				Atorvastatin Calcium (Atorvastatin Calcium)	C		
				Clopidogrel Sulfate (Clopidogrel Sulfate)	C		

Date:07/31/03ISR Number: 4162059-9Report Type:Expedited (15-DaCompany Report #2003031259
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaemia Lymphoedema	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Overdose		All Other Therapeutic Products	SS		
		Petit Mal Epilepsy		Fentanyl (Fentanyl)	C		
		Weight Decreased		Carisoprodol (Carisoprodol)	C		

Diazepam (Diazepam)	C
Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C
All Other Therapeutic Products	C
Trazodone (Trazodone)	C

Date:07/31/03ISR Number: 4162073-3Report Type:Expedited (15-DaCompany Report #2003031179

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Parkinson'S Disease	Consumer	Neurontin (Gabapentin) Quinapril Hydrochloride (Quinapril	PS		

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Hydrochloride) C
 Alendronate Sodium
 (Alendronate Sodium) C

Date:08/01/03ISR Number: 4159367-4Report Type:Expedited (15-DaCompany Report #WAES 0209USA02503
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Enzymes Increased		Vioxx	PS	Merck & Co., Inc	ORAL
Hospitalization -		Completed Suicide		Gabapentin	SS		
Initial or Prolonged		Coordination Abnormal		Ultram	SS		ORAL
Other		Cyanosis					
		Dehydration					
		Electromechanical					
		Dissociation					
		Heart Rate Decreased					
		Hypotension					
		Mental Status Changes					
		Myoglobin Urine Present					
		Renal Failure					
		Somnolence					
		White Blood Cell Count					
		Increased					

Date:08/01/03ISR Number: 4162132-5Report Type:Expedited (15-DaCompany Report #200311636DE
 Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Aspartate	Foreign	Leflunomide (Arava)	PS		ORAL
5 TO 10 MG PO 3 YR		Aminotransferase	Health	Entacapone (Comtan)	SS		ORAL
Initial or Prolonged		Increased	Professional	Gabapentin			
200 MG/DAY PO		Asthenia	Other	(Neurontin)	SS		ORAL
300 MG/DAY PO		Diarrhoea		Selegiline			
		Drug Interaction		Hydrochloride			
		Gamma-Glutamyltransferase		(Selegilin "Teva")	SS		
		Increased		Levodopa	C		
		Pancreatic Insufficiency		Cabergoline			
		Weight Decreased		(Cabaseril)	C		

Date:08/01/03ISR Number: 4162874-1Report Type:Expedited (15-DaCompany Report #2003030919
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged Other	Unevaluable Event	Health Professional	Neurontin (Gabapentin)	PS		

Date:08/01/03ISR Number: 4162919-9Report Type:Expedited (15-DaCompany Report #2003030940
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Dysphagia Gastric Bypass Pharmaceutical Product Complaint	Health Professional	Neurontin (Gabapentin)	PS		

Tongue Disorder
Toothache
Tremor
Urinary Incontinence
Urinary Tract Infection
Visual Acuity Reduced
Vomiting

Date:08/04/03ISR Number: 4163729-9Report Type:Expedited (15-DaCompany Report #2003031371
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 500 MG (TID), ORAL		Hypoglycaemia Syncope Vision Blurred Visual Disturbance	Consumer	Neurontin (Gabapentin) Morphine Sulfate (Morphine Sulfate) Clonazepam	PS C		ORAL

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(Clonazepam) C

Date:08/04/03ISR Number: 4163730-5Report Type:Expedited (15-DaCompany Report #2003024808

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Body Temperature Decreased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL	Catatonia Coma	Company Representative	Alprazolam (Alprazolam)	SS		ORAL
	Intentional Misuse Pupil Fixed Pyrexia		Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride)	SS		

Date:08/04/03ISR Number: 4163793-7Report Type:Expedited (15-DaCompany Report #2003029575

Age:57 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death Hospitalization - Initial or Prolonged Other	Asthenia Blood Glucose Decreased Coma Confusional State Dysphagia Gastrointestinal Motility Disorder Haematocrit Decreased Myocardial Infarction	Consumer	Neurontin (Gabapentin)	PS		
(BID), ORAL			Diltiazem Hydrochloride (Diltiazem Hydrochloride)	SS		
			Rofecoxib (Rofecoxib)	SS		
			Bosentan (Bosentan)	SS		ORAL
INTRAVENOUS	Pneumonia Aspiration Pulmonary Hypertension Pyrexia		Epoprostenol Sodium (Epoprostenol Sodium)	SS		
	Rash Erythematous Respiratory Arrest Sepsis Thrombocytopenia Weight Decreased White Blood Cell Count Decreased		Lansoprazole (Lansoprazole) Spironolactone (Spironolactone) Amitriptyline Hydrochloride (Amitriptyline	C C		

Hydrochloride)	C
Citalopram	
Hydrobromide	
(Citalopram	
Hydrobromide)	C
Oxycodone	
Hydrochloride	
(Oxycodone	
Hydrochloride)	C
Fexofenadine	
Hydrochloride	
(Fexofenadine	
Hydrochloride)	C
Furosemide	
(Furosemide)	C
Warfarin (Warfarin)	C

FDA - Adverse Event Reporting System (AERS)

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Date:08/04/03ISR Number: 4163941-9Report Type:Expedited (15-DaCompany Report #2003028326

Age:91 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 600 MG (TID), Initial or Prolonged ORAL Other	Coma	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Capozide (Hydrochlorothiazide , Captopril) Di-Gesic (Paracetamol, Dextropropoxyphene) Caffeine (Caffeine) Vitamin B1 And B6 (Thiamine, Pyridoxine)	PS C C C C		ORAL

Date:08/04/03ISR Number: 4164291-7Report Type:Expedited (15-DaCompany Report #2003031281

Age:55 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG Other (DAILY), ORAL 80 MG (DAILY), ORAL (BID), ORAL 20 MG	Cyanosis Dermatitis Bullous Lymphoedema Pruritus Rash Erythematous Rash Papular	Foreign Health Professional	Neurontin (Gabapentin) Morphine Sulfate (Morphine Sulfate) Lactulose (Lactulose) Citalopram Hydrobromide (Citalopram Hydrobromide)	PS SS SS SS		ORAL ORAL ORAL ORAL

(DAILY), ORAL

Date:08/04/03ISR Number: 4164388-1Report Type:Expedited (15-DaCompany Report #2003163834GB

Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous Complications Of Maternal	Foreign Health	Detrusitol(Tolterodi ne)	PS		ORAL
2MG, QD, ORAL		Exposure To Therapeutic Drugs	Professional Other	Zopiclone (Zopiclone)	SS		
BID		Maternal Drugs Affecting Foetus		Gabapentin (Gabapentin) Propranolol (Propranolol) Clonazepam(Clonazepa m) Sertraline (Sertraline)	SS SS SS SS		

QD

Date:08/05/03ISR Number: 4165508-5Report Type:Expedited (15-DaCompany Report #2003031544

Age:65 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Balance Disorder Blood Glucose Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Coma Diabetes Mellitus Inadequate Control	Report Source	Product	Role	Manufacturer	Route
(TID), ORAL		Difficulty In Walking Dyspnoea	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Speech Disorder	Professional Company Representative	Fenofibrate (Fenofibrate)	C		
				Indapamide (Indapamide)	C		
				Olanzapine (Olanzapine)	C		
				Thioridazine Hydrochloride (Thioridazine Hydrochloride)	C		
				Ranitidine Hydrochloride (Ranitidine Hydrochloride)	C		
				Gliquidone (Gliquidone)	C		

Date:08/05/03ISR Number: 4165535-8Report Type:Expedited (15-DaCompany Report #2003031747
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
300 MG (TID), ORAL			Professional				
900 MG (TID), ORAL				Oxcarbazepine (Oxcarbazepine)	SS		ORAL

Date:08/05/03ISR Number: 4165537-1Report Type:Expedited (15-DaCompany Report #2003028852
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - ORAL	Initial or Prolonged	Stevens-Johnson Syndrome Toxic Epidermal Necrolysis	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Clonazepam (Clonazepam) Antineoplastic Agents	C		

Date:08/05/03ISR Number: 4165687-XReport Type:Expedited (15-DaCompany Report #2003031753
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1800 MG (600, Other TID),	Cardiac Failure Scrotal Oedema	Health Professional	Neurontin (Gabapentin)	PS		
				Celecoxib (Celecoxib)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/05/03ISR Number: 4165688-1Report Type:Expedited (15-DaCompany Report #2003031542

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Ankle Fracture	Consumer	Neurontin			
Initial or Prolonged	Compression Fracture		(Gabapentin)	PS		ORAL
(DAILY), ORAL						
Other	Fall		Vaseretic			
	Sensory Loss		(Hydrochlorothiazide			
	Spinal Compression		, Enalapril Maleate)	C		
	Fracture		Oxycocet			
			(Paracetamol,			
			Oxycodone			
			Hydrochloride)	C		

Date:08/06/03ISR Number: 4162961-8Report Type:Expedited (15-DaCompany Report #WAES 0307USA01625

Age:57 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Asthenia		Vioxx	PS	Merck & Co., Inc	ORAL
Hospitalization -	Coma		Tracleer	SS		ORAL
53 DAY						
Initial or Prolonged	Confusional State		Tracleer	SS		ORAL
Other	Dyskinesia Oesophageal		Flolan	SS		
INTRAVENOUS						
	Dysphagia		Cardizem	SS		
	Haematocrit Decreased		Neurontin	SS		
	Hyperhidrosis		Prevacid	C		
	Hypoglycaemia		Aldactone	C		
	Hyponatraemia		Elavil	C		ORAL
	Pneumonia Aspiration		Celexa	C		
	Pyrexia		Oxycontin	C		
	Rash Erythematous		Allegra	C		
	Respiratory Arrest		Lasix (Furosemide)	C		
	Thrombocytopenia		Coumadin	C		
	Weight Decreased					
	White Blood Cell Count					
	Decreased					

Date:08/06/03ISR Number: 4165216-0Report Type:Expedited (15-DaCompany Report #2003CG01097

Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Pulmonary Oedema Respiratory Disorder	Foreign Health Professional Other	Mopral Euphylline Neurontin Deroxat Mono-Tildiem	PS SS SS SS SS		ORAL

300 MG DAILY

PO

20 MG DAILY

PO

Date:08/06/03ISR Number: 4166093-4Report Type:Expedited (15-DaCompany Report #2003030521
Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blindness Transient Visual Acuity Reduced	Foreign Health Professional Company Representative	Neurontin (Gabapentin) All Other Therapeutic Products	PS C		ORAL

600 MG (BID),
ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/06/03ISR Number: 4166129-0Report Type:Expedited (15-DaCompany Report #2003030526

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 4800 MG Other (DAILY)		Affective Disorder Aggression Bipolar Disorder Mood Swings Psychotic Disorder Suicide Attempt Weight Decreased Weight Increased	Consumer Health Professional	Neurontin (Gabapentin)	PS		

Date:08/06/03ISR Number: 4166527-5Report Type:Expedited (15-DaCompany Report #2003031828

Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG Other (DAILY)		Anxiety Cognitive Disorder Depression Dizziness Memory Impairment Nausea Somnolence	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:08/06/03ISR Number: 4166531-7Report Type:Expedited (15-DaCompany Report #031-0945-M02000011

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Epilepsy	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG Other (DAILY), ORAL	Angiopathy Confusional State Hallucination, Visual Personality Change	Foreign Health Professional	Neurontin (Gabapentin) Digoxin (Digoxin) Warfarin Sodium (Warfarin Sodium) Cyanocobalamin (Cyanocobalamin) Furosemide (Furosemide) Potassium Chloride (Potassium Chloride) Lansoprazole (Lanzoprazole)	PS C C C C C C C		ORAL

Outcome	PT	Report Source
Other	Drug Interaction Hepatitis	Foreign Health

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Professional Company Representative	Product	Role	Manufacturer	Route
1200 MG			Neurontin (Gabapentin)	PS		ORAL
(TID), ORAL			Carbamazepine (Carbamazepine)	SS		
			Panadeine Co (Codeine Phosphate Paracetamol)	C		
			Quinapril (Quinapril)	C		
			Serenoa Repens (Serenoa Repens)	C		
			Sotalol (Sotalol)	C		

Date:08/07/03ISR Number: 4167103-0Report Type:Expedited (15-DaCompany Report #2003032572
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG, ORAL		Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
			Professional Company Representative	Carbamazepine (Carbamazepine)	C		
				Primidone (Primidone)	C		

Date:08/07/03ISR Number: 4167160-1Report Type:Expedited (15-DaCompany Report #2003032597
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aneurysm	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Abnormal Behaviour Confusional State Disorientation Sedation	Consumer Health Professional	Neurontin (Gabapentin) Oxycocet (Paracetamol, Oxycodone Hydrochloride) Methadone (Methadone) Bupropion Hydrochloride (Bupropion Hydrochloride) Mirtazapine (Mirtazapine)	PS C C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/07/03ISR Number: 4167164-9Report Type:Expedited (15-DaCompany Report #2003024623

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Hypersensitivity Drug Ineffective	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
200 MG (BID), Other ORAL		Dyspnoea	Professional				
		Fibrocystic Breast Disease		Benadryl Injection (Diphenhydramine)	SS		
INTRAVENOUS	INTRAVENOUS	Hysterectomy Infection Menorrhagia Ovarian Cyst Swollen Tongue Throat Lesion Tongue Disorder		Ciprofloxacin (Ciprofloxacin) Paracetamol (Paracetamol)	SS C		

Date:08/07/03ISR Number: 4167165-0Report Type:Expedited (15-DaCompany Report #2003021783

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Toxicity Medication Error	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 MG (ONCE), ORAL		Overdose					
				Lamotrigine (Lamotrigine)	SS		ORAL
200 MG (ONCE), ORAL							
				Carbamazepine (Carbamazepine)	SS		ORAL
400 MG (ONCE), ORAL							
				Human Mixtard (Insulin Human, Insulin Human Injection, Isophane)	C		

Potassium	
(Potassium)	C
Insulin (Insulin)	C
Oxybutynin	
(Oxybutynin)	C
Sertraline	
Hydrochloride	
(Sertraline	
Hydrochloride)	C
Leuprorelin Acetate	
(Leuprorelin	
Acetate)	C
Risperidone	
(Risperidone)	C

Date:08/07/03ISR Number: 4167174-1Report Type:Expedited (15-DaCompany Report #2003031756
Age:49 YR Gender:Female I/FU:I

Outcome	PT
Other	Convulsion
	Eye Swelling
	Medication Error
	Paraesthesia
	Photopsia
	Psychomotor Hyperactivity
	Tendon Disorder

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Freedom Of Information (FOI) Report

Treatment Noncompliance

Dose	Duration		Report Source	Product	Role	Manufacturer	Route
ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Lorazepam (Lorazepam)	C		
				Hydrocodone (Hydrochodone)	C		

Date:08/07/03ISR Number: 4167328-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (BID) Disability , ORAL Other		Application Site Pain Brain Neoplasm	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Cardiac Arrest	Professional				
		Chest Pain Chondromalacia Electrocardiogram P Wave Abnormal Fall Limb Injury Meniscus Lesion Pain In Extremity Scar	Company Representative	Valproate Sodium (Valproate Sodium) Omeprazole (Omeprazole) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Capsaicin (Capsaicin)	SS C C C		

Date:08/07/03ISR Number: 4167330-2Report Type:Expedited (15-DaCompany Report #2003031837
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Hyperhidrosis	Health	Neurontin			

Initial or Prolonged 2100 MG	Muscle Spasms	Professional	(Gabapentin)	PS	
(DAILY),					
10 MG			Escitalopram (Escitalopram)	SS	ORAL
(DAILY), ORAL			Thyroid Hormones	C	

Date:08/07/03ISR Number: 4167331-4Report Type:Expedited (15-DaCompany Report #2003031849
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Medical Device Implantation Unevaluable Event	Consumer	Neurontin (Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:08/07/03ISR Number: 4167332-6Report Type:Expedited (15-DaCompany Report #2003031852

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID), Other ORAL	Blood Pressure - Immeasurable Convulsion Dehydration Influenza Tremor Weight Increased White Blood Cell Count Decreased	Consumer	Neurontin (Gabapentin) Carbamazepine (Carbamazepine) Clorazepate Dipotassium (Clorazepate Dipotassium) Valproate Semisodium (Valproate Semisodium)	PS SS C C		ORAL

Date:08/08/03ISR Number: 4166198-8Report Type:Direct

Company Report #CTU 199613

Age:52 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600MG TID ORAL	Condition Aggravated Speech Disorder		Gabapentin 300mg Parke-Davis/Pfizer Quetiapine Sertraline Cyclobenzaprine	PS C C C	Parke-Davis/Pfizer	ORAL

Date:08/09/03ISR Number: 4166260-XReport Type:Expedited (15-DaCompany Report #2003031855

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - ORAL	Anxiety Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL

Initial or Prolonged	Lung Disorder	Zoloft (Sertraline)	SS	ORAL
ORAL				
Other	Multiple Allergies	All Other		
	Respiratory Disorder	Therapeutic Products	SS	
		Fexofenadine		
		Hydrochloride		
		(Fexofenadine		
		Hydrochloride)	C	
		Alprazolam		
		(Alprazolam)	C	
		Estrogens Conjugated		
		(Estrogens		
		Conjugated)	C	
		Theophylline		
		(Theophylline)	C	
		Salbutamol		
		(Salbutamol)	C	
		Ipratropium Bromide		
		(Ipratropium		
		Bromide)	C	

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Freedom Of Information (FOI) Report

Date:08/09/03ISR Number: 4166262-3Report Type:Expedited (15-DaCompany Report #2003032836

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged UNKNOWN Other (DAILY), ORAL	Depression Dyspepsia Flatulence Herpes Zoster Ophthalmic Hiatus Hernia	Consumer	Lipitor (Atorvastatin)	PS		ORAL
900 MG (THREE TIMES A DAY), ORAL	Irritability Mental Disorder		Neurontin (Gabapentin)	SS		ORAL
	Muscle Spasms Neuropathy Osteoporosis Prostate Cancer		Alendronate Sodium (Alendronate Sodium) Fentanyl (Fentanyl) Paracetamol (Paracetamol) Oxygen (Oxygen) Salbutamol (Salbutamol) Ipratropium Bromide (Ipratropium Bromide) Leuprorelin Acetate (Leuprorelin Acetate) Leuprorelin Acetate (Leuprorelin Acetate)	SS C C C C C C		

Date:08/11/03ISR Number: 4167806-8Report Type:Expedited (15-DaCompany Report #2003032175

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG (100, BID, ORAL	Breast Cancer Drug Effect Decreased Drug Interaction	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL

Gingival Bleeding
Retinal Disorder
Surgery

All Other	
Non-Therapeutic	
Products	SS
Codeine (Codeine)	C
Levothyroxine Sodium	
(Levothyroxine	
Sodium)	C
Pantoprazole	
(Pantoprazole)	C
Cyanocobalamin	
(Cyanocobalamin)	C
All Other	
Therapeutic Products	C
Fenofibrate	
(Fenofibrate)	C
Trazodone	
(Trazodone)	C
Omeprazole	
(Omeprazole)	C
Calcium/Minerals Nos	
(Calcium, Minerals	
Nos)	C

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Freedom Of Information (FOI) Report

Date:08/11/03ISR Number: 4167814-7Report Type:Expedited (15-DaCompany Report #2003032129
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG (BID), ORAL		Blindness Condition Aggravated Vitamin A Deficiency	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Retinol (Retinol)	C		
				Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		
				Bupropion Hydrochloride (Bupropion Hydrochloride)	C		
				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride,	C		

Date:08/11/03ISR Number: 4167855-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200657
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 1800 MG (DAILY), ORAL		Brain Damage Depression Drug Hypersensitivity	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Dysarthria		Lamotrigine (Lamotrigine)	C		
		Dyskinesia		Lithium Carbonate (Lithium Carbonate)	C		
		Eye Movement Disorder		Clonazepam (Clonazepam)	C		
		Gingival Pain		Methylpheniate Hydrochloride(Methyl pheniate			
		Glossodynia		Hydrochloride)	C		
		Malnutrition					
		Medication Error					
		Nerve Injury					
		Nervous System Disorder					
		Prescribed Overdose					

Sensory Disturbance
Speech Disorder
Swollen Tongue
Tooth Disorder

Levothyroxine
Sodium(Levothyroxine
Sodium) C
Liothyronine
Sodium(Liothyronine
Sodium) C
Sertraline
Hydrochloride(Sertra
line Hydrochloride) C
Pilocarpine
Hydrochloride
(Pilocarpine
Hydrochloride) C
Metoprolol Succinate
(Metoprolol
Succinate) C
Lansoprazole
(Lansoprazole) C
Hyoscyamine Sulfate
(Hyoscyamine
Sulfate) C
Diltiazem

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Hydrochloride(Diltiazem Hydrochloride) C
Lamotrigine (Lamotrigine) C

Date:08/12/03ISR Number: 4166438-5Report Type:Direct
Age:43 YR Gender:Female I/FU:I

Company Report #CTU 199775

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Suicide Attempt		Neurontin Pill			
Life-Threatening Hospitalization - Initial or Prolonged Disability				(Warner Lambert/Pfizer 2700 Mgs As Of May 2003	PS	Warner Lambert/Pfizer	
SEE INCLOSED Required ZEROXES Intervention to Prevent Permanent Impairment/Damage							

Date:08/12/03ISR Number: 4168975-6Report Type:Expedited (15-DaCompany Report #2003031759
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abnormal Behaviour Alopecia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG DAILY		Colitis Ulcerative					
ORAL		Diarrhoea Disturbance In Attention		Ibuprofen (Ibuprofen)	SS		ORAL
2400MG		Dizziness					
(DAILY) ORAL		Drug Ineffective Drug Interaction Gastrointestinal Haemorrhage Impaired Driving Ability Intervertebral Disc Protrusion		Valproate Semisodium (Valproate Semisodium)	SS		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	SS		
				Glycopyrronium(Glyco			

Migraine
Pain In Extremity
Road Traffic Accident

pyrronium Bromide) C
Propranolol
Hydrochloride
(Propranolol
Hydrochloride) C
Amitriptyline
Hydrochloride
(Amitriptyline
Hydrochloride) C

Date:08/14/03ISR Number: 4169083-0Report Type:Expedited (15-DaCompany Report #2003025794
Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3200 MG (800, Other QID), ORAL		Delirium	Health Professional Company Representative	Neurontin (Gabapentin) Mirtazapine (Mirtazapine)	PS SS		ORAL ORAL
(DAILY), ORAL				Nortriptyline Hydrochloride (Nortriptyline			

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600 MG (BID),	Feeling Abnormal	(Gabapentin)	SS	ORAL
ORAL	Feeling Hot			
	Flushing	Enalapril		
	Neuropathy	(Enalapril)	C	
	Pancreatitis	All Other		
		Therapeutic Products	C	

Date:08/15/03ISR Number: 4169933-8Report Type:Direct Company Report #CTU 200111
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorgasmia		Gabapentin 300 Mg			
		Discomfort		Cap (Parke Davis)	PS	(Parke Davis)	ORAL
900 MG TID PO		Erythema		Sildenafil	C		
		Hyperaesthesia		Alprazolam	C		
		Penis Disorder					

Date:08/15/03ISR Number: 4170326-8Report Type:Expedited (15-DaCompany Report #2003026291
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bacterial Infection	Health	Neurontin			
		Cardiac Flutter	Professional	(Gabapentin)	PS		ORAL
900 MG (300							

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TID), ORAL

Sumatriptan
 Succinate C
 Amitriptyline C
 Nitrofurantoin C
 Diazepam C
 Oxaprozin C
 Biselect
 (Hydrochlorothiazide
 , Bisoprolol
 Fumarate) C

Date:08/15/03ISR Number: 4170327-XReport Type:Expedited (15-DaCompany Report #2003026124
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Depressed Level Of	Consumer	Neurontin (Tablets)			
Other		Consciousness		(Gabapentin)	PS		

Date:08/15/03ISR Number: 4170332-3Report Type:Expedited (15-DaCompany Report #2003022132
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG DAILY		Somnolence	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other ORAL			Professional				

Furosemide C
 Potassium Chloride C
 Losartan Potassium C
 Pravastatin Sodium C
 Warfarin Sodium C
 Digoxin C
 Seretide Mite
 (Fluticasone
 Propionate,
 Salmeterol
 Xinafoate) C
 Glipizide C
 Entex (Phenylephrine

Hydrochloride,
 Guaifenesin,
 Phenylpropanolamine
 Hydrochloride) C
 Metoprolol Succinate C
 Pantoprazole C
 Combivent
 (Ipratropium
 Bromide, Salbutamol
 Sulfate) C
 All Other
 Therapeutic Products C

Date:08/15/03ISR Number: 4170885-5Report Type:Direct
 Age:15 YR Gender:Male I/FU:I

Company Report #USP 50113

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Neurontin	PS	Parke-Davis	
				Zarontin	SS	Parke-Davis	

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FDA - Adverse Event Reporting System (AERS)

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Date:08/15/03ISR Number: 4172233-3Report Type:Expedited (15-DaCompany Report #2003012425
 Age:76 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2400 MG Other (TID), ORAL	Adrenal Insufficiency	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Clomipramine Hydrochloride (Clomipramine Hydrochloride)	C		

Date:08/18/03ISR Number: 4170647-9Report Type:Direct Company Report #CTU 200147
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Required 1200MG 3 Intervention to TIMES ORAL Prevent Permanent Impairment/Damage	Drug Dependence Irritable Bowel Syndrome Pruritus Stevens-Johnson Syndrome		Neurontin 400mg Parke-Davis	PS	Parke-Davis	ORAL

Date:08/18/03ISR Number: 4172074-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
 Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability Other	Actinic Keratosis Back Pain Blood Urea Nitrogen/Creatinine Ratio Increased Bone Disorder Brain Neoplasm Burning Sensation Cardio-Respiratory Arrest Chest Pain Cognitive Disorder

Conversion Disorder
Convulsion
Cyst
Disease Recurrence
Dizziness
Dysarthria
Dysphemia
Dyspnoea
Ependymoma
Erythema
Fatigue
Feeling Abnormal
Fibromyalgia
Hypoglycaemia
Memory Impairment
Meniscus Lesion
Muscle Rigidity
Nervous System Disorder
Osteopenia
Patellofemoral Pain
Syndrome
Pigmented Naevus
Respiratory Arrest

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (BID),		Somnolence Speech Disorder Tardive Dyskinesia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Tinnitus Tremor Vision Blurred Vomiting	Professional Company Representative	Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin)	SS C C C C		

Date:08/18/03ISR Number: 4172376-4Report Type:Expedited (15-DaCompany Report #2003029798

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2000 MG (BID), ORAL		Eye Haemorrhage Optic Nerve Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Trazadone (Trazadone) ... Methylphenidate (Methylphenidate) Topiramate (Topiramate) Clonazepam (Clonazepam)	C C C C C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Breast Mass	Consumer	Neurontin			
300 MG		Memory Impairment		(Gabapentin)	PS		ORAL
(DAILY), ORAL		Muscle Disorder					
				Depo-Estradiol			
				(Chlorobutanol,			
				Estradiol Cipionate,			
				Cottonseed Oil)	C		
				Zolpidem Tartrate	C		

Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased	Consumer	Neurontin			
2400 MG ,		Pharmaceutical Product	Health	(Gabapentin)	PS		ORAL
ORAL		Complaint	Professional				

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Oxycodone	
Hydrochloride	
(Oxycodone	
Hydrochloride)	C
Valsartan	
(Valsartan)	C
Loratadine	
(Loratadine)	C
Levothyroxine Sodium	
(Levothyroxine	
Sodium)	C
Calcium (Calcium)	C
Glucosamine	
(Glucosamine)	C
Vitamins	C
Acetylsalicylic	
Acid(Acetylsalicylic	
Acid)	C

Date:08/18/03ISR Number: 4173277-8Report Type:Expedited (15-DaCompany Report #2003025783
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening Hospitalization - 900 MG (TID), Initial or Prolonged ORAL	Torsade De Pointes	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other			Enalapril Maleate (Enalapril Maleate)	C		
			Atorvastatin Calcium (Atorvastatin Caldium)	C		
			Capozide (Hydrochlorothiazide , Captopril)	C		
			Hydrochlorothiazide (Hydrochlorothiazide)	C		

Date:08/18/03ISR Number: 4173443-1Report Type:Expedited (15-DaCompany Report #2003033238
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Creatine Phosphokinase Increased	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL Other		Blood Thyroid Stimulating Hormone Increased	Professional	Clomipramine (Clomipramine)	SS		
UNKNOWN		Confusional State Coronary Artery Disease		Clonazepam (Clonazepam)	SS		
		Depression Dizziness Neuralgia Pernicious Anaemia Renal Failure Urinary Tract Infection		Furosemide (Furosemide) Acetylsalicylic Acid (Acetylsalicylic Acid) Spironolactone (Spironolactone) Ramipril (Ramipril) Glyceryl Trinitrate (Glyceryl Trinitrate)	C C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/19/03ISR Number: 4172651-3Report Type:Expedited (15-DaCompany Report #2003033537

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Thought Blocking	Consumer	Neurontin			
Other		Wheelchair User		(Gabapentin)	PS		

Date:08/19/03ISR Number: 4173232-8Report Type:Expedited (15-DaCompany Report #2003017894

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 10 MG (DAILY)		Aphonia	Consumer	Norvasc (Amlodipine)	PS		ORAL
Initial or Prolonged ORAL		Blood Urine	Health				
Other ORAL		Chromaturia	Professional	Neurontin			
		Essential Tremor		(Gabapentin)	SS		ORAL
		Memory Impairment		Acetylsalicylic Acid	C		
		Myocardial Infarction		Multivitamins And Iron	C		
				Losartan Potassium (Losartan Potassium)	C		
				Glipizide (Glipizide)	C		
				Bupropion Hydrochloride (Bupropion Hydrochloride)	C		
				Metoprolol Tartrate (Metoprolol Tartrate)	C		
				Olanzapine (Olanzapine)	C		
				Iron (Iron)	C		
				Isosorbide Monoitrate (Isosorbide Mononitrate)	C		
				Mirtazapine (Mirtazapine)	C		
				Glyceryl Trinitrate (Glyceryl			

Trinitrate)

C

Date:08/19/03ISR Number: 4174517-1Report Type:Expedited (15-DaCompany Report #2003033564
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aura	Foreign	Neurontin			
1200 MG		Iron Deficiency	Health	(Gabapentin)	PS		ORAL
(TID), ORAL		Loss Of Consciousness	Professional				
			Company	Antipsychotics	C		
			Representative	Citalopram			
				Hydrobromide			
				(Citalopram			
				Hydrobrmide)	C		
				All Other			
				Therapeutic Products	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/20/03ISR Number: 4174752-2Report Type:Expedited (15-DaCompany Report #2003028998

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (BID), ORAL		Circulatory Collapse Epilepsy Insomnia Irritability	Foreign Health Professional	Neurontin (Gabapentin) Lamotrigine (Lamotrigine) All Other Therapeutic Products	PS C C		ORAL

Date:08/20/03ISR Number: 4174753-4Report Type:Expedited (15-DaCompany Report #2003033864

Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (DAILY), ORAL		Renal Tubular Necrosis	Foreign Health Professional	Gabapentin (Gabapentin) Metformin (Metformin) Isosorbide Mononitrate (Isosorbine Mononitrate) Acetylsalicylic Acid (Acetylsalicylic Acid) Atorvastatin (Atorvastatin) Prindopril (Prindopril) Metoprolol (Metoprolol) Insulin Human (Insulin Human) Insulin Human Injection, Isophane (Insulin Human	PS C C C C C C C C C C		ORAL

Injection, Isophane) C
Novolin 20/80
(Insulin Human,
Insulin Isophane,
Human Biosynthetic) C

Date:08/20/03ISR Number: 4174808-4Report Type:Expedited (15-DaCompany Report #2003CG01097
Age:76 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged	Pulmonary Oedema	Foreign Health Professional Other	Mopral Euphylline Neurontin Deroxat Mono-Tildiem	PS SS SS SS SS		ORAL
300 MG DAILY						
PO						
20 MG DAILY			Skenan	SS		ORAL
PO						

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Freedom Of Information (FOI) Report

Date:08/20/03ISR Number: 4175741-4Report Type:Expedited (15-DaCompany Report #2003034149

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Benign Breast Neoplasm	Health	Neurontin			
900 MG (300,		Convulsion	Professional	(Gabapentin)	PS		ORAL
TID), ORAL		Head Injury					
		Platelet Count Decreased		Atenolol (Atenolol)	C		
		Trigeminal Neuralgia		Methylphenobarbital			
				(Methylphenobarbital	C		
)			
				Oxycodone			
				(Oxycodone)	C		
				Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	C		
				Multivitamin "Lappe"			
				(Vitamins Nos)	C		
				Ubidecarenone			
				(Ubidecarenone)	C		

Date:08/21/03ISR Number: 4171800-0Report Type:Direct

Company Report #CTU 200426

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Leukopenia		Gabapentin 300 Mg	PS		ORAL
300 MG TID							
Hospitalization -		Pyrexia					
ORAL							
Initial or Prolonged		Rash					
		Thrombocytopenia					

Date:08/21/03ISR Number: 4173975-6Report Type:Expedited (15-DaCompany Report #2003033661

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Confusional State	Foreign	Neurontin			

Initial or Prolonged 900 MG (TID) Other	Coordination Abnormal Dyspnoea Fall	Health Professional Company Representative	(Gabapentin) Oxycodone Hydrochloride (Oxycodone Hydrochloride) Paracetamol (Paracetamol) Verapamil Hydrochloride (Verapamil Hydrochloride) Dexamethasone (Dexamethasone) Furosemide (Furosemide) Spironolactone (Spironolactone) Potassium Chloride (Potassium Chloride) Simvastatin (Simvastatin) Lansoprazole (Lansoprazole) Nitrazepam	PS SS C C C C C C C C C C C
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Freedom Of Information (FOI) Report

(Nitrazepam) C
 Warfarin (Warfarin) C

Date:08/21/03ISR Number: 4177123-8Report Type:Expedited (15-DaCompany Report #2003007273

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 20 MG	Adrenal Adenoma Ageusia	Consumer Health	Lipitor (Atorvastatin)	PS		ORAL
Other (DAILY), ORAL Required Intervention to 300 MG (TID), Prevent Permanent ORAL Impairment/Damage	Arthralgia Back Pain Dysgeusia Fibromyalgia Glossodynia Headache Hypoaesthesia Insomnia Myalgia Neuropathy Peripheral Paraesthesia Stress Uterine Leiomyoma	Professional	Neurontin (Gabapentin) Ezetimibe (Ezetimibe) Amitriptyline (Amitriptyline) Paracetamol (Paracetamol) Fosinopril Sodium (Fosinopril Sodium) Sertraline Hydrochloride (Sertraline Hydrochloride)	SS C C C		ORAL

Date:08/21/03ISR Number: 4177125-1Report Type:Expedited (15-DaCompany Report #2003029269

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Bone Neoplasm Malignant	Consumer	Neurontin (Gabapentin)	PS		

Date:08/21/03ISR Number: 4177129-9Report Type:Expedited (15-DaCompany Report #2003034005

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiac Failure Congestive	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL Other		Dehydration Pain Spinal Disorder Urinary Incontinence		Risedronate Sodium (Risedronate Sodium)	C		
				Esomeprazole (Esomeprazole)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
				Metoprolol Succinate (Metoprolol Succinate)	C		
				Vicodin (Paracetamol, Hydrocodone Bitartrate)	C		
				Amlodipine Besilate (Amlodipine Besilate)	C		

Freedom Of Information (FOI) Report

Simvastatin
 (Simvastatin) C
 Meclozine
 (Meclozine) C
 Alprazolam
 (Alprazolam) C
 Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C

Date:08/21/03ISR Number: 4177131-7Report Type:Expedited (15-DaCompany Report #2003034004
 Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 300 MG (DAILY), ORAL		Body Temperature Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
0.25 MG (DAILY),		Device Failure Difficulty In Walking Feeling Cold		Triazolam (Triazolam)	SS		
		Insomnia Tremor		Oxycocet (Paracetamol, Oxycodone Hydrochloride) Oxycodone (Oxycodone) Ibuprofen (Ibuprofen) Diazepam (Diazepam) Colchicine (Colchicine) Allopurinol (Allopurinol)	C C C C C C		

Date:08/22/03ISR Number: 4173507-2Report Type:Direct
 Age:65 YR Gender:Male I/FU:I

Company Report #CTU 200541

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State		Gabapentin	PS		
		Palpitations		Bumetanide	C		
		Rash		Felodipine	C		
				Meloxicam	C		
				Potassium Chloride	C		
				Levothyroxine Na (Synthroid)	C		
				Gabapentin	C		
				Loratadine	C		

Date:08/25/03ISR Number: 4172831-7Report Type:Expedited (15-DaCompany Report #PHEH2003US07262
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Hyponatraemia Hypotension		Zelnorm	PS	Novartis Sector: Pharma	
6 mg, UNK							
Other				Trileptal	SS		
				Seroquel	SS		
				Effexor-Xr	SS		
				Neurontin	SS		

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Freedom Of Information (FOI) Report

Hyzaar

SS

Date:08/25/03ISR Number: 4177292-XReport Type:Expedited (15-DaCompany Report #2003030521

Age:76 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness Transient	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (BID),			Professional				
ORAL			Company Representative	All Other Therapeutic Products	C		

Date:08/25/03ISR Number: 4177812-5Report Type:Expedited (15-DaCompany Report #2003034963

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neutropenia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Professional Company Representative				

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,		Maternal Drugs Affecting					
BID), ORAL		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 Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (300, BID), ORAL		Caesarean Section Complications Of Maternal Exposure To Therapeutic Drugs	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG (150, BID), ORAL		Pregnancy Test Positive Uterine Dilation And Curettage		Bupropion Hydrochloride (Bupropion Hydrochloride)	SS		ORAL
				Rizatriptan Benzoate (Rizatriptan Benzoate)	SS		

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Date:08/25/03ISR Number: 4177895-2Report Type:Expedited (15-DaCompany Report #2003034280
Age:39 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion	Consumer	Neurontin			
Initial or Prolonged			(Gabapentin)	PS		
Other			Diazepam (Diazepam)	C		
			Oxycocet			
			(Paracetamol,			
			Oxycodone			
			Hydrochloride)	C		
			Ibuprofen			
			(Ibuprofen)	C		

Date:08/25/03ISR Number: 4177897-6Report Type:Expedited (15-DaCompany Report #2003034184
Age:59 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Hypersensitivity	Consumer	Neurontin			
Initial or Prolonged	Neurosis		(Gabapentin)	PS		
	Pharmaceutical Product		Gabapentin			
	Complaint		(Gabapentin)	SS		
	Stress		Antidepressants	C		

Date:08/25/03ISR Number: 4178538-4Report Type:Expedited (15-DaCompany Report #2003034283
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Abdominal Adhesions	Consumer	Neurontin			
Initial or Prolonged	Colitis Ulcerative		(Gabapentin)	PS		ORAL
3000 MG						
Other	Pain					
(TID), ORAL						
			Quetiapine Fumarate			
			(Quetiapine			
			Fumarate)	C		
			Clonazepam			
			(Clonazepam)	C		

Age:35 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Anorexia
Other	Arthralgia
	Atelectasis
	Bacterial Infection
	Blindness
	Blister
	Blood Testosterone
	Decreased
	Blood Urine
	Bronchial Obstruction
	C-Reactive Protein
	Increased
	Candidiasis
	Chest Pain
	Chest X-Ray Abnormal
	Chills
	Decubitus Ulcer

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Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
800 MG (Q6H), GASTROSTOMY TUBE	Dehydration					
	Diarrhoea					
	Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		
	Dysaesthesia					
	Dyspepsia					
	Dyspnoea					
	Eczema					
	Faecal Incontinence		Amitriptyline (Amitriptyline)	C		
	Fatigue		Fluoxetine Hydrochloride			
	Gastritis Viral		(Fluoxetine Hydrochloride)	C		
	Guillain-Barre Syndrome		Rofecoxib (Rofecoxib)	C		
	Hallucination, Visual		Docosate Sodium (Docosate Sodium)	C		
	Headache		Bisacodyl (Bisacodyl)	C		
	Iron Deficiency Anaemia		All Other Therapeutic Products	C		
	Ketoacidosis		Testosterone Cipionate			
	Lip Disorder		(Testosterone Cipionate)	C		
	Mass		Warfarin Sodium (Warfarin Sodium)	C		
	Metabolic Acidosis		Salbutamol (Salbutamol)_	C		
	Nausea					
	Neck Pain					
	Nephrolithiasis					
Neutrophil Percentage Decreased						
Oedema Peripheral						
Optic Neuritis						
Pain						
Pain In Extremity						
Paraesthesia						
Platelet Count Increased						
Pneumonia						
Pruritus						
Pyrexia						
Rash						
Rash Maculo-Papular						
Rash Pustular						
Red Blood Cell						
Sedimentation Rate Increased						
Sensation Of Pressure						
Sinus Tachycardia						
Skin Ulcer						
Staphylococcal Infection						
Swelling						

Swelling Face
Tongue Disorder
Toothache
Trismus
Ulcer
Urethral Disorder
Urinary Incontinence
Urinary Tract Infection
Visual Acuity Reduced
Vomiting

Date:08/26/03ISR Number: 4178507-4Report Type:Expedited (15-DaCompany Report #2003005182
Age:62 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
ORAL		Condition Aggravated Hepatic Steatosis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Hepatitis Hepatomegaly	Professional	Clomipramine (Clomipramine)	C		
				Bloprss Plus (Hydrochlorothiazide , Candesartan Cilexetil)	C		
				Pravastatin (Pravastatin)	C		
				Atenolol (Atenolol)	C		
				Clopidogrel (Clopidogrel)	C		
				Urapidil (Urapidil)	C		

Date:08/26/03ISR Number: 4179704-4Report Type:Expedited (15-DaCompany Report #2003028825
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Blood Cholesterol Increased	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Blood Triglycerides Condition Aggravated Euphoric Mood Foot Fracture Headache Lipids Abnormal Weight Increased	Professional	Pravastatin Sodium (Pravastatin Sodium)	C		
				Tamoxifen (Tamoxifen)	C		
				Pantoprazole (Pantoprazole)	C		
				Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		
				Fioricet W/Codeine (Codeine Phosphate, Caffeine, Butalbital, Paracetamol)	C		

Date:08/26/03ISR Number: 4179706-8Report Type:Expedited (15-DaCompany Report #2003034181
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ejaculation Failure Surgery	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:08/26/03ISR Number: 4179707-XReport Type:Expedited (15-DaCompany Report #2003032836
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Other	Constipation Depression Dyspepsia Flatulence Herpes Zoster

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hiatus Hernia Irritability Mental Status Changes	Consumer Health	Lipitor (Atorvastatin)	PS		ORAL
DAILY, ORAL		Muscle Spasms Neuropathy	Professional	Neurontin (Gabapentin)	SS		ORAL
900 MG (THREE TIMES A DAY), ORAL		Osteoporosis Prostate Cancer		Alendronate Sodium (Alendronate Sodium)	SS		
				Fentanyl (Fentanyl)	C		
				Paracetamol (Paracetamol)	C		
				Oxygen (Oxygen)	C		
				Salbutamol (Salbutamol)	C		
				Ipratropium Bromide (Ipratropium Bromide)	C		
				Leuprorelin Acetate (Leuprorelin Acetate)	C		

Date:08/26/03ISR Number: 4179708-1Report Type:Expedited (15-DaCompany Report #2003034528

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Drug Level Below Therapeutic	Health Professional	Neurontin (Gabapentin)	PS		
		Pain		All Other Therapeutic Products	C		
				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
				Hydrocodone (Hydrocodone)	C		

Propacet
 (Paracetamol,
 Dextropropoxyphene
 Napsilate) C
 Arthrotex
 (Diclofenac Sodium,
 Misoprostol) C

Date:08/27/03ISR Number: 4180533-6Report Type:Expedited (15-DaCompany Report #DSA_23189-2003
 Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 300 MG PER Hospitalization - 24HR PO Initial or Prolonged		Acute Pulmonary Oedema	Foreign Health	Mono-Tildiem	PS		ORAL
			Professional Other	Euphylline Neurontin Deroxat Mopral Skenan	SS SS SS SS SS		ORAL
20 MG PER 24HR PO							

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Date:08/27/03ISR Number: 4180917-6Report Type:Expedited (15-DaCompany Report #2003034727

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Sinusitis					
900 MG (TID),							
ORAL				Pravastatin Sodium (Pravastatin Sodium)	C		
				Cimetidine (Cimetidine)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		

Date:08/27/03ISR Number: 4181248-0Report Type:Expedited (15-DaCompany Report #2003034659

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Malaise	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		
Other		Mental Disorder					

Date:08/27/03ISR Number: 4181254-6Report Type:Expedited (15-DaCompany Report #2003034840

Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
Other		Disorientation Memory Impairment					
200 MG DAILY			Professional				
ORAL		Speech Disorder		Rofecoxib	C		
				Acetylsalicylic Acid	C		
				Tocopherol	C		

Date:08/28/03ISR Number: 4180896-1Report Type:Expedited (15-DaCompany Report #2003034902

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 3200 MG, ORAL Other	Hospitalisation	Health Professional	Neurontin (Gabapentin) All Other Therapeutic Products	PS SS		ORAL

Date:08/28/03ISR Number: 4180970-XReport Type:Expedited (15-DaCompany Report #2003034837

Age:55 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 100 MG DAILY ORAL	Anxiety Muscle Spasms Muscle Tightness Sedation Somnolence	Consumer	Neurontin (Gabapentin) Nefazodone Hydrochloride (Nefazodone Hydrochloride)	PS C		ORAL

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Freedom Of Information (FOI) Report

Clonazepam (Clonazepam)	C
Olanzapine (Olanzapine)	C
Esomeprazole (Esomeprazole)	C
Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C

Date:08/28/03ISR Number: 4180977-2Report Type:Expedited (15-DaCompany Report #2003031373
Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 4500 MG	Grand Mal Convulsion Hyponatraemia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other (1500, TID)	Syncope					
ORAL			Oxcarbazepine (Oxcarbazepine)	SS		ORAL
(300, BID)						
ORAL			Terazosin Hydrochloride (Terazosin Hydrochloride)	C		
			Loratadine (Loratadine)	C		
			Omeprazole (Omeprazole)	C		
			Modafinil (Modafinil)	C		
			Estrogens Conjugated (Estrogens Conjugated)	C		
			Rofecoxib (Rofecoxib)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG Other (TID), ORAL		Neuropathic Pain	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
				Tramadol Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 30 MG QD PO Initial or Prolonged 30 MG QD PO		Hypotension	Health Professional	Avinza Avinza	PS SS		ORAL ORAL
300 MG QD PO				Gabapentin Gabapentin	SS C		ORAL ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/02/03ISR Number: 4178856-XReport Type:Direct
 Age:46 YR Gender:Male I/FU:I

Company Report #CTU 201097

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Neurontin 100 Mg	PS		ORAL
100 MG 3X							
DAILY ORAL							

Date:09/02/03ISR Number: 4183268-9Report Type:Expedited (15-DaCompany Report #DEU-2003-0000549
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased Drug Interaction Skin Exfoliation	Foreign Consumer Other	Oxygesic 10 Mg (Oxycodone Hydrochloride) Cr Tablet	PS		ORAL
10 MG, BID,							
ORAL	61 DAY						
				Mtx (Methotrexate Sodium)	SS		
				Neurontin (Gabapentin)	SS		
61 DAY							

Date:09/02/03ISR Number: 4183442-1Report Type:Expedited (15-DaCompany Report #2003030954
 Age:94 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Coordination Abnormal	Consumer	Neurontin (Tablets)			
Other		Drug Ineffective	Health	(Gabapentin)	PS		ORAL
200 MG (BID),							
ORAL		Dysarthria	Professional				
		Mental Impairment		Atenolol (Atenolol)	C		
		Oedema Peripheral		Nifedipine			
		Sedation		(Nifedipine)	C		
		Sleep Disorder		Levothyroxine Sodium (Levothyroxine			

Sodium) C
 Furosemide C
 (Furosemide)
 Prednisone C
 (Prednisone)
 Alendronate Sodium C
 (Alendronate Sodium)
 Combivent
 (Ipratropium
 Bromide, Salbutamol
 Sulfate) C

Date:09/02/03ISR Number: 4183536-0Report Type:Expedited (15-DaCompany Report #2003035920
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Brain Abscess	Consumer	Neurontin (Tablets)			
Initial or Prolonged		Difficulty In Walking		(Gabapentin)	PS		
Other		Extradural Abscess		All Other			
		Pain		Therapeutic Products	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/02/03ISR Number: 4183538-4Report Type:Expedited (15-DaCompany Report #2003035343
 Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	900 MG (QID), ORAL	Catheterisation Cardiac Confusional State Feeling Abnormal Feeling Cold Flat Affect Heart Rate Decreased Mental Impairment	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:09/02/03ISR Number: 4183662-6Report Type:Expedited (15-DaCompany Report #USA-2002-0001677
 Age:41 YR Gender:Female I/FU:F

Outcome	PT
Death	Alopecia
Hospitalization - Initial or Prolonged	Angiopathy Anxiety Asthenia Back Pain Cardiac Enzymes Increased Chest Pain Coma Depressed Mood Diabetes Mellitus Disturbance In Attention Drug Ineffective Dyspnoea Emphysema Fatigue Finger Deformity Flushing Hilar Lymphadenopathy Hyperglycaemia Hyperhidrosis Hyperventilation Hypoxia Influenza Insomnia Irritability Libido Decreased

Lobar Pneumonia
Lung Disorder
Nausea
Nervousness
Night Sweats
Overdose
Pain
Peripheral Occlusive
Disease
Pleurisy
Portal Triaditis
Pruritus
Pulmonary Congestion
Pulmonary Oedema
Pulse Absent
Rash Pruritic
Rash Scaly
Somnolence
Troponin Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Upper Respiratory Tract Infection Urinary Incontinence Weight Increased	Report Source	Product	Role	Manufacturer	Route
10 MG			Consumer Health Professional Other	Oxycontin Tablets 10 Mg(Oxycodone Hydrochloride) Cr Tablet	PS		
20 MG	1008 DAY			Oxycontin Tablets 20 Mg (Oxycodone Hydrochloride) Cr Tablet	SS		
1008 DAY				Hydrocodone Bitartrate(Similar To Ind 59,175)(Hydrocodone Bitartrate)	SS		
1008 DAY				Alprazolam(Alprazolam)	SS		
1008 DAY				Ephedrine(Ephedrine)	SS		
1008 DAY				Pseudoephedrine(Pseudoephedrine)	SS		
75 MG, DAILY	1008 DAY			Effexor (Venlafaxine Hydrochloride)	SS		
1008 DAY				Codeine (Codeine)	SS		
100 MG, TID	1008 DAY			Neurontin (Gabapentin)	SS		
1008 DAY				Metoclopramide (Metoclopramide)	SS		
1008 DAY				Quetiapine (Quetiapine)	SS		
				Claritin (Loratadine)	C		
				Vitamin C (Ascorbic Acid)	C		
				Celebrex (Celecoxib)	C		

Prilosec	
(Omeprazole)	C
Daypro (Oxaprozin)	C
Trandate (Labetalol	
Hydrochloride)	C
Axid (Nizatidine)	C
Paxil (Paroxetine	
Hydrochloride)	C
Medrol	
(Methylprednisolone)	C
Macro Antioxidant	
(Ascorbic Acid,	
Cystine, Tocopherol,	
Calcium Ascorbate,	
Betacarotene,	C
Zocor (Simvastatin)	C
Risperidal	
(Risperidone)	C
Fioricet	
(Butalbital)	C
Flonase (Fluticasone	
Propionate)	C
Phenobarbital	
(Phenobarbital)	C
Donnatal (Atropine	
Sulfate, Hyoscine	

Freedom Of Information (FOI) Report

Hydrobromide, Hyoscyamine Sulfate, Phenobarbital)	C
Bellergal-S (Belladonna Alkaloids, Ergotamine Tartrate, Phenobarbital)	C
Weight Loss Supplement (Does Not Code)	C
Ibuprofen	C
Ambien (Zolpidem Tartrate)	C
Baclofen (Baclofen)	C
Trandate (Labetalol Hydrochloride)	C
Celexa (Citalopram Hydrobromide)	C
Cortisone (Cortisone)	C
Atarax (Hydroxyzine Hydrochloride)	C
Proventil Tablet (Salbutamol Sulfate)	C
Amitriptyline (Amitriptyline)	C
Cyclobenzaprine (Cyclobenzaprine)	C
Augmentin (Amoxicillin Trihydrate, Clavulanate Potassium)	C
Ergobel (Nicergoline)	C
Glucotrol XL (Glipizide) Tablet	C
Humuline Nph (Insulin Human Injection, Isophane) Injectable	C
Capoten Tablet	C
Albuterol (Salbutamol) Inhaler	C
Azmacort (Triamcinolone Acetonide) Inhaler	C

Aciphex (Rabeprazole
Sodium) Tablet C
Seroquel
(Quetiapine) Tablet C

Date:09/02/03ISR Number: 4184743-3Report Type:Expedited (15-DaCompany Report #2003022679

Age: Gender:Female I/FU:F

Outcome PT
Hospitalization - Fall
Initial or Prolonged Hypotension
Malaise
Mydriasis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Somnolence

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG ORAL		Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Professional Company Representative	Loprazolam All Other Therapeutic Products	C		

Date:09/03/03ISR Number: 4184833-5Report Type:Expedited (15-DaCompany Report #2003027782
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	3600 MG (DAILY), ORAL	Conversion Disorder Drug Tolerance	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Medication Error		Fentanyl (Fentanyl)	C		
		Pain		Ramipril (Ramipril)	C		
		Paraesthesia		Allopurinol (Allopurinol)	C		
		Self-Medication		Acetazolamide (Acetazolamide)	C		
				Sodium Bicarbonate (Sodium Bicarbonate)	C		
				Colchicine (Colchicine)	C		
				Tolterodine L-Tartrate (Tolterodine L-Tartrate)	C		
				Calcium (Calcium)	C		
				Hydrochlorothiazide (Hydrochlorothiazide)	C		
				Pethidine Hydrochloride (Pethidine Hydrochloride)	C		
				Montelukast Sodium (Montelukast Sodium)	C		

Ranitidine	
Hydrochloride	
(Ranitidine	
Hydrochloride)	C
Cetirizine	
Hydrochloride	
(Cetirizine	
Hydrochloride)	C
Diphenhydramine	
Hydrochloride	
(Diphenhydramine	
Hydrochloride)	C
Salmeterol Xinafoate	
(Salmeterol	
Xinafoate)	C
Fluticasone	
Propionate	
(Fluticasone	
Propionate)	C
Salbutamol	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Salbutamol)	C
Epinephrine	
(Epinephrine)	C
Prednisone	
(Prednisone)	C
Vicodin	
(Paracetamol,	
Hydrocodone	
Bitartrate)	C

Date:09/03/03ISR Number: 4184954-7Report Type:Expedited (15-DaCompany Report #2003032895
 Age:40 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 100 MG (PRN), Initial or Prolonged ORAL	Diabetes Mellitus	Consumer	Viagra (Sildenafil)	PS		ORAL
Other 600 MG (BID), ORAL	Drug Ineffective Drug Interaction Feeling Abnormal Feeling Hot	Health Professional	Neurontin (Gabapentin)	SS		ORAL
	Flushing Neuropathy Pancreatitis Pharmaceutical Product Complaint		Enalapril (Enalapril)	C		

Date:09/03/03ISR Number: 4184956-0Report Type:Expedited (15-DaCompany Report #2003026124
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Death Other 1800 MG (TID)	Depressed Level Of Consciousness	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		
		Professional				

Date:09/04/03ISR Number: 4178982-5Report Type:Expedited (15-DaCompany Report #FR-ROCHE-345497
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Withdrawal Syndrome Status Epilepticus		Rivotril Neurontin Neurontin Sedative	PS SS SS C	Roche	ORAL ORAL ORAL
UNKNOWN		7 DAY					

Date:09/04/03ISR Number: 4179191-6Report Type:Expedited (15-DaCompany Report #WAES 0308USA02664
Age:78 YR Gender:Male I/FU:I

Outcome	PT
Death	Acidosis
Hospitalization - Initial or Prolonged	Anion Gap Increased Blood Creatinine Increased Blood Ph Decreased Bradycardia Cardiac Arrest Completed Suicide Diarrhoea Electrolyte Imbalance Hypoglycaemia Hypotension

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route

Date:09/04/03ISR Number: 4181142-5Report Type:Direct Company Report #USP 042234
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
CAPSULE							

Date:09/04/03ISR Number: 4183802-9Report Type:Expedited (15-DaCompany Report #2003036042
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
Required							
ORAL							

Date:09/04/03ISR Number: 4184984-5Report Type:Expedited (15-DaCompany Report #2003035672
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Neurontin			
Other		Basedow'S Disease		(Gabapentin)	PS		
		Emotional Disorder					
		Feeling Drunk					

Date:09/05/03ISR Number: 4180205-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0410183A
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety		Requip	PS	Glaxosmithkline	ORAL
50MG See		Chest Pain					
dosage text	1 YR	Disorientation		Neurontin	SS		
UNKNOWN		Pyrexia		Effexor	SS		
UNKNOWN		Tremor		Serzone	SS		
UNKNOWN				Flexeril	SS		
UNKNOWN				Adderall	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/05/03ISR Number: 4185128-6Report Type:Expedited (15-DaCompany Report #2003035747
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300 MG DAILY	Eye Rolling Loss Of Consciousness	Consumer	Neurontin (Gabapentin)	PS		
				Propacet (Paracetamol, Dextropropoxyphene Napsilate)	SS		
				Esomperazole (Esomeprazole)	C		
				Sucralfate (Sucralfate)	C		
				Paracetamol (Paracetamol)	C		

Date:09/05/03ISR Number: 4185131-6Report Type:Expedited (15-DaCompany Report #2003031542
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged DAILY, ORAL		Ankle Fracture Compression Fracture	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other		Fall Sensory Loss Spinal Compression Fracture	Professional	Vaseretic (Hydrochlorothiazide , Enalapril Maleate) Oxycocet (Paracetamol, Oxycodone Hydrochloride)	C		

Date:09/08/03ISR Number: 4183524-4Report Type:Direct Company Report #CTU 201414
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	4 @ NIGHT 2 @	Abnormal Dreams		Neurontin 100 Mg	PS		

MORING

Feeling Abnormal
Insomnia
Nervous System Disorder
Pain
Psychotic Disorder
Sleep Terror
Stress
Thinking Abnormal

Sulfasalazine C
Indomethacin C
Clonazepam C
Luvox C
Clomipramine C
Ranitidine C
Lotensin C

Date:09/08/03ISR Number: 4185655-1Report Type:Expedited (15-DaCompany Report #2003036328
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	900 MG (TID),	Paraesthesia Sensory Loss	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
ORAL				Levothyroxine (Levothyroxine)	C		
				Diclofenac (Diclofenac)	C		
				Kliogest "Novo Industri" (Estradiol,			

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Norethisterone
Acetate) C

Date:09/08/03ISR Number: 4185664-2Report Type:Expedited (15-DaCompany Report #2003035893
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Chest Injury Confusional State	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Haemothorax Status Epilepticus	Professional	Clonazepam (Clonazepam)	SS		ORAL

Date:09/08/03ISR Number: 4185807-0Report Type:Expedited (15-DaCompany Report #2003031759
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG (DAILY), ORAL		Abnormal Behaviour Alopecia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
2400 MG (DAILY), ORAL		Colitis Ulcerative Depressed Level Of Consciousness		Ibuprofen (Ibuprofen)	SS		ORAL
		Disturbance In Attention Dizziness Drug Ineffective Drug Interaction Intervertebral Disc Protrusion Medication Error Pain In Extremity Road Traffic Accident		Valproate Semisodium (Valproate Semisodium) Acetylsalicylic Acid (Acetylsalicylic Acid) Mesalazine (Mesalazine) Glycopyrronium Bromide (Glycopyrronium Bromide) Propranolol Hydrochloride	SS SS C		

(Propranolol
Hydrochloride) C
Amitriptyline
Hydrochloride
(Amitriptyline
Hydrochloride) C

Date:09/08/03ISR Number: 4186059-8Report Type:Expedited (15-DaCompany Report #2003150492US
Age:38 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Accident
Initial or Prolonged	Arthritis
Other	Eye Disorder
	Eye Movement Disorder
	Fibromyalgia
	Herpes Zoster
	Intervertebral Disc Degeneration
	Intervertebral Disc Protrusion
	Nerve Compression
	Nerve Injury

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FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	Neuralgia Neuropathy Peripheral Oedema Peripheral	Report Source	Product	Role	Manufacturer	Route
200 MG, BID, ORAL		Tremor	Consumer	Celebrex (Celecoxib)	PS		ORAL
1500MG/DAILY, ORAL				Neurontin (Gabapentin)	SS		ORAL
175 MG, ORAL EPIDURAL	EPIDURAL			Topamax (Topiramate)	SS		ORAL
				Unspecified Drug()	SS		
				Luvox	C		
				Xanax	C		
				Skelaxin (Metaxolone)	C		
				Prozac (Fluoxetine Hydrochloride)	C		

Date:09/08/03ISR Number: 4186486-9Report Type:Expedited (15-DaCompany Report #2003036045

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other (BID), ORAL		Burning Sensation Cardiac Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Condition Aggravated Coronary Artery Occlusion		Carbamazepine (Carbamazepine)	C		
		Epistaxis		All Other			
		Hypoaesthesia		Therapeutic Products	C		
		Insomnia					
		Trigeminal Neuralgia					

Date:09/08/03ISR Number: 4186619-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome PT

Hospitalization -
Initial or Prolonged
Disability
Other

Abdominal Pain Upper
Abdominal Tenderness
Acrochordon
Actinic Keratosis
Albumin Globulin Ratio
Amnesia
Anxiety
Arthralgia
Arthropathy
Blood Calcium Decreased
Blood Cholesterol
Increased
Blood Glucose Increased
Blood Triglycerides
Increased
Blood Urea
Nitrogen/Creatinine Ratio
Increased
Brain Neoplasm
Burning Sensation
Cardiac Arrest
Chest Pain
Cholecystitis
Cognitive Disorder
Coma

Freedom Of Information (FOI) Report

Conversion Disorder
Convulsion
Coordination Abnormal
Deafness
Depressed Level Of
Consciousness
Depression
Dermal Cyst
Diplopia
Disturbance In Attention
Dizziness
Drug Level Above
Therapeutic
Drug Level Increased
Dysphagia
Dysphemia
Dyspnoea
Emotional Distress
Eosinophil Count
Increased
Erectile Dysfunction
Erythema
Exostosis
Fall
Fatigue
Feeling Abnormal
Fibromyalgia
Gallbladder Disorder
Gastroesophageal Reflux
Disease
Gynaecomastia
Haematocrit Decreased
Headache
Hearing Impaired
Hepatic Steatosis
High Density Lipoprotein
Decreased
Hypoalbuminaemia
Hypoglycaemia
Insomnia
Learning Disorder
Limb Injury
Memory Impairment
Meniscus Lesion
Monocyte Count Increased
Nausea
Neurological Examination

Abnormal
Nuchal Rigidity
Oedema
Osteoarthritis
Osteopenia
Pain
Patellofemoral Pain
Syndrome
Pigmented Naevus
Red Blood Cell Count
Decreased
Red Blood Cell
Sedimentation Rate
Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (BID), ORAL		Respiratory Arrest Spinal X-Ray Abnormal Sports Injury Tension Tinnitus Toe Deformity Tremor Vision Blurred Vomiting X-Ray Limb Abnormal	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin)	PS SS C C C C		ORAL

Date:09/09/03ISR Number: 4188269-2Report Type:Expedited (15-DaCompany Report #2003036352

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Anxiety Condition Aggravated Depression Diarrhoea Hypertension Multiple Sclerosis Nausea Oesophageal Spasm Pain Photophobia Weight Decreased	Consumer	Neurontin (Gabapentin) Prozac (Fluoxetine Hydrochloride) Baclofen (Baclofen)	PS C C		ORAL

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiac Failure	Health	Neurontin			
Other		Congestive	Professional	(Gabapentin)	PS		
300 MG (TID)		Condition Aggravated		Sucralfate			
		Fluid Retention		(Sucralfate)	C		
		Treatment Noncompliance		Omeprazole	C		
				(Omeprazole)			
				Docusate Calcium			
				(Docusate Calcium)	C		
				Prochlorperazine			
				Edisylate			
				(Prochlorperazine			
				Edisylate)	C		
				Amlodipine Besilate			
				(Amlodipine			
				Besilate)	C		
				Metoprolol Succinate			
				(Metoprolol			
				Succinate)	C		

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Freedom Of Information (FOI) Report

Isosorbide
 Mononitrate
 (Isosorbide
 Mononitrate) C
 Glyceryl Trinitrate
 (Glyceryl
 Trinitrate) C
 Fentanyl (Fentanyl) C
 Vicodin
 (Paracetamol,
 Hydrocodone
 Bitartrate) C
 Ibuprofen
 (Ibuprofen) C
 Paroxetine
 Hydrochloride
 (Paroxetine
 Hydrochloride) C
 Cetirizine
 Hydrochloride
 (Cetirizine
 Hydrochloride) C
 Cyanocobalamin
 (Cyanocobalamin) C
 Zolpidem Tartrate
 (Zolpidem Tartrate) C
 Oxygen (Oxygen) C
 Nystatin (Nystatin) C
 Polytrim
 (Trimethoprim,
 Polymyxin B Sulfate) C

Date:09/10/03ISR Number: 4188629-XReport Type:Expedited (15-DaCompany Report #2003036714
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Lacrimation Increased Vision Blurred	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG		Visual Acuity Reduced					
(DAILY), ORAL				Candesartan Cilexetil (Candesartan			

Cilexetil)	C
Omeprazole	
(Omeprazole)	C
Dorzolamide	
Hydrochloride	
(Dorzolamide	
Hydrochloride)	C
Brimonidine Tartrate	
(Brimonidine	
Tartrate)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/10/03ISR Number: 4188630-6Report Type:Expedited (15-DaCompany Report #2003036715

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2000 MG Other (DAILY)		Cardiac Failure Circulatory Collapse Fall Myocardial Infarction Neuropathy	Consumer	Neurontin (Gabapentin) Metformin Hydrochloride (Metformin Hydrochloride)	PS C		

Date:09/10/03ISR Number: 4188631-8Report Type:Expedited (15-DaCompany Report #2003034149

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (300, TID), ORAL		Benign Breast Neoplasm Convulsion Dysplasia Fibrocystic Breast Disease Head Injury Platelet Count Decreased Trigeminal Neuralgia	Health Professional	Neurontin (Gabapentin) Atenolol (Atenolol) Methylphenobarbital (Methylphenobarbital) Oxycodone (Oxycodone) Oxycodone Hydrochloride (Oxycodone Hydrochloride) Multivitamin (Vitamins Nos) Ubidecarenone (Ubidecarenone)	PS C C C C C C		ORAL Lappe

Date:09/10/03ISR Number: 4189060-3Report Type:Expedited (15-DaCompany Report #2003036800

Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG		Carpal Tunnel Syndrome Surgery	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
(DAILY), ORAL				All Other Therapeutic Products	C		

Date:09/11/03ISR Number: 4189388-7Report Type:Expedited (15-DaCompany Report #2003036790
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG		Anxiety Chest Discomfort	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL		Circulatory Collapse	Professional				
		Tachycardia	Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/11/03ISR Number: 4189395-4Report Type:Expedited (15-DaCompany Report #2003037044

Age:81 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (TID), ORAL	Peripheral Nerve Operation	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
			Colestyramine (Colestyramine)	C		
			Gliclazide (Gliclazide)	C		
			Atenolol (Atenolol)	C		
			Amlodipine (Amlodipine)	C		

Date:09/11/03ISR Number: 4189987-2Report Type:Expedited (15-DaCompany Report #2003030108

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL 120 MG, ORAL	Dizziness Hypotension Prostate Cancer	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Propranolol Hydrochloride (Propranolol Hydrochloride)	SS		ORAL
			Valproate Semisodium (Valproate Semisodium)	C		

Date:09/11/03ISR Number: 4189991-4Report Type:Expedited (15-DaCompany Report #2003037078

Age:1 DY Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Disability 1600 MG	Anorectal Disorder Cardiac Murmur	Consumer	Neurontin (Gabapentin)	PS		ORAL

Congenital Anomaly	Dyspnoea
(BID), ORAL	
Other	Feeding Problem In Newborn Intercostal Retraction Maternal Drugs Affecting Foetus Neonatal Disorder Respiration Abnormal Respiratory Arrest Tracheomalacia

Date:09/11/03ISR Number: 4200786-5Report Type:Periodic Company Report #2003164678US
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNK, UNK;		Asthenia Difficulty In Walking	Health Professional	Bextra (Valdecoxib) Tablet	PS		ORAL
ORAL		Flushing					
UNKNOWN	UNK, UNK, UNK	Muscle Twitching Myalgia		Celebrex (Celecoxib, Celecoxib) Capsule	SS		
UNKNOWN	300 UNK,	Nausea Renal Failure Acute		Viread (Tenofovir)	SS		
UNK, UNK							
UNKNOWN	UNK, UNK, UNK	Somnolence Vomiting		Fortovase (Saquinavir)	SS		

Freedom Of Information (FOI) Report

UNK, UNK, UNK		Zerit (Stavudine)	SS
UNKNOWN	UNK, UNK, UNK	Kaletra (Lopinavir/Ritonavir)	SS
UNKNOWN	UNK, UNK, UNK	Phenergan (Promethazine)	SS
UNKNOWN	UNK, UNK, UNK	Ambien (Zolpidem Tartrate)	SS
UNKNOWN	UNK, UNK, UNK	Ensure (Vitamins Nos)	SS
UNKNOWN	UNK, UNK, UNK	Tricor (Fenofibrate)	SS
UNKNOWN	UNK, UNK, UNK	Dulcolax (Bisacodyl)	SS
UNK, UNK, UNK		Marinol (Dronabinol)	SS
UNKNOWN	UNK, UNK, UNK	Soma (Carisoprodol)	SS
UNKNOWN	UNK, UNK, UNK	Bactrim Ds (Sulfamethoxazole Trimethoprim)	SS
UNKNOWN	UNK, UNK, UNK	Biaxin (Clarithromycin)	SS
UNKNOWN	UNK, UNK, UNK	Protonix ()	SS
UNKNOWN	UNK, UNK, UNK	Oxandrin (Oxlandrolone)	SS
UNKNOWN	UNK, UNK, UNK	Neurontin (Gabapentin)	SS
UNKNOWN	UNK, UNK, UNK	Colchicine (Colchicine)	SS

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dysphagia	Health Professional	Neurontin (Gabapentin)	PS		
200 MG, BID							

Date:09/12/03ISR Number: 4190224-3Report Type:Expedited (15-DaCompany Report #2003033661
 Age:81 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Confusional State Coordination Abnormal	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other		Dyspnoea	Professional				
ORAL		Fall Rib Fracture	Company Representative	Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		
15 MG (TID)							
				Paracetamol (Paracetamol)	C		
				Verapamil Hydrochloride (Verapamil Hydrochloride)	C		
				Dexamethasone (Dexamethasone)	C		
				Furosemide (Furosemide)	C		
				Spironolactone (Spironolactone)	C		
				Potassium Chloride (Potassium Chloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Simvastatin
 (Simvastatin) C
 Lansoprazole
 (Lansoprazole) C
 Nitrazepam
 (Nitrazepam) C
 Warfarin (Warfarn) C

Date:09/16/03ISR Number: 4187540-8Report Type:Expedited (15-DaCompany Report #WAES 0309USA01127
 Age:36 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization - Initial or Prolonged	Alanine Aminotransferase Increased Blood Alkaline Phosphatase Increased		Indocin (Indomethacin) Neurontin Copaxone	PS SS SS		ORAL
570 DAY						
	Drug Interaction Pancreatitis		Copaxone Elavil Actiq Oxycontin	SS C C C	Merck & Co., Inc	ORAL

Date:09/16/03ISR Number: 4190774-XReport Type:Expedited (15-DaCompany Report #2003037152
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Death Other	Death	Health Professional Company Representative	Neurontin (Gabapentin) All Other Therapeutic Products Ethanol (Ethanol)	PS SS SS		

Date:09/16/03ISR Number: 4190781-7Report Type:Expedited (15-DaCompany Report #2003008068
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Disability Other	Activities Of Daily Living Impaired	Consumer	Neurontin (Gabapentin)	PS		

Arthralgia	Acetylsalicylic Acid	
Balance Disorder	(Acetylsalicylic	
Contusion	Acid)	SS
Disturbance In Attention	Vitamins With	
Dry Mouth	Minerals	SS
Fall		
Headache		
Mental Impairment		
Muscle Spasms		
Nerve Injury		
Nervousness		
Pain		
Panic Attack		
Paresis		
Thermal Burn		
Walking Aid User		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/16/03ISR Number: 4190824-0Report Type:Expedited (15-DaCompany Report #2003037153

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional Company Representative	Neurontin (Gabapentin) All Other Therapeutic Products	PS		
Other					SS		

Date:09/16/03ISR Number: 4191276-7Report Type:Expedited (15-DaCompany Report #2003037418

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Acne Intracranial Pressure Increased Weight Increased	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:09/17/03ISR Number: 4191438-9Report Type:Expedited (15-DaCompany Report #2003027172

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Arthralgia Dizziness Extremity Contracture	Foreign Consumer Health	Neurontin (Gabapentin) (See Image)	PS		ORAL
600 MG (BID);							
ORAL		Fear Of Disease	Professional				
		Hyperaesthesia Joint Stiffness Pain In Extremity Paraesthesia Paralysis					

Date:09/17/03ISR Number: 4192914-5Report Type:Expedited (15-DaCompany Report #2003037445

Age:24 MON Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 10 MG		Abnormal Behaviour Medication Error	Consumer	Doxepin (Conc.) (Doxepin)	PS		ORAL
(DAILY), ORAL		Sleep Disorder					
1.5 ML (TID), ORAL		Staphylococcal Infection		Neurontin (Ped Oral Susp) (Gabepentin)	SS		ORAL

Date:09/22/03ISR Number: 4193762-2Report Type:Expedited (15-DaCompany Report #2003030174
Age:35 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Other	Abdominal Pain Anorexia Blindness Blood Blister Blood Urine Chills Culture Urine Culture Urine Positive Dehydration Dyspepsia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
800 MG (Q6H), GASTROSTOMY TUBE)		Consumer	Neurontin (Gabapentin)	PS		

Date:09/22/03ISR Number: 4194023-8Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 202190

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose Other 100MG TID ORAL		PT	Gabapentin 100mg	PS		ORAL

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (TID), Other ORAL	Blood Potassium Decreased Blood Pressure Decreased Dehydration	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL	Dizziness Drug Ineffective Hyperaesthesia Pain		Vicodin (Paracetamol, Hydrocodone Bitartrate)	SS		ORAL
ORAL	Red Blood Cell Sedimentation Rate Increased Sensory Disturbance		Oxycocet (Paracetamol, Oxycodone Hydrochloride)	SS		ORAL
	Skin Burning Sensation Syncope Vomiting		Panadeine Co (Codeine Phosphate, Paracetamol) Lisinopril (Lisinopril) Hydrochlorothiazide (Hydrochlorothiazide)	SS C		

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Freedom Of Information (FOI) Report

) C
 Valdecoxib
 (Valdecoxib) C

Date:09/22/03ISR Number: 4194278-XReport Type:Expedited (15-DaCompany Report #2003038101

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 (THREE TIMES A DAY), ORAL		Balance Disorder Coordination Abnormal Drug Toxicity Fall Feeling Abnormal Pain Rash Erythematous Rash Maculo-Papular	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Atorvastatin (Atorvastatin)	C		
				Valproate Semisodium (Valproate Semisodium)	C		
				Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		
				Amlodipine Besilate (Amlodipine Besilate)	C		
				Enalapril Maleate (Enalapril Maleate)	C		
				Bupropion Hydrochloride (Bupropion Hydrochloride)	C		
				Spironolactone (Spironolactone)	C		
				Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
				Furosemide (Furosemide)	C		
				Mirtazapine (Mirtazapine)	C		
				Temazepam (Temazepam)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder	Consumer	Neurontin			
2400 MG		Coordination Abnormal		(Gabepentin)	PS		ORAL
(DAILY), ORAL		Drug Toxicity					
		Fall		Antihypertensives	C		
		Rash Erythematous		Valproate Semisodium			
		Rash Maculo-Papular		(Valproate			
		Renal Cell Carcinoma		Semisodium)	C		
		Stage Unspecified		Mirtazapine			
				(Mirtazapine)	C		
				All Other			
				Therapeutic Products	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/22/03ISR Number: 4194730-7Report Type:Expedited (15-DaCompany Report #USA-2002-0002855

Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Asthenia Blood Creatine Phosphokinase	Consumer Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride)Cr Tablet			ORAL
MG, ORAL; MG, ORAL		Blood Creatine Phosphokinase Mb Cardiac Failure Congestive Chronic Obstructive		Oxycontin Tablets 20 Mg (Oxycodone Hydrochloride) Cr Tablet	PS		ORAL
20 GM, ORAL; 40 MG, ORAL		Pulmonary Disease Coagulopathy Coma Confusional State Decreased Appetite		Hydrocodone Cp (Phenylephrine, Chlorphenamine Maleate) Syrup	SS		ORAL
ORAL		Depression Disturbance In Attention		Darvocet-N (Paracetam ol) Tablet	SS		ORAL
ORAL	7 DAY	Drug Dependence Drug Withdrawal Syndrome		Cyclobenzaprine (Cycl obenzaprine) Tablet	SS		ORAL
ORAL		Dysuria Encephalopathy Fall		Hydroxyzine Pamoate (Hydroxyzine Embonate)	SS		ORAL
MG, ORAL		Headache Hepatic Necrosis		Valium (Diazepam) Tabl et	SS		ORAL
ORAL	7 DAY	Hyperhidrosis Hypokalaemia		Neurontin (Gabapentin) Tablet	SS		ORAL
ORAL	7 DAY	Hypotension Insomnia		Tegretal (Carbamazepi ne) Tablet	SS		ORAL
ORAL	7 DAY	Liver Function Test Abnormal Mood Swings		Celexa Tablet (Citalopram Hydrobromide) Tablet	SS		ORAL

ORAL	7	DAY	Multi-Organ Failure Multiple Drug Overdose	Sonata (Zaleplon) Tablet	SS	ORAL
ORAL	7	DAY	Nightmare Oedema Peripheral	Trazodone (Trazodone) Tablet	SS	ORAL
7	DAY		Pain Paranoia Pneumonia	Lorcet (Paracetamol, Hydrocodone Bitartrate)	SS	
			Pneumonia Aspiration Pneumonia Klebsiella Prothrombin Level Psychomotor Retardation Renal Failure Respiratory Failure Rhabdomyolysis Sepsis Speech Disorder Tendonitis Tremor Vision Blurred	Promethazine (Promethazine) Tablet Clindamycin (Clindamycin) Tablet Prednisone (Prednisone) Tablet Premarin Tablet (Estrogens Conjugated) Tablet Xenical (Orlistat) Flurazepam (Flurazepam) Tablet Prevacid (Lansoprazole) Tablet	C C C C C C C	

Freedom Of Information (FOI) Report

Date:09/22/03ISR Number: 4194758-7Report Type:Expedited (15-DaCompany Report #2003017238

Age:25 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 1800 MG (600, Initial or Prolonged TID) Other 160 MG	Activated Partial Thromboplastin Time Prolonged Anaemia	Health Professional	Neurontin (Gabapentin)	PS		
4 MG	Angioneurotic Oedema Anorexia Asthenia Blood Albumin Decreased		Geodon (Ziprasidone) Benzatropine Mesilate (Benzatropine Besilate)	SS		
6 MG (3, BID)	Blood Chloride Increased Blood Fibrinogen Abnormal		Risperidone (Risperidone)	SS		
75 MG (DAILY)	Blood Iron Decreased Blood Urea Decreased Cough Drug Toxicity		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS		
	Drug Withdrawal Syndrome Dyspnoea Hallucination Hyperhidrosis Hyperplasia Hyperpyrexia Insomnia Lymphadenopathy Lymphoma Muscular Weakness Nausea Paranoia Psychomotor Hyperactivity Rash Red Cell Distribution Width Increased Serum Ferritin Decreased Tachycardia Thrombocytopenia Vomiting Weight Decreased White Blood Cell Count		Clonazepam (Clonazepam) Haloperidol (Haloperidol) Lamotrigine (Lamotrigine) Lansoprazole (Lansoprazole) Ranitidine Hydrochloride (Ranitidine Hy Drochloride) Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride) Trihexyphenidyl Hydrochloride (Trihexyphenidyl Hydrochloride) Chlorpromazine Hydrochloride	C C C C C C C C C		

Decreased

(Chlorpromazine	
Hydrochloride)	C
Olanzapine	
(Olanzapine)	C
Lorazepam	
(Lorazepam)	C
Temazepam	
(Temazepam)	C
Zolpidem Tartrate	
(Zolpidem Tartrate)	C

Date:09/22/03ISR Number: 4194795-2Report Type:Expedited (15-DaCompany Report #2003034669

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiac Disorder	Consumer	Neurontin (Tablets)			
Other		Diabetes Mellitus		(Gabapentin)	PS		ORAL
1800 MG		Non-Insulin-Dependent					
(Q4H), ORAL		Erectile Dysfunction					
		Lung Disorder					
		Weight Increased					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/22/03ISR Number: 4194796-4Report Type:Expedited (15-DaCompany Report #2003036352

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Disorder Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Depression Diarrhoea Hyperreflexia Hypertension Inner Ear Disorder Lung Disorder Multiple Sclerosis Muscle Spasms Nausea Nervous System Disorder Oesophageal Spasm Pain Panic Attack Periodontitis Photophobia Weight Decreased		Prozac (Fluoxetine Hydrochloride) Baclofen (Baclofen)	C C		

Date:09/22/03ISR Number: 4194798-8Report Type:Expedited (15-DaCompany Report #2003038069

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Drug Ineffective Haemorrhage	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							
Other		Hypertension Pain Weight Fluctuation		Clopidogrel Sulfate (Clopidogrel Sulfate) Glyceryl Trinitrate (Glyceryl Trinitrate) Risedronate Sodium (Risedronate Sodium) Isosorbide Mononitrate (Isosorbide Monomitrate)	SS C C C		

Calcium Carbonate
(Calcium Carbonate) C
Atenolol (Atenolol) C
Lorazepam
(Lorazepam) C
Simvastatin
(Simvastatin) C
Zolpidem Tartrate
(Zolpidem Tartrate) C

Date:09/22/03ISR Number: 4194801-5Report Type:Expedited (15-DaCompany Report #2003018755

Age: Gender:Female I/FU:F

Outcome PT
Disability Arthralgia
Other Brain Damage
Cerebral Artery Occlusion
Cerebrovascular Accident
Convulsion

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Emotional Distress Eye Pain Migraine	Consumer	Neurontin (Gabapentin)	PS		
2400 MG (600, FOUR TIMES A DAY)		Oral Pain Peroneal Nerve Palsy		Loestrin (Anovlar) (Norethindrone Acetate, Ethinyl Estradiol)	SS		

Date:09/22/03ISR Number: 4195266-XReport Type:Expedited (15-DaCompany Report #2003037308
Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNKNOWN Other (UNKNOWN), ORAL		Dialysis Parkinson'S Disease	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL

Date:09/22/03ISR Number: 4195476-1Report Type:Expedited (15-DaCompany Report #2003037862
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1200 MG (TID), ORAL		Hepatic Neoplasm Malignant Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Clotrimazole (Clotrimazole) Spironolactone (Spironolactone)	C C		

Cefalexin Monohydrate (Cefalexin Monohydrate)	C
Levothyroxine Sodium (Levothyroxine Sodium)	C
Omeprazole (Omeprazole)	C
Metronidazole (Metronidazole)	C
Anti-Diabetics	C
Codeine (Codeine)	C
Promethazine Hydrochloride (Promethazine Hydrochloride)	C
Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride)	C
Dicycloverine Hydrochloride (Dicycloverine Hydrochloride)	C
Temazepam (Temezepam)	C

Freedom Of Information (FOI) Report

Oxycodone
 Hydrochloride
 (Oxycodone
 Hydrochloride) C
 Glimepiride
 (Glimepiride) C
 Furosemide
 (Furosemide) C
 Nitrofurantoin
 (Nitrofurantoin) C

Date:09/22/03ISR Number: 4195815-1Report Type:Expedited (15-DaCompany Report #2003038770

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alcohol Use	Foreign	Neurontin			
		Metastases To Bone	Health	(Gabapentin)	PS		ORAL
1200 MG							
(TID), ORAL		Psychotic Disorder	Professional				
				Ethanol (Ethanol)	SS		
				Opioids	C		

Date:09/22/03ISR Number: 4195838-2Report Type:Expedited (15-DaCompany Report #2003037701

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
Initial or Prolonged		Dermatitis Exfoliative	Foreign	Neurontin			
300 MG (TID)		Skin Inflammation	Health	(Gabapentin)	PS		
Other							
		Toxic Epidermal	Professional	Amitriptyline			
		Necrolysis	Company	(Amitriptyline)	C		
			Representative	Analgesics	C		

Date:09/22/03ISR Number: 4196331-3Report Type:Expedited (15-DaCompany Report #2003038198

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Dysphagia Foreign Neurontin
Initial or Prolonged Face Oedema Health (Gabapentin) PS
900 MG (TID)

Professional
Company
Representative

Date:09/22/03ISR Number: 4196363-5Report Type:Expedited (15-DaCompany Report #2003038300

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Congenital Anomaly	Congenital Anomaly Maternal Drugs Affecting	Foreign Health	Neurontin (Gabapentin)	PS		
TRANSPLACENTAL 2400 MG (DAILY), PLACENTAL	Foetus Toe Deformity	Professional	Folic Acid (Folic Acid)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/23/03ISR Number: 4196759-1Report Type:Expedited (15-DaCompany Report #2003036042

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Foreign	Neurontin			
Required		Dyskinesia	Health	(Gabapentin)	PS		ORAL
ORAL							
Intervention to		Myoclonus	Professional				
Prevent Permanent							
Impairment/Damage							

Date:09/23/03ISR Number: 4197090-0Report Type:Expedited (15-DaCompany Report #2003038071

Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Alkaline	Foreign	Gabapentin			
300 MG		Phosphatase Abnormal	Literature	(Gabapentin)	PS		
(DAILY),		Hepatic Function Abnormal	Health				
		Liver Function Test	Professional				
		Abnormal					

Date:09/23/03ISR Number: 4197239-XReport Type:Expedited (15-DaCompany Report #2003038560

Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Ammonia Increased	Health	Neurontin			
Initial or Prolonged			Professional	(Gabapentin)	PS		ORAL
200 MG (BID),							
ORAL				Warfarin (Warfarin)	C		
				Oxycocet			
				(Paracetamol,			
				Oxycodone			
				Hydrochloride)	C		
				Lomotil (Atropine			
				Sulfate,			
				Diphenoxylate			

Hydrochloride) C
 All Other
 Therapeutic Products C
 Zinc (Zinc) C
 Ascorbic Acid
 (Ascorbic Acid C

Date:09/23/03ISR Number: 4198135-4Report Type:Expedited (15-DaCompany Report #2003017741

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Drunk	Foreign	Neurontin (Tablets)			
		Somnolence	Health	(Gabapentin)	PS		
400 MG (TID)		Vertigo	Professional	Clonazepam			
				(Clonazepam)	C		

Date:09/23/03ISR Number: 4198147-0Report Type:Expedited (15-DaCompany Report #2003037071

Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Drug Interaction	Foreign	Neurontin (Tablets)			
Other		Sudden Death	Health	(Gabapentin)	PS		ORAL
600 MG (BID),			Professional				
ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

All Other
 Therapeutic Products SS
 Phenytoin Sodium
 (Phenytoin Sodium) C
 Phenobarbital
 (Phenobarbital) C
 Diazepam (Diazepam) C
 Tetrazepam
 (Tetrazepam) C

Date:09/23/03ISR Number: 4198343-2Report Type:Expedited (15-DaCompany Report #2003038573
 Age:91 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Coma	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
300 MG DAILY		Respiratory Arrest	Professional				
ORAL				Fludrocortisone	C		
				Carbamazepine	C		

Date:09/24/03ISR Number: 4198393-6Report Type:Expedited (15-DaCompany Report #2003038820
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Growth Hormone Increased	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
800 MG (BID),		Hypothalamo-Pituitary Disorders					
ORAL		Neoplasm					

Date:09/24/03ISR Number: 4199418-4Report Type:Expedited (15-DaCompany Report #2003038562
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Hospitalisation Health Professional Neurontin (Gabapentin) PS ORAL
Initial or Prolonged
ORAL
Other

Date:09/25/03ISR Number: 4196919-XReport Type:Direct Company Report #CTU 202626
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Neurontin	PS	Park Davis	

Date:09/25/03ISR Number: 4199594-3Report Type:Expedited (15-DaCompany Report #2003039144
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dry Mouth	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Glossodynia					
300 MG		Tongue Blistering					
(DAILY), ORAL		Tongue Dry		Vicodin (Paracetamol, Hydrocodone Bitartrate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/25/03ISR Number: 4199595-5Report Type:Expedited (15-DaCompany Report #2003031129

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Hyperthyroidism	Health	Neurontin			
Initial or Prolonged	Thyroid Neoplasm	Professional	(Gabapentin)	PS		ORAL
2400 MG						
(TID), ORAL	Thyrototoxic Crisis	Company				
		Representative				

Date:09/25/03ISR Number: 4200234-5Report Type:Expedited (15-DaCompany Report #2003038888

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death	Agitation	Foreign	Gabapentin			
Hospitalization -	Cardio-Respiratory Arrest	Health	(Gabapentin)	PS		ORAL
400 MG						
Initial or Prolonged	Coma	Professional				
(DAILY), ORAL						
	Insomnia		Bezafibrate			
	Medication Error		(Bezafibrate)	C		
			Sil-Norboral			
			(Metformin,			
			Glibenclamide)	C		
			Flurazepam			
			(Flurazepam)	C		

Date:09/26/03ISR Number: 4199466-4Report Type:Expedited (15-DaCompany Report #2003038887

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Cholecystitis Chronic	Health	Neurontin			
Initial or Prolonged	Microlithiasis	Professional	(Gabapentin)	PS		
Other	Pancreatitis		Glatiramer Acetate			
			(Glatiramer Acetate)	SS		
			Indometacin			
			(Indometacin)	SS		
			Oxycodone			
			Hydrochloride			

(Oxycodone
Hydrochloride) C
Fentanyl Citrate
(Fentanyl Citrate) C
Amitriptyline
Hydrochloride
(Amitriptyline
Hydrochloride) C

Date:09/26/03ISR Number: 4199903-5Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20030904444
Age:42 YR Gender: I/FU:I

Outcome PT
Death Alanine Aminotransferase
Increased
Asthenia
Blood Urea Increased
Drug Screen Positive
Lethargy
Mental Impairment
Nervous System Disorder
Overdose
Pco2 Decreased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Po2 Decreased Pyrexia Rhabdomyolysis					
ORAL		Sinus Tachycardia	Literature Health Professional	Ultram Tablets (Tramadol Hydrochloride)	PS		ORAL
			Distributor	Acetaminophen/Hydroc odone Tablets	SS		
				Cyclobenzaprine (Cyclobenzaprine)	SS		
				Pioglitazone (Pioglitazone)	SS		
				Orphenadrine (Orphenadrine)	SS		
				Metaxalone (Metaxalone)	SS		
				Furosemide (Furosemide)	SS		
				Gabapentin (Gabapentin)	SS		

Date:09/26/03ISR Number: 4199956-4Report Type:Expedited (15-DaCompany Report #2003034963
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 900 MG (TID), ORAL		Klebsiella Sepsis Neutropenia	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
			Company Representative	Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
				Doxazosin (Doxazosin)	C		
				Amlodipine (Amlodipine)	C		
				Diethylstilbestrol (Diethylstilbestrol)	C		
				Dexamethasone (Dexamethasone)	C		

Acetylsalicylic Acid
(Acetylsalicylic
Acid) C

Date:09/26/03ISR Number: 4199995-3Report Type:Expedited (15-DaCompany Report #2003039085
Age:58 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 100 MG (DAILY), ORAL	Confusional State Sedation	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL

Baclofen (Baclofen) C
Citalopram
(Citalopram) C
Omeprazole
(Omeprazole) C
Metoclopramide
(Metoclopramide) C
Paracetamol

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Freedom Of Information (FOI) Report

(Paracetamol) C
 Morphine Sulfate
 (Morphine Sulfate) C

Date:09/26/03ISR Number: 4200035-8Report Type:Expedited (15-DaCompany Report #2003025783
 Age:68 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 900 MG (TID), Initial or Prolonged ORAL Other	Bundle Branch Block Left Pulmonary Arterial Pressure Increased Supraventricular Extrasystoles Torsade De Pointes Tricuspid Valve Incompetence Ventricular Dyskinesia Ventricular Extrasystoles Ventricular Tachycardia	Foreign Health Professional	Neurontin (Gabapentin) Enalapril Maleate (Enalapril Maleate) Atorvastatin Calcium (Atorvastatin Calcium) Capozide (Hydrochlorothiazide , Captopril) Hydrochlorthiazide (Hydrochlorothiazide)	PS C C C		ORAL

Date:09/26/03ISR Number: 4200037-1Report Type:Expedited (15-DaCompany Report #2003039084
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG Other (DAILY), ORAL 40 MG (PRN), ORAL 400 MG	Alanine Aminotransferase Increased Aspartate Aminotransferase Increased Serum Ferritin Increased	Foreign Health Professional	Neurontin (Gabapentin) Relpax (Eletriptan) Valproate Sodium (Valproate Sodium)	PS SS SS		ORAL ORAL ORAL

(DAILY), ORAL

Date:09/29/03ISR Number: 4201649-1Report Type:Expedited (15-DaCompany Report #S03-FRA-03138-01
Age:55 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 20 MG QD PO	Cyanosis Dermatitis Bullous	Foreign Health	Seropram (Citalopram)	PS		ORAL
80 MG QD PO	Lymphoedema Pruritus	Professional Other	Skenan(Morphine Sulfate)	SS		ORAL
60 MG QD PO	Rash Erythematous Rash Papular		Skenan (Morphine Sulfate)	SS		ORAL
300 MG QD PO			Neurontin (Gabapentin)	SS		ORAL
			Lactulose	SS		
			Duphalac (Lactulose)	C		
			Climaston	C		
			Atarax (Hydroxyzine Hydrochloride)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/29/03ISR Number: 4202755-8Report Type:Expedited (15-DaCompany Report #2003033244

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), Other ORAL		Fall Loss Of Consciousness Memory Impairment Nervous System Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:09/29/03ISR Number: 4202817-5Report Type:Expedited (15-DaCompany Report #2003039135

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100 MG (DAILY), ORAL (TID), ORAL		Abdominal Pain Upper Coma Cough Depression Diarrhoea Flushing Grand Mal Convulsion Hyperhidrosis Hypoaesthesia Menopausal Symptoms Retching Tremor	Consumer	Zoloft (Sertraline) Neurontin (Gabapentin)	PS SS		ORAL ORAL

Date:09/29/03ISR Number: 4202824-2Report Type:Expedited (15-DaCompany Report #KII-2001-0001303

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional Company Representative	Morphine Sulfate (Similar To Nda 19-516) (Morphine Sulfate) Other Neurontin(Gabapentin	PS		

)

SS

Date:09/29/03ISR Number: 4206523-2Report Type:Expedited (15-DaCompany Report #US-SHR-03-012130
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Dizziness Liver Function Test Abnormal	Consumer	Betaseron (Interferon Beta - 1b) Injection	PS		
SUBCUTANEOUS EVERY 2D, SUBCUTANEOUS	5.6 MIU,	Pain Tremor					
				Prednisone (Prednisone) Neurontin (Gabapentin)	SS SS		
300 MG, 4X/DAY ; 300 MG, 3X/DAY; 300 MG, 1X/DAY,				Prempro (Estrogens Conjugated) Flomax (Morniflumate)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/30/03ISR Number: 4203666-4Report Type:Expedited (15-DaCompany Report #064-0945-M0100008

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abortion Induced	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Congenital Anomaly		Complications Of Maternal	Professional Company Representative	Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
900 MG, ORAL		Exposure To Therapeutic Drugs Exomphalos Maternal Drugs Affecting Foetus Multiple Congenital Abnormalities Pregnancy					

Date:09/30/03ISR Number: 4203750-5Report Type:Expedited (15-DaCompany Report #2003039442

Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Renal Failure	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG							
(DAILY), ORAL							

Date:10/01/03ISR Number: 4202133-1Report Type:Direct Company Report #CTU 202963

Age:13 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Aggression		Neurontin	PS		ORAL
300MG PO X 1							
Intervention to Prevent Permanent Impairment/Damage		Confusional State Hostility					

Date:10/01/03ISR Number: 4203511-7Report Type:Expedited (15-DaCompany Report #2003039650

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (TID), ORAL		Drug Withdrawal Syndrome Flushing	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other		Insomnia					

Date:10/02/03ISR Number: 4201449-2Report Type:Direct Company Report #CTU 203053
 Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ESCALATING		Gun Shot Wound Intentional Self-Injury		Gabapentin 300mg	PS		
DOSE FROM 1-9 PILLS THEN DOSE TAPER TO CROSSOVER				Placebo	SS		

Date:10/02/03ISR Number: 4201597-7Report Type:Direct Company Report #CTU 203063
 Age:82 YR Gender:Male I/FU:I

Outcome	PT
Disability	Deafness Micturition Urgency

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1CAP 3 X			Neurontin 100 Mg Parke-Davis	PS	Parke-Davis	
DAILY - 2 WKS						
THEN 2 CAPS 3						
X DAILY						

Date:10/02/03ISR Number: 4201697-1Report Type:Direct Company Report #CTU 203074
 Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG PO BID Initial or Prolonged		Hyponatraemia		Neurontin	PS		ORAL

Date:10/02/03ISR Number: 4204806-3Report Type:Expedited (15-DaCompany Report #2003033889
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (TID), Other ORAL		Abnormal Behaviour Anxiety Cerebrovascular Accident	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Gangrene		Isosorbide (Isosorbide)	C		
		Irritability		Lasix (Furosemide)	C		
		Leg Amputation		Norvasc (Amlodipine)	C		
		Memory Impairment		Dilantin (Phenytoin Sodium)	C		
		Palpitations		All Other			
		Skin Ulcer		Therapeutic Products	C		
		Tachycardia		Valium (Diazepam)	C		
		Tinnitus		Lisinopril			

(Lisinopril)

C

Date:10/02/03ISR Number: 4204807-5Report Type:Expedited (15-DaCompany Report #2003031759

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia	Health	Neurontin			
300 MG		Carpal Tunnel Syndrome	Professional	(Gabapentin)	PS		ORAL
(DAILY), ORAL		Colitis Ulcerative					
		Disturbance In Attention		Ibuprofen			
2400 MG		Dizziness		(Ibuprofen)	SS		ORAL
(DAILY), ORAL		Drug Ineffective					
		Drug Interaction		Valproate Semisodium			
		Feeling Abnormal		(Valproate			
		Feeling Drunk		Semisodium)	SS		
		Intervertebral Disc		Acetylsalicylic Acid			
		Protrusion		(Acetylsalicylic			
		Medication Error		Acid)	SS		
		Neuralgia		Mesalazine			
		Pain		(Mesalazine)	SS		
		Pain In Extremity		Glycopyrronium			
		Road Traffic Accident		Bromide			
				(Glycopyrronium			
				Bromide)	C		
				Propranolol			

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Freedom Of Information (FOI) Report

Hydrochloride
 (Propranolol
 Hydrochloride) C
 Amitriptyline
 Hydrochloride
 (Amitriptyline
 Hydrochloride) C

Date:10/03/03ISR Number: 4204589-7Report Type:Expedited (15-DaCompany Report #2003039721
 Age:34 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Hydroxyzine Hydrochloride (Tablet) (Hydroxyzine Hydrochloride)	SS		ORAL
ORAL				Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		ORAL

Date:10/03/03ISR Number: 4204593-9Report Type:Expedited (15-DaCompany Report #2003039898
 Age:52 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose	Literature Health Professional	Doxepin (Caps) (Doxepin)	PS		
				Gabapentin (Gabapentin)	SS		
				Sertraline (Sertraline)	SS		
				All Other Therapeutic Products	SS		

Date:10/03/03ISR Number: 4204603-9Report Type:Expedited (15-DaCompany Report #2003039897
Age:27 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Completed Suicide	Literature Health Professional	Doxepin (Doxepin) Gabapentin (Gabapentin) Olanzapine (Olanzapine)	PS SS SS		

Date:10/03/03ISR Number: 4204730-6Report Type:Expedited (15-DaCompany Report #2003039719
Age:54 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose	Literature Health Professional	Gabapentin (Gabapentin) Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	PS SS		ORAL ORAL

ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL				Vicodin (Paracetamol, Hydrocodone Bitartrate)	SS		ORAL
				All Other Therapeutic Products	SS		

Date:10/03/03ISR Number: 4204737-9Report Type:Expedited (15-DaCompany Report #2003040285
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Chest Pain Conduction Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG (800 MG TID), ORAL		Electrocardiogram Abnormal					

Date:10/03/03ISR Number: 4205961-1Report Type:Expedited (15-DaCompany Report #2003037071
Age:15 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Drug Interaction Sudden Death	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
600 MG BID, ORAL			Professional				

Phenytoin Sodium (Phenytoin Sodium)	C
Phenobarbital (Phenobarbital)	C
Diazepam (Diazepam)	C
Tetrazepam (Tetrazepam)	C

Date:10/03/03ISR Number: 4206161-1Report Type:Expedited (15-DaCompany Report #2003040746
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Blood Alcohol Increased Face Injury	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Intentional Misuse Laceration Loss Of Consciousness	Professional Company Representative	Ethanol (Ethanol)	C		

Date:10/03/03ISR Number: 4206348-8Report Type:Expedited (15-DaCompany Report #2003039720
Age:50 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Cardio-Respiratory Arrest Completed Suicide Overdose	Literature Health Professional	Gabapentin (Gabapentin) Methadone (Methadone) Olanzapine (Olanzapine)	PS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/06/03ISR Number: 4201501-1Report Type:Expedited (15-DaCompany Report #DE-BRISTOL-MYERS SQUIBB COMPANY-12331336
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abdominal Pain		Taxol Inj Syn	PS	Bristol-Myers Squibb Company	
Life-Threatening		Back Pain					
INTRAVENOUS							
Hospitalization -		Confusional State		Ribocarbo	SS		
INTRAVENOUS	AUC 2						
Initial or Prolonged		Diaphragmatic Paralysis		Gabapentin	SS		
		Dyspnoea					
		General Physical Health					
		Deterioration					
		Malnutrition					
		Memory Impairment					
		Motor Dysfunction					
		Neoplasm Progression					
		Overdose					
		Pneumonia					
		Visual Disturbance					

Date:10/06/03ISR Number: 4204955-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000697
 Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Acrochordon
Disability	Actinic Keratosis
Other	Alanine Aminotransferase
	Increased
	Aspartate
	Aminotransferase
	Increased
	Blood Urea
	Nitrogen/Creatinine Ratio
	Increased
	Bone Disorder
	Brain Neoplasm
	Chondromalacia
	Cognitive Disorder
	Conversion Disorder
	Convulsion
	Cyst
	Depressed Level Of

Consciousness
Dizziness
Dysphemia
Ependymoma
Fatigue
Fibromyalgia
Folliculitis
Gamma-Glutamyltransferase
Increased
Hypersomnia
Hypoglycaemia
Memory Impairment
Meniscus Lesion
Nervous System Disorder
Osteopenia
Pain
Pigmented Naevus
Seborrhoeic Keratosis
Skin Irritation

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Somnolence Speech Disorder Tremor			Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Vision Blurred			Professional Company Representative	Valproate Sodium (Valproate Sodium)	SS		
				Buspirone Hydrochloride (Buspirone Hydrochloride)	C		
				Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		
				Omeprazole (Omeprazole)	C		
				Capsaicin (Capsaicin)	C		

Date:10/06/03ISR Number: 4205092-0Report Type:Expedited (15-DaCompany Report #2003040741
Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other			Consumer	Neurontin (Gabapentin)	PS		ORAL
Dry Mouth							
Fatigue Impaired Driving Ability							

Date:10/06/03ISR Number: 4205135-4Report Type:Expedited (15-DaCompany Report #2003034005
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Consumer	Neurontin			

Initial or Prolonged	Disease			
ORAL			(Gabapentin)	PS
Other	Cardiac Failure		Risedronate Sodium	
	Congestive		(Risedronate Sodium)	C
	Dehydration		Esomeprazole	
	Fatigue		(Esomeprazole)	C
	Feeling Abnormal		Levothyroxine Sodium	
	Pain		(Levothyroxine	
	Urinary Incontinence		Sodium)	C
			Clopidogrel Sulfate	
			(Clopidogrel	
			Sulfate)	C
			Metoprolol Succinate	
			(Metoprolol	
			Succinate)	C
			Vicodin	
			(Paracetamol,	
			Hydrocodone	
			Bitartrate)	C
			Amlodipine Besilate	
			(Amlodipine	
			Besilate)	C
			Simvastatin	
			(Simvastatin)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Meclozine
(Meclozine) C
Alprazolam
(Alprazolam) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C

Date:10/06/03ISR Number: 4205142-1Report Type:Expedited (15-DaCompany Report #2002058034
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hypoaesthesia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other	1200 MG						
(TID), ORAL							

Fentanyl (Fentanyl) C
Amitriptyline
(Amitriptyline) C

Date:10/06/03ISR Number: 4205149-4Report Type:Expedited (15-DaCompany Report #2003040742
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1800 MG	Balance Disorder Blindness	Consumer	Neurontin (Gabapentin)	PS		ORAL
Sleep Attacks							
(600,TID),							

ORAL

Date:10/06/03ISR Number: 4205196-2Report Type:Expedited (15-DaCompany Report #03P-083-0234836-00
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Hepatic Enzyme Increased	Foreign	Depakine (Sodium			

Initial or Prolonged	Serum Ferritin Increased	Health	Valproate)	PS	ORAL
200 MG 2 IN 1		Professional			
D, ORAL			Gabapentin	SS	ORAL
900 MG, 1 IN					
1 D, ORAL			Eletriptan	C	

Date:10/07/03ISR Number: 4206989-8Report Type:Expedited (15-DaCompany Report #2003040984
 Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Respiratory Arrest	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG							
(DAILY), ORAL				All Other Therapeutic Products	C		

Date:10/07/03ISR Number: 4209548-6Report Type:Periodic Company Report #2002133111US
 Age:19 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Literature Health	Xanax (Alprazolam) Tablet	PS		ORAL
ORAL			Professional	Meprobamate			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL (Meprobamate) SS ORAL

ORAL Gabapentin (Gabapentin) SS ORAL

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
Dose		Caesarean Section	Consumer	Neurontin			
Other		Complications Of Maternal	Health	(Gabapentin)	PS		ORAL
600 MG (300,		Exposure To Therapeutic	Professional				
BID), ORAL		Drugs		Bupropion			
		Maternal Drugs Affecting		Hydrochloride			
		Foetus		(Bupropion			
300 MG (150,		Pregnancy Test Positive		Hydrochloride)	SS		ORAL
BID), ORAL				Rizatriptan Benzoate			
				(Rizatriptan			
				Benzoate)	SS		

Date:10/08/03ISR Number: 4207679-8Report Type:Expedited (15-DaCompany Report #2003030062
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amputation	Consumer	Lipitor			
Other		Surgery	Health	(Atorvastatin)	PS		
			Professional	Neurontin(Gabapentin			
) (Gabapentin)	SS		
				Cilostazol			
				(Cilostazol)	C		

Date:10/08/03ISR Number: 4207680-4Report Type:Expedited (15-DaCompany Report #2003040766
Age:5 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Autism	Health	Neurontin			
		Maternal Drugs Affecting	Professional	(Gabapentin)	PS		
		Foetus					

Date:10/08/03ISR Number: 4207681-6Report Type:Expedited (15-DaCompany Report #2002072914
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin			
				(Gabapentin)	PS		

Date:10/09/03ISR Number: 4208177-8Report Type:Expedited (15-DaCompany Report #2003041153
Age:77 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Ankle Fracture
Initial or Prolonged	Cerebellar Atrophy
Other	Cerebrovascular Accident
	Chest Pain
	Confusional State
	Difficulty In Walking
	Disorientation

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Disturbance In Attention Drug Interaction Encephalopathy	Report Source	Product	Role	Manufacturer	Route
900 MG		Fall Fatigue	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL		Hallucination, Visual	Professional				
TRANSDERMAL	20 MG	Herpes Zoster Myoclonus		Buprenorphine (Buprenorphine)	SS		
(DAILY),		Pain					
TRANSDERMAL		Speech Disorder					
200 MG				Amiodarone Hydrochloride (Amiodarone Hydrochloride)	SS		ORAL
(DAILY),							
ORAL, YEARS							
200 MG				Tramadol Hydrochloride (Tramadol Hydrochloride)	SS		ORAL
(50,QID),							
ORAL							
				Omeprazole (Omeprazole)	C		
				Pravastatin Sodium (Pravastatin Sodium)	C		
				Furosemide (Furosemide)	C		
				Glyceryl Trinitrate (Glyceryl Trinitrate)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Acenocoumarol			

Date:10/09/03ISR Number: 4208307-8Report Type:Expedited (15-DaCompany Report #2003038770
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcohol Use	Foreign	Neurontin			
Other		Metastases To Bone	Health	(Gabapentin)	PS		ORAL
1200 MG		Psychotic Disorder	Professional				
(TID), ORAL			Company Representative	Ethanol (Ethanol) Opioids	SS C		

Date:10/09/03ISR Number: 4208453-9Report Type:Expedited (15-DaCompany Report #2003041138
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Renal Impairment	Health	Neurontin			
Other			Professional	(Gabapentin)	PS		
1800 (TID)			Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/09/03ISR Number: 4208928-2Report Type:Direct
Age:38 YR Gender:Male I/FU:I

Company Report #CTU 203553

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dehydration		Lithium	PS		
		Large Intestinal Obstruction		Quetiapine	SS		
		Shock		Gabapentin	SS		
		Vomiting		Clonazepam	SS		

Date:10/10/03ISR Number: 4206164-7Report Type:Expedited (15-DaCompany Report #FR-BRISTOL-MYERS SQUIBB COMPANY-12402749
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Re-started		Amyotrophy Asthenia		Sustiva	PS	Bristol-Myers Squibb Company	
Other 08-Jun-2001		Calculus Bladder					
		Dehydration		Zerit Caps	SS	Bristol-Myers Squibb Company	ORAL
		Gynaecomastia					
		Hepatic Steatosis		Videx	SS	Bristol-Myers Squibb Company	
		Hepatitis					
		Hepatomegaly		Indinavir	SS		
		Hyperlactacidaemia		Bactrim	SS		
		Hypersensitivity		Plaquenil	SS		
		Hypoaesthesia		Interferon	SS		
units		Lactic Acidosis		Ribavirin	SS		
		Lipoatrophy		L-Carnitine	SS		
		Liver Function Test Abnormal		Vitamin B2	SS		
		Muscle Spasms		Vitamin B1	SS		
		Nervous System Disorder		Zidovudine	SS		
		Pancreatitis		Lamivudine	SS		
		Paraesthesia		Neurontin	SS		
		Phlebothrombosis		Vitamin E	SS		
		Pyrexia		Selenium	SS		
		Tinnitus		Kaletra	SS		
		Vertigo					
		Vomiting					
		Weight Decreased					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bladder Spasm	Health	Neurontin			
Other		Complex Regional Pain	Professional	(Gabapentin)	PS		ORAL
300 MG (TID),		Syndrome					
ORAL		Condition Aggravated		Zonisamide			
		Feeling Abnormal		(Zonisamide)	SS		
		Laboratory Test Abnormal		Ropinirole			
		Medication Error		Hydrochloride			
		Vaginitis Bacterial		(Ropinirole			
		White Blood Cell Count		Hydrochloride)	SS		
		Decreased		Tizanidine			
				Hydrochloride			
				(Tizanidine			
				Hydrochloride)	SS		
				Montelukast Sodium			
				(Montelukast Sodium)	C		
				All Other			

Freedom Of Information (FOI) Report

Therapeutic Products C
 Multivitamins
 (Ergocalciferol,
 Ascorbic Acid, Folic
 Acid, Thiamine
 Hydrochloride, C
 Magnesium
 (Magnesium) C
 Calcium (Calcium) C
 Ascorbic Acid
 (Ascorbic Acid) C
 Zinc (Zinc) C
 Chromium (Chromium) C
 Policosanol
 (Policosanol) C
 Lactobacillus
 Acidophilus
 (Lactobacillus
 Acidophilus) C

Date:10/10/03ISR Number: 4208805-7Report Type:Expedited (15-DaCompany Report #2003041397
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other PRN, ORAL		Depression Petit Mal Epilepsy	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Suicidal Ideation Tremor		Dilantin (Phenytoin Sodium)	SS		
				Retinol (Retinol)	C		
				Folic Acid (Folic Acid)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Testosterone Cipionate (Testosterone Cipionate)	C		
				Oxycocet (Paracetamol, Oxycodone Hydrochloride)	C		
				Oxycodone Hydrochloride			

(Oxycodone Hydrochloride)	C
Tocopherol (Tocopherol)	C
Furosemide (Furosemide)	C
Potassium (Potassium)	C
Hydroxyzine Embonate (Hydroxyzine Embonate)	C
Flurazepam Hydrochloride (Flurazepam Hydrochloride)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/10/03ISR Number: 4208806-9Report Type:Expedited (15-DaCompany Report #2003037139
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract	Health	Neurontin			
		Convulsion	Professional	(Gabapentin)	PS		
		Deafness Unilateral		Antivert (Meclizine)	SS		
		Electroencephalogram		Celecoxib			
		Abnormal		(Celecoxib)	SS		
400 MG (BID)		Headache		Vicodin			
		Hyporeflexia		(Paracetamol,			
		Insomnia		Hydrocodone			
		Medication Error		Bitartrate)	C		
		Syncope					
		Treatment Noncompliance					

Date:10/10/03ISR Number: 4208808-2Report Type:Expedited (15-DaCompany Report #2003041305
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Acute Respiratory Failure	Health	Neurontin			
Initial or Prolonged		Pneumonia Aspiration	Professional	(Gabapentin)	PS		ORAL
4800 MG							
(QID), ORAL				Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	C		

Date:10/10/03ISR Number: 4208811-2Report Type:Expedited (15-DaCompany Report #2003034283
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Abdominal Adhesions	Consumer	Neurontin			
Initial or Prolonged		Colitis Ulcerative	Health	(Gabapentin)	PS		ORAL
3000 MG							
Other		Pain	Professional				
(TID), ORAL							

Quetiapine Fumarate
 (Quetiapine Fumarate) C
 Clonazepam
 (Clonazepam) C

Date:10/10/03ISR Number: 4208863-XReport Type:Expedited (15-DaCompany Report #2003028077
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged SEE IMAGE	Anxiety Asthenia	Consumer Health	Neurontin (Gabapentin)	PS		
Disability Other	Circulatory Collapse Dysgeusia Emotional Distress Lethargy Malaise Nausea Pain Pneumonia	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/10/03ISR Number: 4209153-1Report Type:Expedited (15-DaCompany Report #2003041585
Age:92 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Respiratory Arrest	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG							
(DAILY), ORAL							

Date:10/10/03ISR Number: 4209235-4Report Type:Expedited (15-DaCompany Report #2003110066
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Activities Of Daily Living Impaired	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
1800 MG		Asthenia					
(DAILY), ORAL							
		Balance Disorder		Morphine (Morphine)	C		
		Bedridden		Pinaverium Bromide			
		Bradyphrenia		(Pinaverium Bromide)	C		
		Confusional State		Dimenhydrinate			
		Decreased Activity		(Dimenhydrinate)	C		
		Diplopia		Desipramine			
		Disturbance In Attention		(Desipramine)	C		
		Dizziness		Estradiol			
		Dysarthria		(Estradiol)	C		
		Dysphemia		Ranitidine			
		Emotional Disorder		Hydrochloride			
		Fatigue		(Ranitidine			
		Gait Disturbance		Hydrochloride)	C		
		Headache					
		Hearing Impaired					
		Impaired Driving Ability					
		Impaired Work Ability					
		Insomnia					
		Lack Of Spontaneous Speech					
		Learning Disorder					
		Muscle Spasms					
		Myalgia					
		Nasopharyngitis					

Nervous System Disorder
Pain
Speech Disorder
Vision Blurred

Date:10/14/03ISR Number: 4207923-7Report Type:Expedited (15-DaCompany Report #FR-ROCHE-348810
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	140 DAY	Agitation		Valium	PS	Roche	ORAL
	56 DAY	Fall		Neurontin	SS		ORAL
	56 DAY	Sudden Death		Deroxat	SS		ORAL
	18 DAY			Noctamide	SS		ORAL
	120 DAY			Gardenal	SS		ORAL
INTRAMUSCULAR			1 DAY	Tercian	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4209463-8Report Type:Expedited (15-DaCompany Report #2003041236

Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anorgasmia	Consumer	Viagra (Sildenafil)	PS		ORAL
50 MG (PRN), Other ORAL		Body Height Decreased					
		Drug Effect Decreased		Gabapentin			
		Ejaculation Delayed		(Gabapentin)	SS		
TOPICAL	TOPICAL	Exostosis		Strattera			
		Intervertebral Disc Protrusion		(Atomoxetine Hydrochloride)	SS		
		Libido Decreased		Rofecoxib			
		Mental Disorder		(Rofecoxib)	C		
		Pain In Extremity		Esomeprazole			
		Penis Disorder		(Esomeprazole)	C		
		Sciatic Nerve Injury					
		Spinal Cord Injury					
		Spinal Deformity					
		Spinal Disorder					
		Spondylolisthesis					
		Acquired					

Date:10/14/03ISR Number: 4209629-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Acrochordon
Disability	Actinic Keratosis
Other	Acute Sinusitis
	Alanine Aminotransferase
	Increased
	Alcoholism
	Anaphylactic Shock
	Aphasia
	Aspartate
	Aminotransferase
	Increased
	Blood Urea
	Nitrogen/Creatinine Ratio
	Increased

Brain Neoplasm Benign
Cerebellar Atrophy
Cerebral Astrocytoma, Low
Grade
Chondropathy
Cognitive Disorder
Condyloma Acuminatum
Cyst
Depressed Level Of
Consciousness
Depression
Disturbance In Attention
Drug Hypersensitivity
Dysphemia
Ependymoma
Fatigue
Fibromyalgia
Folliculitis
Gamma-Glutamyltransferase
Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (BID), ORAL		Hallucination Hypersomnia Hypoglycaemia Impaired Work Ability Memory Impairment Meniscus Lesion Mental Impairment Nervous System Disorder Neurodegenerative Disorder Osteopenia Paraesthesia Paraneoplastic Syndrome Patellofemoral Pain Syndrome Pigmented Naevus Seborrheic Keratosis Sleep Disorder Speech Disorder Suicidal Ideation Tinnitus Vision Blurred	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin)	PS SS C C C C		ORAL

Date:10/14/03ISR Number: 4209799-0Report Type:Expedited (15-DaCompany Report #2003041715
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG, ORAL		Balance Disorder Dizziness Dysstasia Joint Swelling Muscle Spasms Pain	Consumer	Neurontin (Gabapentin) Carbamazepine (Carbamazepine) Sinemet (Levodopa, Carbidopa) Triamterene (Triamterene) Lisinopril (Lisinopril) Pravastatin Sodium (Pravastatin Sodium) Phenobarbital	PS C C C C C		ORAL

(Phenobarbital) C
Baclofen (Baclofen) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C
Hydrochlorothiazide
(Hydrochlorothiazide
) C

Date:10/14/03ISR Number: 4209803-XReport Type:Expedited (15-DaCompany Report #2003018755

Age:37 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Arthralgia
Initial or Prolonged	Brain Damage
Disability	Cerebral Artery Occlusion
Other	Cerebral Infarction
	Cerebrovascular Accident
	Convulsion

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Difficulty In Walking Emotional Distress Eye Pain	Consumer	Neurontin (Gabapentin)	PS		
2400 MG (600 FOUR TIMES A DAY)		Injury Migraine Oral Pain Pain Peroneal Nerve Palsy Self-Medication		Loestrin (Anovlar) (Norethindrone Acetate, Ethinyl Estradiol) Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS SS		

Date:10/14/03ISR Number: 4210188-3Report Type:Expedited (15-DaCompany Report #2003110281
Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiovascular Disorder Face Injury Fall Haematoma Muscle Twitching Pyrexia	Foreign Health Professional	Neurontin (Gabapentin) Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Calcium Acetate (Calcium Acetate) Folic Acid (Folic Acid) Doxepin Hydrochloride (Doxepin Hydrochloride) Insulin Human Injection, Isophane (Insulin Human Injection, Isophane) Esomeprazole	PS C C C C C		

(Esomeprazole)	C
Panadeine Co	
(Codeine Phosphate, Paracetamol)	C
Antiinfectives	C
Piritramide	
(Piritramide)	C
Ferric Sodium Gluconate Complex	
(Ferric Sodium Gluconate Complex)	C
Epoetin Alfa	
(Epoetin Alfa)	C
Carbamazepine	
(Carbamazepine)	C
Diazepam (Diazepam)	C
Vancomycin	
(Vancomycin)	C
Blinded Therapy	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211410-XReport Type:Periodic
 Age: Gender:Male I/FU:F

Company Report #2002052813

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2700 MG (TID), ORAL		Pneumonia	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)	C	
Gatifloxacin(Gatiflo xacin)	C	
Lansoprazole (Lansoprazole)	C	
Metoclopramide (Metoclopramide)	C	
Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C	
Celecoxib (Celecoxib)	C	
Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C	

Date:10/14/03ISR Number: 4211412-3Report Type:Periodic
 Age: Gender:Female I/FU:F

Company Report #2002053813

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 500 MG (QID), ORAL		Tremor	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Oxybutynin (Oxybutynin) Quetiapine Fumarate	C	
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(Quetiapine
 Fumarate) C
 Nefazodone
 Hydrochloride
 (Nefazodone
 Hydrochloride) C
 Trihexyphenidyl
 (Trihexyphenidyl) C

Date:10/14/03ISR Number: 4211415-9Report Type:Periodic Company Report #001-0945-M0100468
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211417-2Report Type:Periodic
Age: Gender: I/FU:F

Company Report #001-0945-M0100469

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211418-4Report Type:Periodic
Age: Gender: I/FU:F

Company Report #001-0945-M0100471

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211421-4Report Type:Periodic
Age: Gender:Male I/FU:F

Company Report #001-0945-M0101129

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	2400 MG (600 MG, QID), ORAL	Hallucination Intentional Misuse Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL

Fosinopril Sodium (Fosinopril Sodium)	C
Simvastatin (Simvastatin)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C
Ranitidine (Ranitidine)	C
Amitriptyline (Amitriptyline)	C
Metoprolol Succinate (Metoprolol Succinate)	C

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
		Difficulty In Walking	Health	(Gabapentin)	PS		ORAL
900 MG (TID),							
ORAL		Fatigue	Professional				
		Headache		Didanosine			
		Increased Appetite		(Didanosine)	C		
		Intentional Misuse		Nevirapine			
		Mood Altered		(Nevirapine)	C		
		Neuropathy Peripheral		Stavudine			
		Oedema Peripheral		(Stavudine)	C		
		Pain		Clonazepam			
		Somnolence		(Clonazepam)	C		
		Speech Disorder		Warfarin Sodium			
		Vision Blurred		(Warfarin Sodium)	C		
		Weight Increased		Metoprolol Succinate			
				(Metoprolol			
				Succinate)	C		
				Trandolapril			
				(Trandolapril)	C		

Freedom Of Information (FOI) Report

Omeprazole
(Omeprazole) C

Date:10/14/03ISR Number: 4211425-1Report Type:Periodic
Age:74 YR Gender:Female I/FU:F

Company Report #2002050045

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 400 MG (200 MG, TWICE DAILY), ORAL	Back Pain Blood Cholesterol Increased Cataract	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
10 MG (DAILY), ORAL	Dizziness Pain In Extremity		Atorvastatin (Atorvastatin)	SS		ORAL
			Latanoprost (Latanoprost)	C		
			Repaglinide (Repaglinide)	C		
			Metformin Hydrochloride (Metformin Hydrochloride)	C		
			Loratadine (Loratadine)	C		
			Pirbuterol Acetate (Pirbuterol Acetate)	C		
			Budesonide (Budesonide)	C		
			Naproxen Sodium (Naproxen Sodium)	C		
			Mometasone Furoate (Mometasone Furoate)	C		
			Felodipine (Felodipine)	C		
			Acetylsalicylic Acid (Acetylsalicylic Acid)	C		

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident	Consumer	Neurontin (Gabapentin)	PS		
300 MG		Fatigue					
		Nervousness		Celecoxib (Celecoxib)	SS		ORAL
100 MG (QD),		Pain					
ORAL		Sensory Disturbance		Valdecoxib (Valdecoxib)	SS		ORAL
10 MG (QD),							
ORAL				Atorvastatin (Atorvastatin)	C		
				Clopidogrel Sulfate (Clopidogrel Sulfate)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211430-5Report Type:Periodic
Age:77 YR Gender:Female I/FU:F

Company Report #2002051328

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), ORAL		Cerebrovascular Accident Feeling Abnormal	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Rabeprazole (Rabeprazole)	C		
				Celecoxib (Celecoxib)	C		
				Clopidogrel (Clopidogrel)	C		
				Potassium Chloride (Potassium Chloride)	C		
				Furosemide (Furosemide)	C		
				Losartan Potassium (Losartan Potassium)	C		
				Alendronate Sodium (Alendronate Sodium)	C		
				Famotidine (Famotidine)	C		
				Benzonatate (Ebnzonatate)	C		

Date:10/14/03ISR Number: 4211433-0Report Type:Periodic
Age: Gender:Male I/FU:F

Company Report #2002052304

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG (600 MG, THREE TIMES DAILY), ORAL		Blood Triglycerides Increased Cerebrovascular Accident Coordination Abnormal Neuropathy Peripheral Osteopenia Vision Blurred	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Irbesartan (Irbesartan)	C		

Date:10/14/03ISR Number: 4211436-6Report Type:Periodic Company Report #2003033157
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Mania	Consumer	Neurontin (Gabapentin)	PS		ORAL
UNKNOWN, ORAL							

Date:10/14/03ISR Number: 4211437-8Report Type:Periodic Company Report #2003033496
 Age:66 YR Gender:Male I/FU:I

Outcome	PT
Other	Agitation Balance Disorder Disorientation Disturbance In Attention Dizziness Dysphagia Dyspnoea Illogical Thinking Insomnia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
		Irritability Mood Altered Rash	Consumer	Neurontin (Gabapentin)	PS		ORAL
UNKNOWN, ORAL		Restlessness Somnolence		Antihypertensives Cholesterol-And Triglyceride Reducers Alprazolam (Alprazolam) Analgesics Atenolol (Atenolol) Captopril (Captopril) Metformin Hydrochloride (Metformin Hydrochloride)	C C C C C C		

Date:10/14/03ISR Number: 4211440-8Report Type:Periodic Company Report #2003034183
Age:51 YR Gender:Female I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
1800 MG (DAILY), ORAL		Fluid Retention Oedema Peripheral Weight Increased	Consumer	Neurontin (Gabapentin)	PS		
				Nortriptyline (Nortriptyline)	C		

Date:10/14/03ISR Number: 4211443-3Report Type:Periodic Company Report #001-0945-M0100029
Age:77 YR Gender:Female I/FU:F

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
300 MG (QHS),		Alopecia Constipation	Consumer Health	Neurontin (Gabapentin)	PS		ORAL

ORAL	Deafness	Professional			
	Depression	Company	Depo-Medrol		
	Diarrhoea	Representative	(Methylprednisolone		
	Dry Mouth		Acetate)	SS	
UNKNOWN	Eye Pruritus		All Other		
	Insomnia		Therapeutic Products	SS	
UNKNOWN	Lacrimation Increased		Doxazosin Mesilate		
	Muscle Spasms		(Doxazosin Mesilate)	C	
	Nervousness		Cortisone		
	Visual Acuity Reduced		(Cortisone)	C	
	Vomiting		Paracetamol		
			(Paracetamol)	C	

Date:10/14/03ISR Number: 4211445-7Report Type:Periodic Company Report #001-0945-M0100462
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin			
Other				(Gabapentin)	PS		
UNKNOWN							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211447-0Report Type:Periodic
 Age: Gender: I/FU:F

Company Report #001-0945-M0100463

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							
UNKNOWN							

Date:10/14/03ISR Number: 4211449-4Report Type:Periodic
 Age: Gender: I/FU:I

Company Report #001-0945-M0100464

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							
UNKNOWN							

Date:10/14/03ISR Number: 4211450-0Report Type:Periodic
 Age: Gender: I/FU:F

Company Report #001-0945-M0100465

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							
UNKNOWN							

Date:10/14/03ISR Number: 4211452-4Report Type:Periodic
 Age: Gender: I/FU:F

Company Report #001-0945-M0100466

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							
UNKNOWN							

Date:10/14/03ISR Number: 4211453-6Report Type:Periodic
 Age: Gender: I/FU:F

Company Report #001-0945-M0100467

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							
UNKNOWN							

Date:10/14/03ISR Number: 4211455-XReport Type:Periodic Company Report #2003027730
 Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abnormal Behaviour Amnesia Blood Glucose Increased Hallucination, Visual Speech Disorder	Consumer	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211456-1Report Type:Periodic Company Report #2003028162
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (TID)		Blood Pressure Decreased Dehydration Somnolence	Health Professional	Neurontin (Gabapentin)	PS		
				Furosemide (Furosemide) Betaseron (Glucose, Albumin Human, Interferon Beta)	SS SS		

SUBCUTANEOUS EVERY 2 DAYS

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

SUBCUTANEOUS

Modafinil
(Modafinil) C

Date:10/14/03ISR Number: 4211458-5Report Type:Periodic Company Report #2003028400
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Memory Impairment Renal Failure Acute	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211459-7Report Type:Periodic Company Report #2003029017
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Thrombocytopenia	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211461-5Report Type:Periodic Company Report #2003030372
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
Other							

Date:10/14/03ISR Number: 4211462-7Report Type:Periodic Company Report #2003031037
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intentional Misuse	Health	Neurontin			

3600 MD	Medication Error	Professional	(Gabapentin)	PS
	Pancreatitis		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS

Date:10/14/03ISR Number: 4211463-9Report Type:Periodic Company Report #2003031373
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Sodium Decreased Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
4500 MG (1500 TID) ORAL		Medication Error Overdose		Oxcarbazepine (Oxcarbazepine)	SS		ORAL
600 MG (300 BID) ORAL				Terazosin (Hydrochloride)	C		
				Loratadine (Loratadine)	C		
				Omeprazole	C		
				Modafinil (Modafinil)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Estrogens Conjugated C
 Rofecoxib C

Date:10/14/03ISR Number: 4211465-2Report Type:Periodic
 Age:32 YR Gender:Female I/FU:I

Company Report #2003031961

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID) Other		Stevens-Johnson Syndrome	Consumer	Gabapentin (Gabapentin)	PS		
				Zidovudine (Zidovudine)	C		
				Pyrimethamine	C		
				Calcium Folate	C		
				Clindamycin	C		

Date:10/14/03ISR Number: 4211466-4Report Type:Periodic
 Age:58 YR Gender:Female I/FU:I

Company Report #2003032347

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (DAILY), ORAL		Alopecia Convulsion Feeling Abnormal Rash Tremor	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:10/14/03ISR Number: 4211467-6Report Type:Periodic
 Age:73 YR Gender:Male I/FU:I

Company Report #2003032864

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG (DAILY) ORAL		Coordination Abnormal Cough Deafness Unilateral Disturbance In Attention	Consumer	Neurontin (Gabapentin)	PS		ORAL

Dizziness
 Dry Mouth
 Eyelid Disorder
 Feeling Abnormal
 Headache
 Lacrimation Increased
 Listless
 Mood Altered
 Pain
 Pharyngolaryngeal Pain
 Pyrexia
 Somnolence
 Vision Blurred

Date:10/14/03ISR Number: 4211470-6Report Type:Periodic Company Report #2003024052
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abasia	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG		Back Pain					
(DAILY), ORAL		Hypokinesia		Rofecoxib (Rofecoxib) Alosetron	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Alosetron) C

Date:10/14/03ISR Number: 4211472-XReport Type:Periodic
 Age:74 YR Gender:Male I/FU:I

Company Report #2003024059

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (DAILY), ORAL		Pain Pneumonia Somnolence	Consumer	Neurontin (Gabapentin)	PS		ORAL

Atorvastatin
(Atorvastatin) C
 Quinapril
Hydrochloride
(Quinapril
Hydrochloride) C
 Isosorbide
(Isosorbide) C
 Metformin
Hydrochloride
(Metformin
Hydrochloride) C
 Tamsulosin
Hydrochloride
(Tamsulosin
Hydrochloride) C
 Ibuprofen
(Ibuprofen) C

Date:10/14/03ISR Number: 4211473-1Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #2003024120

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300, Q8H)		Thrombocytopenia	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211476-7Report Type:Periodic Company Report #2003024486
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211477-9Report Type:Periodic Company Report #2003024959
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Incoherent Tremor Vomiting	Consumer	Neurontin (Gabapentin) Hypericum Perforatum (Hypericum Perforatum) Kava-Kava Rhizoma (Kava-Kava Rhizoma)	PS C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211478-0Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2003025236

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Neurontin (Gabapentin)	PS		
				Baclofen (Baclofen)	SS		OTHER
OTHER				Atenolol (Atenolol)	C		
				Bisacodyl (Bisacodyl)	C		
				Docusate Sodium (Docusate Sodium)	C		
				Vitamins	C		
				Magnesium Gluconate (Magnesium Gluconate)	C		

Date:10/14/03ISR Number: 4211479-2Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2003025603

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1500 MG	Atrial Fibrillation Fatigue	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other (TID), ORAL		Herpes Zoster	Professional				
500 MCG (250, BID), ORAL		Urinary Tract Infection		Tikosyn (Dofetilide)	SS		ORAL
				Carvedilol (Carvedilol)	C		
				All Other Therapeutic Products	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		

Date:10/14/03ISR Number: 4211480-9Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #2003025831

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicidal Ideation Tremor	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
100 MG				Lansoprazole (Lansoprazole)	C		
(DAILY), ORAL				Estradiiol (Estradiol)	C		

Date:10/14/03ISR Number: 4211483-4Report Type:Periodic Company Report #2003026010
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness	Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL				Celecoxib (Celecoxib)	SS		ORAL
200 MG				Folic Acid (Folic Acid) Tolterodine	C		
(DAILY), ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

L-Tartrate
 (Tolterodine
 L-Tartrate) C
 Levothyroxine Sodium
 (Levothyroxine
 Sodium) C
 Amitriptyline
 Hydrochloride
 (Amitriptyline
 Hydrochloride) C
 Alendronate Sodium
 (Alendronate Sodium) C
 Quinapril
 Hydrochloride
 (Quinapril
 Hydrochloride) C

Date:10/14/03ISR Number: 4211486-XReport Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2003026753

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), Other ORAL		Suicidal Ideation	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Naproxen (Naproxen) C
 Pantoprazole
 (Pantoprazole) C

Date:10/14/03ISR Number: 4211488-3Report Type:Periodic
 Age:68 YR Gender:Female I/FU:I

Company Report #2003027303

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG (TID), ORAL		Angioneurotic Oedema Swelling	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Bactrim
 (Sulfamethoxazole,

(BID)

Trimethoprim)	SS
Sertraline Hydrochloride (Sertraline Hydrochloride)	C
Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C

Date:10/14/03ISR Number: 4211492-5Report Type:Periodic Company Report #2003020119
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mania	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211496-2Report Type:Periodic
Age:40 YR Gender:Female I/FU:I

Company Report #2003021209

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cataract Visual Disturbance	Health Professional	Neurontin (Gabapentin)	PS		ORAL
400 MG (200, BID), ORAL				All Other Therapeutic Products Sertraline Hydrochloride Clonazepam	SS C C		

Date:10/14/03ISR Number: 4211499-8Report Type:Periodic
Age:49 YR Gender:Female I/FU:I

Company Report #2003021325

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Asthenia Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (BID), ORAL		Fall Feeling Abnormal Head Discomfort Headache Hypersomnia Musculoskeletal Stiffness Nausea Nervousness Pain Paraesthesia Tremor		Propacet (Paracetamol, Dextropropoxyphene Napsilate) Lorazepam	C C		

Date:10/14/03ISR Number: 4211500-1Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2003021715

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Haematochezia	Consumer	Neurontin			

ORAL	Haemoglobin Decreased	(Gabapentin)	PS	ORAL
	Neuropathy Peripheral	Rofecoxib	C	
	Rash	All Other		
	Refractory Anaemia	Therapeutic Products	C	

Date:10/14/03ISR Number: 4211503-7Report Type:Periodic Company Report #2003022165
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Pain Upper	Health	Neurontin			
Other		Chest Pain	Professional	(Gabapentin)	PS		
200 MG		Euphoric Mood					
(DAILY)		Fatigue		Topamax (Topiramate)	SS		
		Feeling Abnormal		Nortriptyline			
		Hepatic Enzyme Increased		(Nortriptyline)	SS		
		Hepatitis		Oxcarbazepine			
		Hepatomegaly		(Oxcarbazepine)	SS		
		Hypotension		Tiagabine			
		Infectious Mononucleosis		Hydrochloride			
		Logorrhoea		(Tiagabine			
		Sleep Disorder		Hydrochloride)	SS		
		Syncope		Carbamazepine			
		Trigeminal Neuralgia		(Carbamazepine)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ascorbic Acid C
 Tocopherol C
 Multivitamin "Lappe"
 (Vitamins Nos) C
 Glucosamine
 W/Chondroitin
 Sulfates (Ascorbic
 Acid, Manganese,
 Chondroitin Sulfate, C

Date:10/14/03ISR Number: 4211507-4Report Type:Periodic Company Report #2003022761
 Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (100, TID), ROAL		Convulsion Vomiting	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Phenytoin Sodium	C		
				Warfarin Sodium	C		
				Oxacarbazepine	C		
				Pantoprazole	C		
				Temazepam	C		
				Simvastatin	C		
				Metoprolol Succinate	C		

Date:10/14/03ISR Number: 4211509-8Report Type:Periodic Company Report #2003022769
 Age:87 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (400, TID)		Confusional State	Health Professional Company Representative	Neurontin (Gabapentin)	PS		
				Ipratropium Bromide (Ipratropium Bromide)	C		
				Rofecoxib (Rofecoxib)	C		

Atorvastatin
(Atorvastatin) C
Levothyroxine Sodium
(Levothyroxine
Sodium) C

Date:10/14/03ISR Number: 4211512-8Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2003023515

Outcome PT
Other Anorexia
Apathy
Body Height Decreased
Chest Pain
Crying
Decreased Activity
Dehydration
Depression
Feeling Abnormal
Headache
Insomnia

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Nightmare Tremor Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Clonazepam (Clonazepam)	C		
				Atorvastatin (Atorvastatin)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		

Date:10/14/03ISR Number: 4211513-XReport Type:Periodic Company Report #2003023518
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG		Anxiety Dyspnoea Pain	Consumer	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211517-7Report Type:Periodic Company Report #2003023568
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Amnesia Anxiety	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Balance Disorder		Fentanyl (Fentanyl)	C		
		Confusional State		Fluoxetine			
		Convulsion		Hydrochloride (Fluoxetine			
		Dyspnoea		Hydrochloride)	C		
		Intentional Misuse		Alprazolam			
		Medication Error		(Alprazolam)	C		
		Multiple Sclerosis		Trimipramine			
		Muscle Spasms		(Trimipramine)	C		
		Palpitations		Calcium (Calcium)	C		
		Panic Attack					

Vision Blurred
Vomiting
Weight Decreased

Vitamins

C

Date:10/14/03ISR Number: 4211522-0Report Type:Periodic Company Report #2003015778
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (DAILY), ORAL		Agitation Depression Disorientation	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
10 MG (DAILY)		Dissociation Drug Interaction	Company Representative	Lipitor (Atorvastatin)	SS		
30 MG (DAILY), ORAL		Dyspnoea Headache Memory Impairment Suicidal Ideation		Paroxetine Hydrochloride (Paroxetine Hydrochloride)	SS		ORAL
				Tolterodine L-Tartrate (Tolterodine			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
8 MG (TWICE DAILY)					L-Tartrate)	SS		
2 MG (DAILY)					Estradiol (Estradiol)	SS		
DAILY					Hydrochlorothiazide (Hydrochlorothiazide)	SS		
					Enalapril (Enalapril)	C		
					Alprazolam (Alprazolam)	C		

Date:10/14/03ISR Number: 4211523-2Report Type:Periodic Company Report #2003015900
 Age:56 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	2700 MG (900, THREE TIMES DAILY), ORAL		Hypokinesia Incoherent Intentional Misuse Medication Error Memory Impairment Oedema Pain Sleep Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:10/14/03ISR Number: 4211525-6Report Type:Periodic Company Report #2003016749
 Age: Gender:Unknown I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Death	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211526-8Report Type:Periodic
Age:44 YR Gender:Female I/FU:I

Company Report #2003017140

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyskinesia	Health	Neurontin			
300 MG		Migraine	Professional	(Gabapentin)	PS		ORAL
(DAILY), ORAL							
				Alprazolam			
				(Alprazolam)	C		
				Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	C		
				Analgesics	C		

Date:10/14/03ISR Number: 4211528-1Report Type:Periodic
Age:55 YR Gender:Male I/FU:I

Company Report #2003017838

Outcome	PT
Hospitalization -	Gastrooesophageal Reflux
Initial or Prolonged	Disease
Other	Hypoaesthesia
	Intentional Misuse
	Medication Error

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Freedom Of Information (FOI) Report

Pain

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
3200 MG		Consumer Health	Neurontin (Gabapentin)	PS		ORAL
(QID), ORAL		Professional	Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
			Pethidine Hydrochloride (Pethidine Hydrochloride)	C		
			Metoprolol (Metoprolol)	C		
			Simvastatin (Simvastatin)	C		

Date:10/14/03ISR Number: 4211531-1Report Type:Periodic Company Report #2003018129
 Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other		Aggression	Professional				
600 MG (TID),		Dizziness		Baclofen (Baclofen)	C		
ORAL		Dreamy State		Desipramine (Desipramine)	C		
		Fatigue		Promethazine (Promethazine)	C		
		Feeling Abnormal					
		Loss Of Consciousness					
		Somnolence					

Date:10/14/03ISR Number: 4211533-5Report Type:Periodic Company Report #2003018673
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other

Pancreatitis

Health
Professional
Company
Representative

Neurontin
(Gabapentin)

PS

Date:10/14/03ISR Number: 4211536-0Report Type:Periodic
Age:76 YR Gender:Female I/FU:I

Company Report #2003018800

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other 3600 MG (TID), ORAL		Depression Intentional Misuse Medication Error Suicide Attempt	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Donnatal (Phenobarbital, Atropine Sulfate, Hyoscine Hydrobromide, Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride,	C C		

Freedom Of Information (FOI) Report

Combivent
 (Ipratropium
 Bromide, Salbutamol
 Sulfate) C
 Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C
 Celecoxib
 (Celecoxib) C
 Ascorbic Acid
 (Ascorbic Acid) C
 Nicotine (Nicotine) C
 Fentanyl (Fentanyl) C
 Levothyroxine Sodium
 (Levothyroxine
 Sodium) C
 Lansoprazole
 (Lasoprazole) C

Date:10/14/03ISR Number: 4211538-4Report Type:Periodic Company Report #2003008954
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Convulsion Migraine	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:10/14/03ISR Number: 4211540-2Report Type:Periodic Company Report #2003009355
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG (DAILY)		Anxiety Blood Thyroid Stimulating Hormone Increased	Consumer	Neurontin (Gabapentin)	PS		
		Cardiac Failure Congestive Decreased Appetite Depersonalisation Depression		Levothyroxine Sodium (Levothyroxine Sodium) Lisinopril (Lisinopril)	C C		

Dyskinesia
Extremity Contracture
Haemorrhage Intracranial
Hostility
Pain
Paranoia
Tongue Blistering
Tooth Discolouration
Visual Disturbance

Date:10/14/03ISR Number: 4211546-3Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2003009705

Outcome PT
Other Alopecia
Decreased Appetite
Dermatitis Contact
Dizziness
Eustachian Tube Disorder
Hunger

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Mucosal Dryness Pain Paraesthesia					
ORAL		Peripheral Coldness Pruritus Generalised	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Psychotic Disorder Sedation Sinus Disorder Somnolence Vision Blurred Visual Disturbance	Professional	Hyoscyamine Sulfate (Hyoscyamine Sulfate) All Other Non-Therapeutic Products	SS SS		
TOPICAL	TOPICAL	Weight Decreased		Estrogens Conjugated (Estrogens Conjugated) Fexofenadine Hydrochloride (Fexofenadine Hydrochloride) Midrid (Paracetamol, Dichloralphenazone, Isometheptene) Estropipate (Estropipate) Cyclobenzaprine (Cyclobenzaprine)	C C C C C		

Date:10/14/03ISR Number: 4211547-5Report Type:Periodic Company Report #2003009898
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Pneumonia Staphylococcal Somnolence	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211549-9Report Type:Periodic Company Report #2003009955
Age: Gender:Female I/FU:I

Outcome	PT
Other	Amnesia

Asthenia
Blood Urea Decreased
Cachexia
Confusional State
Conversion Disorder
Convulsion
Drug Level Decreased
Dysphagia
Eating Disorder
Emotional Distress
Fatigue
Feeling Abnormal
Full Blood Count Abnormal
Headache
Hypersensitivity
Hypoaesthesia
Hypoaesthesia Oral
Nervous System Disorder
Pco2 Decreased
Speech Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG	(DAILY), ORAL	Tinnitus Tremor Vision Blurred	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Trazodone (Trazodone)	C		
				Tramadol Hydrochloride (Tramadol Hydrochloride)	C		
				Naproxen (Naproxen)	C		
				Salbutamol (Salbutamol)	C		
				Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)	C		

Date:10/14/03ISR Number: 4211550-5Report Type:Periodic Company Report #2003010145
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Blindness Diarrhoea	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Dry Mouth Eye Pain Eyelid Ptosis Skin Disorder Somnolence		Cyanocobalamin (Cyanocobalamin)	C		
				Glucosamine (Glucosamine)	C		

Date:10/14/03ISR Number: 4211553-0Report Type:Periodic Company Report #2003010617
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Renal Failure	Consumer	Neurontin		
ORAL	Renal Impairment		(Gabapentin)	PS	ORAL

Date:10/14/03ISR Number: 4211555-4Report Type:Periodic Company Report #2003010675
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder	Health	Neurontin			
300 MG		Deafness	Professional	(Gabapentin)	PS		ORAL

(DAILY), ORAL

Fluoxetine	
Hydrochloride	
(Fluoxetine	
Hydrochloride)	SS
Trazodone	
(Trazodone)	C
Prednisone	
(Prednisone)	C

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Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211557-8Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2003013016

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cellulitis	Health	Neurontin			
ORAL		Polyneuropathy	Professional	(Gabapentin)	PS		ORAL
				Lamivudine			
				(Lamivudine)	C		
				Zidovudine			
				(Zidovudine)	C		
				Lansoprazole			
				(Lansoprazole)	C		
				Nevirapine			
				(Nevirapine)	C		
				Zolpidem (Zolpidem)	C		
				Oxycocet			
				(Paracetamol,			
				Oxycodone			
				Hydrochloride)	C		
				Paracetamol			
				(Paracetamol)	C		

Date:10/14/03ISR Number: 4211560-8Report Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #2003014030

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Glucose Decreased	Consumer	Neurontin			
Initial or Prolonged		Overdose		(Gabapentin)	PS		ORAL
ORAL							
Other							

Date:10/14/03ISR Number: 4211562-1Report Type:Periodic
Age:15 YR Gender:Female I/FU:I

Company Report #2003003090

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Insomnia	Health	Neurontin			
900 MG (300		Muscle Contractions	Professional	(Gabapentin)	PS		ORAL

MG, TID),
 ORAL
 Involuntary
 Paraesthesia
 Swollen Tongue
 Diphenhydramine
 Hydrochloride
 (Diphenhydramine
 Hydrochloride) C
 Ibuprofen
 (Ibuprofen) C
 Methylphenidate
 Hydrochloride
 (Methylphenidate
 Hydrochloride) C

Date:10/14/03ISR Number: 4211565-7Report Type:Periodic Company Report #2003003423
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2700 MG		Hepatic Enzyme Increased Intentional Misuse	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other (TID), ORAL		Medication Error Nausea Vomiting	Company Representative	Sumatriptan Succinate (Sumatriptan			

Freedom Of Information (FOI) Report

Succinate) C
 Cyclobenzaprine
 Hydrochloride
 (Cyclobenzaprine
 Hydrochloride) C

Date:10/14/03ISR Number: 4211568-2Report Type:Periodic
 Age:41 YR Gender:Female I/FU:I

Company Report #2003003628

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Bladder Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Blood Cortisol Abnormal Blood Pressure Decreased Blood Prolactin Abnormal Constipation Coordination Abnormal Cough Fatigue Gastrointestinal Disorder Heart Rate Decreased Hepatic Function Abnormal Hyperhidrosis Hypersomnia Hypoglycaemia Immune System Disorder Neurological Symptom Nocturia Pain Respiratory Rate Decreased Unevaluable Event Urinary Hesitation Urinary Retention Vomiting		Insulin Human Injection, Isophane (Insulin Human Injection Isophane) Vitamins	C C		

Date:10/14/03ISR Number: 4211570-0Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2003003629

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Constipation	Consumer	Neurontin			

ORAL	Pruritus	(Gabapentin)	PS	ORAL
	Somnolence	Psyllium (Psyllium)	SS	ORAL
ORAL	Throat Tightness			

Date:10/14/03ISR Number: 4211578-5Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2003003880

Outcome	PT
Other	Asthenia
	Diarrhoea
	Eructation
	Gingival Disorder
	Intentional Misuse
	Malaise
	Nausea
	Pain
	Pain In Jaw
	Tooth Abscess

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Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
ORAL		Toothache Vomiting	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Tramadol Hydrochloride (Tramadol Hydrochloride)	C		

Date:10/14/03ISR Number: 4211580-3Report Type:Periodic Company Report #2003003985
Age:75 YR Gender:Female I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
Other		Burning Sensation Diarrhoea Rosacea Thrombosis Vein Disorder Vomiting	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211582-7Report Type:Periodic Company Report #2003004299
Age: Gender:Female I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
1800 MG, ORAL		Creatinine Renal Clearance Decreased Fluid Retention	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:10/14/03ISR Number: 4211586-4Report Type:Periodic Company Report #2003006520
Age:69 YR Gender:Female I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
Other		Deafness	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211588-8Report Type:Periodic Company Report #2003007399
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Heart Rate Increased Iron Deficiency Anaemia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
1200 MG (400 MG, TID) ORAL		Myalgia	Professional				
		Nervousness Pain In Extremity		Nortriptyline (Nortriptyline)	SS		ORAL
25 MG (DAILY), ORAL		Tremor					
				Metoprolol Succinate (Metoprolol Succinate)	C		
				Estrogen Nos (Estrogen Nos)	C		
				Progesterone (Progesterone)	C		

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Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211591-8Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2003008953

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Consumer	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211595-5Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2002058716

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG		Fatigue Myocardial Infarction	Consumer	Neurontin (Gabapentin)	PS		ORAL

(DAILY), ORAL

Levothyroxine Sodium (Levothyroxine Sodium)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C
Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Tocopherol (Tocopherol)	C
Ascorbic Acid (Ascorbic Acid)	C
Calcium Carbonate (Calcium Carbonate)	C
Atenolol (Atenolol)	C

Date:10/14/03ISR Number: 4211596-7Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2002059031

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Asthenia	Consumer	Neurontin			

Other	Diabetes Mellitus	(Gabapentin)	PS
900 MG (300			
MG, THREE	Hypothyroidism		
TIMES DAILY),	Nerve Injury		
UNKNOWN	Pain		
	Somnolence	Amitriptyline	
		Hydrochloride	
		(Amitriptyline	
		Hydrochloride)	SS
		Glibomet (Metformin	
		Hydrochloride,	
		Glibenclamide)	SS
10 MG/1000MG			
(TWICE			
DAILY),			
UNKNOWN			
		Insulin Human	
		Injection, Isophane	
		(Insulin Human	
		Injection, Isophane)	SS
25 UNITS			
(DAILY),			
UNKNOWN			
		Levothyroxine Sodium	
		(Levothyroxine	

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Sodium) SS
 Paroxetine
 Hydrochloride
 (Paroxetine
 Hydrochloride) SS

Date:10/14/03ISR Number: 4211598-0Report Type:Periodic
 Age:40 YR Gender:Female I/FU:I

Company Report #2002059376

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Accidental Overdose	Health	Neurontin			
		Dysarthria	Professional	(Gabapentin)	PS		ORAL
900 MG TID		Medication Error					
ORAL							
		Palpitations		Citalopram			
		Paraesthesia		Hydrobromide			
				(Citalopram			
				Hydrobromide)	C		
				Zyrexia (Cetirizine			
				Hydrochloride)	C		
				Clonazepam	C		
				Amlodipine	C		

Date:10/14/03ISR Number: 4211599-2Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #2002059379

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Health	Neurontin			
		Panic Attack	Professional	(Gabapentin)	PS		
		Suicidal Ideation					

Date:10/14/03ISR Number: 4211602-XReport Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2002059417

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Angle Closure Glaucoma	Consumer	Neurontin			
		Balance Disorder		(Gabapentin)	PS		ORAL
ORAL							

Feeling Abnormal
Headache
Mental Status Changes
Pain
Tremor

Simvastatin
(Simvastatin) C
Nizatidine
(Nizatidine) C
Metoprolol Succinate
(Metoprolol
Succinate) C
Lorazepam
(Lorazepam) C
Pilocarpine
Hydrochloride
(Pilocarpine
Hydrochloride) C
Latanoprost
(Latanoprost) C

Date:10/14/03ISR Number: 4211603-1Report Type:Periodic Company Report #2002060604
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Hyperlipidaemia Liver Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1800 MG							

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(DAILY), ORAL

Vicodin
 (Paracetamol,
 Hydrocodone
 Bitartrate) C
 Amitriptyline
 (Amitriptyline) C

Date:10/14/03ISR Number: 4211606-7Report Type:Periodic Company Report #2002060722
 Age:84 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Failure Congestive	Consumer	Neurontin (Gabapentin)	PS		ORAL
800 MG (BID), ORAL		Somnolence					
		Unevaluable Event Weight Increased		Karvea Hct (Hydrochlorothiazide , Irbesartan) Atorvastatin (Atorvastatin) Amlodipine (Amlodipine) Isosorbide Mononitrate (Isosorbide Mononitrate) Furosemide (Furosemide) Atenolol (Atenolol)	C C C C C C		

Date:10/14/03ISR Number: 4211607-9Report Type:Periodic Company Report #2002061353
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety Asthenia	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL		Blood Pressure					

Fluctuation
 Blood Pressure Increased
 Dyspnoea
 Fatigue
 Heart Rate Irregular
 Influenza Like Illness
 Pain In Extremity

Levothyroxine Sodium
 (Levothyroxine
 Sodium) C
 Vicodin
 (Paracetamol,
 Hydrocodone
 Bitartrate) C

Date:10/14/03ISR Number: 4211610-9Report Type:Periodic
 Age:53 YR Gender:Female I/FU:I

Company Report #2002062194

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG Other (DAILY), ORAL	Anxiety Balance Disorder Blood Urine Cholelithiasis Coordination Abnormal Headache Speech Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Levothyroxine Sodium (Levothyroxine Sodium) C Liothyronine Sodium (Liothyronine Sodium) C Acetylsalicylic Acid			

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Freedom Of Information (FOI) Report

(Acetylsalicylic
Acid) C
Paracetamol
(Paracetamol) C
Bupropion
Hydrochloride
(Bupropion
Hydrochloride) C

Date:10/14/03ISR Number: 4211612-2Report Type:Periodic
Age:53 YR Gender:Male I/FU:I

Company Report #2002062287

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Diplopia	Consumer	Neurontin (Gabapentin)	PS		ORAL
1800 MG (TID), ORAL		Impatience					
		Ivth Nerve Paralysis					
		Visual Disturbance		Omeprazole (Omeprazole)	C		
				Sotalol Hydrochloride (Sotalol Hydrochloride)	C		

Date:10/14/03ISR Number: 4211613-4Report Type:Periodic
Age:77 YR Gender:Female I/FU:I

Company Report #2002062352

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anorexia	Health	Neurontin (Gabapentin)	PS		ORAL
300 MG (DAILY), ORAL		Cerebrovascular Accident	Professional				
		Difficulty In Walking					
		Dizziness		Medroxyprogesterone (Medroxyprogesterone)	C		
		Fall					
		Nausea					
		Sedation					
		Vision Blurred					
		Weight Decreased					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intentional Misuse Medication Error	Health Professional	Neurontin (Gabapentin)	PS		ORAL
3600 MG (1200 MG, TID),		Thrombocytopenia	Company Representative				
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Intentional Misuse Medication Error	Health Professional	Neurontin (Gabapentin)	PS		ORAL
2400 MG		Pneumonia					
Other (TID), ORAL				Morphine (Morphine)	C		

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Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211616-XReport Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2002063614

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams	Consumer	Neurontin			
		Fatigue		(Gabapentin)	PS		
		Hallucination		Zoloft (Sertraline)	SS		
		Somnolence		Methadone			
				(Methadone)	C		

Date:10/14/03ISR Number: 4211619-5Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2002064113

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Diabetes Mellitus	Health	Neurontin			
Initial or Prolonged		Gynaecomastia	Professional	(Gabapentin)	PS		
900 MG (300							
MG, TID),							
UNKNOWN							

Date:10/14/03ISR Number: 4211621-3Report Type:Periodic
Age:59 YR Gender:Male I/FU:I

Company Report #2002064295

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Confusional State	Health	Neurontin			
Initial or Prolonged			Professional	(Gabapentin)	PS		
				Duragesic (Fentanyl)	SS		
				Carisoprodol			
				(Carisoprodol)	C		
				Doxepin (Doxepin)	C		
				Lidocaine			
				Hydrochloride			
				(Lidocaine			
				Hydrochloride)	C		
				Testosterone			
				(Testosterone)	C		
				Simvastatin			
				(Simvastatin)	C		

Valdecoxib (Valdecoxib)	C
Tramadol Hydrochloride (Tramadol Hydrochloride)	C
Morphine Sulfate (Morphine Sulfate)	C
Risperidone (Risperidone)	C

Date:10/14/03ISR Number: 4211624-9Report Type:Periodic Company Report #2002066725
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache Hypertension	Health Professional	Neurontin (Gabapentin)	PS		
900 MG (THREE TIMES DAILY)		Visual Disturbance	Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211625-0Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2002067087

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (QD), Other ORAL		Cardiac Failure Congestive	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Hydralazine (Hydralazine)	C		
				Verapamil (Verapamil)	C		
				Atenolol (Atenolol)	C		
				Estrogens Conjugated (Estrogens Conjugated)	C		
				Simvastatin (Simvastatin)	C		
				Levetiracetam (Levetiracetam)	C		
				Ondansetron (Ondansetron)	C		
				Lorazepam (Lorazepam)	C		
				Pantoprazole (Pantoprazole)	C		

Date:10/14/03ISR Number: 4211641-9Report Type:Periodic
Age: Gender: I/FU:I

Company Report #2002067630

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Pancreatitis	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211643-2Report Type:Periodic
Age: Gender: I/FU:I

Company Report #2002067642

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Pancreatitis	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/14/03ISR Number: 4211648-1Report Type:Periodic Company Report #2002068311
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Agitation Chest Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, BID), ORAL		Convulsion Headache Hyperhidrosis Pain		Kaletra (Ritonavir, Lopinavir) Tenofovir Disoproxil Fumarate (Tenofovir Disoproxil Fumarate) Zalcitabine (Zalcitabine)	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lorazepam
 (Lorazepam) C
 Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C

Date:10/14/03ISR Number: 4211652-3Report Type:Periodic Company Report #2002068672
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrioventricular Block	Health Professional Company Representative	Neurontin (Gabapentin) Rofecoxib (Rofecoxib)	PS SS		

Date:10/14/03ISR Number: 4211654-7Report Type:Periodic Company Report #2002068853
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Glucose Increased Ill-Defined Disorder Pain Peripheral Coldness	Consumer Health Professional	Neurontin (Gabapentin) Pravastatin Sodium (Pravastatin Sodium) Metformin Hydrochloride (Metformin Hydrochloride) Insulin Human Injection, Isophane (Insulin Human Ijection, Isophane) Insulin Lispro (Insulin Lispro)	PS C C C C		ORAL
300 MG (DAILY), ORAL							

Date:10/14/03ISR Number: 4211657-2Report Type:Periodic Company Report #2002069510
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cerebrovascular Accident Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Lethargy Pain		Celecoxib (Celecoxib)	SS		ORAL
ORAL				Tramadol Hydrochloride (Tramadol Hydrochloride)	SS		ORAL
ORAL				Isradipine (Isradipine)	C		

Date:10/14/03ISR Number: 4211659-6Report Type:Periodic
Age:67 YR Gender:Female I/FU:I

Company Report #2002073113

Outcome	PT
Other	Dizziness Oral Pain Sciatica

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tooth Injury

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
(DAILY), ORAL		Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Professional	Amiodarone (Amiodarone)	SS		
			Benazepril Hydrochloride (Benazepril Hydrochloride)	C		
			Atenolol (Atenolol)	C		
			Warfarin Sodium (Warfarin Sodium)	C		

Date:10/14/03ISR Number: 4211660-2Report Type:Periodic Company Report #2002073224
 Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Back Pain Chest Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (THREE TIMES DAILY), ORAL		Chills Cough					
		Depression		Valsartan (Valsartan)	C		
		Diarrhoea		Amitriptyline (Amitriptyline)	C		
		Dizziness		Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
		Dyspepsia		Glibomet (Metformin Hydrochloride, Glibenclamide)	C		
		Dyspnoea		Diazepam (Diazepam)	C		
		Gastric Disorder		Vicodin (Paracetamol, Hydrocodone Bitartrate)	C		
		Headache					
		Hyperhidrosis					
		Insomnia					
		Memory Impairment					
		Neck Pain					
		Pain					
		Rash					
		Retching					
		Weight Increased					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Consumer	Neurontin			
Other		Dyskinesia		(Gabapentin)	PS		
		Feeling Hot		Carbamazepine			
		Hyperhidrosis		(Carbamazepine)	SS		
		Malaise		Fluoxetine			
		Respiratory Disorder		Hydrochloride			
				(Fluoxetine			
				Hydrochloride)	SS		
				All Other			
				Therapeutic Products	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211667-5Report Type:Periodic
Age:65 YR Gender:Female I/FU:I

Company Report #2003000514

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (TID), ORAL	Dry Mouth Hypersensitivity Urticaria	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL			Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride)	SS		ORAL
			Cardura (Doxazosin) (Doxazosin Mesilate) Levothyroxine Sodium (Levothyroxine Sodium) Acetylsalicylic Acid (Acetylsalicylic Acid) Vicodin (Paracetamol, Hydrocodone Bitartrate)	C C C C		

Date:10/14/03ISR Number: 4211669-9Report Type:Periodic
Age:82 YR Gender:Female I/FU:I

Company Report #2003000955

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Asthenia Fall Nausea Vomiting	Consumer	Neurontin (Gabapentin) Vicodin (Paracetamol, Hydrocodone Bitartrate) Sertraline Hydrochloride (Sertraline Hydrochloride) Amitriptyline (Amitriptyline)	PS C C C		

Rivastigmine
 Tartrate
 (Rivastigmine
 Tartrate) C
 Donepezil
 Hydrochloride
 (Donepezil
 Hydrochloride) C

Date:10/14/03ISR Number: 4211672-9Report Type:Periodic Company Report #2003001245
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other (TID), ORAL		Laboratory Test Abnormal Syncope	Health Professional Company Representative	Neurontin (Gabapentin) Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride)	PS C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211678-XReport Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2002054216

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Somnolence	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:10/14/03ISR Number: 4211679-1Report Type:Periodic
 Age:80 YR Gender:Male I/FU:I

Company Report #2002054218

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anticonvulsant Drug Level Below Therapeutic	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, THREE TIMES DAILY),		Cerebrovascular Accident					
ORAL		Convulsion					
		Lethargy					
ORAL		Pyrexia Staphylococcal Infection		Dilantin (Phenytoin Sodium)	SS		ORAL
		Weight Decreased		Warfarin Sodium (Warfarin Sodium)	C		

Date:10/14/03ISR Number: 4211682-1Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2002054242

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract Diabetes Mellitus	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Non-Insulin-Dependent Dizziness		Ibuprofen (Ibuprofen)	C		
		Ill-Defined Disorder		Propranolol Hydrochloride			
		Intentional Misuse		(Propranolol Hydrochloride)	C		
		Medication Error					
		Neuralgia					

Pain
Pain In Extremity
Weight Increased

Hydrochlorothiazide
(Hydrochlorothiazide
) C

Date:10/14/03ISR Number: 4211684-5Report Type:Periodic Company Report #2002054672
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Blindness	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other	900 MG (THREE TIMES A DAY), ORAL	Cerebrovascular Accident					

Verapamil
(Verapamil) C
Metformin
(Metformin) C
Glipizide
(Glipizide) C
Unknown Ointment C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211688-2Report Type:Periodic
Age:31 YR Gender:Male I/FU:I

Company Report #2002055132

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (400 Other MG, THREE TIMES DAILY), ORAL		Hallucination Sleep Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Clonazepam (Clonazepam)	C		
				Nefazodone Hydrochloride (Nefazodone Hydrochloride)	C		
				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
				Celecoxib (Celecoxib)	C		

Date:10/14/03ISR Number: 4211691-2Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2002056353

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 300 MG (100 MG, TID), ORAL		Amnesia Fatigue Loss Of Consciousness Somnolence	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:10/14/03ISR Number: 4211693-6Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2002056555

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Intentional Misuse Medication Error	Health Professional	Neurontin (Gabapentin)	PS		ORAL
2400 MG, ORAL		Red Blood Cell Count Decreased White Blood Cell Count Decreased		Isosorbide Dinitrate (Isosorbide Dinitrate) Insulin (Insulin) Acetylsalicyclic Acid (Acetylsalicyclic Acid)	C C C		

Date:10/14/03ISR Number: 4211695-XReport Type:Periodic Company Report #2002056631
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Depression Grand Mal Convulsion	Consumer Health	Neurontin (Gabapentin)	PS		
900 MG (300 MG,TID)		Insomnia Myalgia Somnolence	Professional	Amitriptyline (Amitriptyline)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/03ISR Number: 4211697-3Report Type:Periodic
Age:64 YR Gender:Male I/FU:I

Company Report #2002057057

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fluid Retention	Consumer	Neurontin			
2400 MG, (800		Intentional Misuse		(Gabapentin)	PS		ORAL
MG,TID), ORAL		Unevaluable Event					
				Quinapril			
				Hydrochloride			
				(Quinapril			
				Hydrochloride)	C		

Date:10/14/03ISR Number: 4211699-7Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #2002057069

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Face Oedema	Consumer	Neurontin			
Initial or Prolonged		Pharyngeal Oedema	Company	(Gabapentin)	PS		
Other			Representative				

Date:10/15/03ISR Number: 4208719-2Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0311384A
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Agitation		Deroxat	PS	Glaxosmithkline	ORAL
20MG Three							
Life-Threatening		Death					
times per day				Noctamid	SS		ORAL
2MG Twice per		Fall					
day		Sudden Death					
				Valium	SS		ORAL
10MG Three							
times per day				Tercian	SS		
INTRAMUSCULAR	100MG Per day						

300MG Five		Neurontin	SS	ORAL
times per day				
50MG Unknown		Gardenal	SS	ORAL

Date:10/15/03ISR Number: 4210435-8Report Type:Expedited (15-DaCompany Report #2003035218
 Age:48 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG Other (TID), ORAL	Condition Aggravated Neuropathic Pain Pain In Extremity Peripheral Nerve Operation	Foreign Consumer	Gabapentin (Gabapentin) Tramadol Fluoxetine Hydrochloride	PS C C		ORAL

Date:10/15/03ISR Number: 4210446-2Report Type:Expedited (15-DaCompany Report #2003028218
 Age:30 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG (600 Other ,TID),ORAL	Abortion Spontaneous Complications Of Maternal Exposure To Therapeutic Drugs Intra-Uterine Death	Foreign Health Professional	Neurontin (Gabapentin) Folic Acid (Folic Acid) Phenobarbital(Phenob arbital) Ferrous Sulfate	PS C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Ferrous Sulfate) C

Date:10/15/03ISR Number: 4210447-4Report Type:Expedited (15-DaCompany Report #2003111363

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abortion Spontaneous	Foreign	Neurontin			
Congenital Anomaly		Congenital Anomaly	Health				
1800 MG		Intra-Uterine Death	Professional	(Gabapentin)	PS		ORAL
(TID), ORAL		Maternal Drugs Affecting					
		Foetus		Folic Acid			
				(Folic Acid)	C		
				Phenobarbital			
				(Phenobarbital)	C		
				Ferrous Sulfate			
				(Ferrous Sulfate)	C		

Date:10/15/03ISR Number: 4210697-7Report Type:Expedited (15-DaCompany Report #2003041722

Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Coordination Abnormal	Consumer	Neurontin			
Other		Dystonia		(Gabapentin)	PS		ORAL
600 MG		Memory Impairment					
(DAILY), ORAL		Tremor		Lipitor			
				(Atorvastatin)	C		
				Clopidogrel Sulfate			
				(Clopidogrel			
				Sulfate)	C		
				Toprol (Metoprolol)	C		
				Ace Inhibitor Nos	C		

Date:10/15/03ISR Number: 4210700-4Report Type:Expedited (15-DaCompany Report #2003041701

Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - ORAL		Cardiac Arrest Grand Mal Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
Initial or Prolonged Other		Systemic Lupus Erythematosus		Dilantin (Phenytoin Sodium) Carbamazepine (Carbamazepine) Ciprofloxacin (Ciprofloxacin) Antiinfectives	SS C C C		

Date:10/15/03ISR Number: 4210751-XReport Type:Expedited (15-DaCompany Report #2003035920
Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability Other		Brain Abscess Extradural Abscess	Consumer	Neurontin (Tablets) (Gabapentin) All Other Therapeutic Products	PS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/03ISR Number: 4210861-7Report Type:Expedited (15-DaCompany Report #2003110156
Age:15 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Basedow'S Disease	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/15/03ISR Number: 4210862-9Report Type:Expedited (15-DaCompany Report #2003110159
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bladder Disorder Cystitis Interstitial	Consumer	Neurontin (Gabapentin)	PS		ORAL

3200 MG (800,

FOUR TIMES A

DAY), ORAL

Furosemide	C
Sertraline	
Hydrochloride	C
All Other	
Therapeutic Products	C
Potassium	C
Trazodone	C
Modafinil	C
Methylphenidate	
Hydrochloride	C
Topiramate	C
Interferon Beta	C
Mitoxantrone	
Hydrochloride	C

Date:10/15/03ISR Number: 4211248-3Report Type:Expedited (15-DaCompany Report #2003109879
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatinine Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL

2400 MG, ORAL

Company
Representative

Date:10/15/03ISR Number: 4211253-7Report Type:Expedited (15-DaCompany Report #2003032129
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blindness	Consumer	Neurontin			
Other		Condition Aggravated	Health	(Gabapentin)	PS		ORAL
200 MG (BID),		Vitamin A Decreased	Professional				
ORAL				Retinol (Retinol)	C		
				Venlafaxine			
				Hydrochloride			
				(Venlafaxine			
				Hydrochloride)	C		
				Bupropion			
				Hydrochloride			
				(Bupropion			
				Hydrochloride)	C		
				Multivitamins			
				(Ergocalciferol,			
				Ascorbic Acid, Folic			
				Acid, Thiamine			

Freedom Of Information (FOI) Report

Hydrochloride, C

Date:10/15/03ISR Number: 4211566-9Report Type:Expedited (15-DaCompany Report #2003110353
Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG (BID); Other ORAL	Alanine Aminotransferase Aspartate Aminotransferase Asthenia Hepatotoxicity Lethargy Nausea	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Benzodiazepine Derivatives Hypericum Perforatum (Hypericum Perforatum) Herbal Preparation Ingredients Unknown	C C C		

Date:10/15/03ISR Number: 4211815-7Report Type:Expedited (15-DaCompany Report #2003039144
Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 1200 MG (DAILY), ORAL	Carpal Tunnel Syndrome Condition Aggravated Dry Mouth Fungal Infection Glossodynia Pharmaceutical Product Complaint Postoperative Infection Tongue Blistering Tongue Dry	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Vicodin (Paracetamol, Hydrochloride Bitartrate) Cefazolin Sodium (Cefazolin Sodium)	C C		

Date:10/15/03ISR Number: 4211817-0Report Type:Expedited (15-DaCompany Report #2003110583
Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Delirium	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other ORAL		Drug Interaction					
		Gastrointestinal Carcinoma		Morphine Sulfate (Morphine Sulfate)	SS		ORAL
ORAL		Hypotension		Calcium (Calcium)	C		
		Weight Decreased		Centrum Silver (Ascorbic Acid, Tocopheryl Acetate, Retinol, Zinc, Calcium, Vitamins Atenolol (Atenolol)	C C		
				Docusate Sodium (Docusate Sodium)	C		
				Limbitrol (Chlordiazepoxide, Amitriptyline Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/03ISR Number: 4212154-0Report Type:Expedited (15-DaCompany Report #2003110422

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	3000 MG	Convulsion Hallucination	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other (TID), ORAL				Dilantin (Phenytoin Sodium)	SS		ORAL
200 MG (BID), ORAL				Phenobarbital (Phenobarbital)	SS		ORAL
90 MG (BID), ORAL							

Date:10/15/03ISR Number: 4212155-2Report Type:Expedited (15-DaCompany Report #2003110413

Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abdominal Pain Arthralgia Back Pain Drug Ineffective Dysuria Kidney Infection Menstruation Irregular Metrorrhagia Nephrolithiasis Oedema Peripheral Ovarian Cyst Ruptured Pain In Extremity Peripheral Coldness Pregnancy Test Positive Skin Discolouration Stress	Consumer	Neurontin (Gabapentin) Zonisamide (Zonisamide) Cilest (Ethinylestradiol, Norgestimate)	PS SS C		

Date:10/15/03ISR Number: 4212156-4Report Type:Expedited (15-DaCompany Report #2003110107
Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 3600 MG Other (TID), ORAL	Blood Glucose Increased Breast Cancer Renal Impairment	Health Professional	Neurontin (Gabapentin) Hydrocodone (Hydrocodone)	PS C		ORAL

Date:10/15/03ISR Number: 4213490-4Report Type:Expedited (15-DaCompany Report #2003110336
Age:75 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Balance Disorder Cerebellar Syndrome	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/03ISR Number: 4213534-XReport Type:Expedited (15-DaCompany Report #2003168755FR

Age:63 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG, BID, ORAL	Anaemia Erythema Platelet Count Increased	Foreign Health Professional	Zyvoxid(Linezolid) Tablet	PS		ORAL
300 MG, BID, ORAL	Thrombocytopenia	Other	Neurontin(Gabapentin)	SS		ORAL
500 MG, ORAL			Doliprane(Paracetamo l)	SS		ORAL
1 DF, BID, ORAL			Urbanyl(Clobizam)	SS		ORAL
400 MG, BID, ORAL			Peflacine(Pefloxacin Mesilate)	SS		ORAL
			Domperidone (Domperidone)	C		
			Cetirizine (Cetirizine)	C		
			Amphotericin B	C		

Date:10/16/03ISR Number: 4211583-9Report Type:Expedited (15-DaCompany Report #2003110073

Age:35 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Life-Threatening 1500 MG (5 Hospitalization - TIMES A DAY), Initial or Prolonged ORAL	Fall Nervous System Disorder Restlessness	Foreign Health Professional Other	Neurontin (Gabapentin)	PS		ORAL

Other		Lormetazepam (Lormetazepam)	SS	ORAL
4 MG (BID),				
ORAL		Diazepam (Diazepam)	SS	ORAL
30 MG (TID),				
ORAL		Cyamemazine (Cyamemazine)	SS	
INTRAMUSCULAR	100 MG			
(ONCE),				
INTRAMUSCULAR				
		Phenobarbital Sodium (Phenobarbital Sodium)	SS	ORAL
50 MG, ORAL				
		Paroxetine Hydrochloride (Paroxetine Hydrochloride)	SS	ORAL
60 MG (TID),				
ORAL				

Date:10/16/03ISR Number: 4212631-2Report Type:Expedited (15-DaCompany Report #US-SHR-03-009149
Age:57 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Arrhythmia
Initial or Prolonged	Asthenia
	Condition Aggravated
	Dehydration
	Depressed Level Of
	Consciousness
	Fatigue

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Event	Report Source	Product	Role	Manufacturer	Route
		Feeling Cold Hypotension Loss Of Consciousness					
		Medication Error Multiple Sclerosis Nausea	Consumer Health Professional	Betaseron (Interferon Beta-1b) Injection	PS		
SUBCUTANEOUS	8 M IU, EVERY	Respiration Abnormal					
2D,		Somnolence					
SUBCUTANEOUS		Sudden Onset Of Sleep		Lasix (Furosemide)	SS		
40 MG, 1X/DAY		Syncope Vomiting		Neurontin (Gabapentin)	SS		
SEE IMAGE				Provigil (Modafinil)	C		
				Macrochantin (Nitrofurantoin)	C		
				Mirapex "Pharmacia-Upjohn" (Pramipexole Dihydrochloride)	C		
				Bextra (Bucindolol Hydrochloride)	C		
				Wellbutrin (Amfebutamone Hydrochloride)	C		
				Zanaflex (Tizanidine Hydrochloride)	C		
				Sinemet (Carbidopa, Levodopa)	C		
				Potassium Chloride	C		
				Baclofen	C		
				Percocet (Oxycodone Hydrochloride)	C		
				Celexa (Citalopram Hydrobromide)	C		

Date:10/17/03ISR Number: 4210721-1Report Type:Expedited (15-DaCompany Report #WAES 0310USA01437
Age:46 YR Gender:Male I/FU:I

Outcome PT
Hospitalization - Amyotrophy

Initial or Prolonged

Asthenia
Calculus Bladder
Dehydration
Gynaecomastia
Hepatic Steatosis
Hepatitis
Hepatomegaly
Hyperlactacidaemia
Hypersensitivity
Hypoaesthesia
Lactic Acidosis
Lipoatrophy
Liver Function Test
Abnormal
Muscle Spasms
Nervous System Disorder
Pancreatitis
Paraesthesia
Phlebothrombosis
Pyrexia
Tinnitus

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Vertigo Vomiting Weight Decreased	Report Source	Product	Role	Manufacturer	Route
				Crixivan	PS		ORAL
				Videx	SS	Merck & Co., Inc	
				Sustiva	SS		ORAL
				Sustiva	SS		ORAL
				Lamivudine	SS		
				Kaletra	SS		
				Riboflavin	SS		
				Zerit	SS		
				Thiamine	SS		
				Zidovudine	SS		
				Vitamin E	SS		
				Bactrim	SS		
				Selenium			
				(Unspecified)	SS		
157	DAY			Ribavirin	SS		
				L-Carnitine	SS		
				Interferon			
				(Unspecified)	SS		
157	DAY			Neurontin	SS		
				Plaquenil	SS		

Date:10/17/03ISR Number: 4213885-9Report Type:Expedited (15-DaCompany Report #2003110823
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 10 MG Other (DAILY), ORAL		Drug Interaction - Erythema	Health Professional	Lipitor (Atorvastatin)	PS		ORAL
300 MG (DAILY), ORAL		Flushing Nausea Nervousness		Neurontin (Gabapentin)	SS		ORAL
1000 MG		Pain Of Skin Post Procedural Complication		Nicotinic Acid (Nicotinic Acid)	SS		ORAL

(DAILY), ORAL	Radicular Pain		
	Skin Warm	Oxycocet	
	Spinal Column Stenosis	(Paracetamol,	
	Surgical Procedure	Oxycodone	
	Repeated	Hydrochloride)	SS
(Q4H), ORAL			ORAL
	Ulcer	Ezetimibe	
	Weight Decreased	(Ezetimibe)	C
		Multivitamins	
		(Ergocalciferol,	
		Ascorbic Acid, Folic	
		Acid, Thiamine	
		Hydrochloride,	C
		Glucosamine	
		W/Chondroitin	
		Sulfates (Ascorbic	
		Acid, Manganese,	
		Chondroitin Sulfate,	C
		Tocopherol	
		(Tocopherol)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/17/03ISR Number: 4214007-0Report Type:Expedited (15-DaCompany Report #2003110465

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Depressed Level Of Consciousness	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
	Drug Toxicity Simple Partial Seizures	Professional				

Date:10/17/03ISR Number: 4214232-9Report Type:Expedited (15-DaCompany Report #2003110574

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 900 MG (DAILY), ORAL	Neuroleptic Malignant Syndrome	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Professional				
			Thioridazine Hydrochloride (Thioridazine Hydrochloride)	SS		ORAL
			Co-Diovan (Hydrochlorothiazide , Valsartan)	SS		
			Clodronate Sodium (Clodronate Sodium)	SS		
			Oxybutynin (Oxybutynin)	SS		ORAL
25 MG (DAILY), ORAL						
			Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
			Fentanyl (Fentanyl)	C		
10 MG (BID), ORAL			Bisoprolol Fumarate (Bisoprolol			

Fumarate) C
 Esomeprazole C
 (Esomeprazole)
 Magnesium
 Trisilicate
 (Magnesium
 Trisilicate) C
 Nulytely (Sodium
 Bicarbonate,
 Potassium Chloride,
 Sodium Chloride,
 Macrogol) C

Date:10/17/03ISR Number: 4214233-0Report Type:Expedited (15-DaCompany Report #2003110281
 Age:83 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cardiovascular Disorder Fall Haematoma Injury Muscle Twitching Pyrexia	Foreign Health Professional	Neurontin (Gabapentin) Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride,	PS C		

Freedom Of Information (FOI) Report

Calcium Acetate (Calcium Acetate)	C
Folic Acid (Folic Acid)	C
Doxepin Hydrochloride (Doxepin Hydrochloride)	C
Insulin Human Injection, Isophane (Insulin Human Injection, Isophane)	C
Esomeprazole (Esomeprazole)	C
Panadeine Co (Codeine Phosphate, Paracetamol)	C
Antiinfectives Piritramide (Piritramide)	C
Ferric Sodium Gluconate Complex (Ferric Sodium Gluconate Complex)	C
Epoetin Alfa (Epoetin Alfa)	C
Carbamazepine (Carbamazepine)	C
Diazepam (Diazepam)	C
Vancomycin (Vancomycin)	C
Blinded Therapy	C

Date:10/17/03ISR Number: 4214584-XReport Type:Expedited (15-DaCompany Report #2003111053

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Abasia Cerebrovascular Accident	Health Professional	Dilantin Infatabs (Phenytoin Sodium)	PS		ORAL
300 MG (TID),		Drug Ineffective Movement Disorder		Neurontin (Tablets) (Gabapentin)	SS		ORAL

ORAL

Myocardial Infarction

Night Cramps

Pain

Pain In Extremity

Poor Peripheral

Circulation

Sensory Disturbance

Phenobarbital

(Phenobarbital)

SS

Clopidogrel Sulfate

(Clopidogrel

Sulfate)

C

Metoprolol Tartrate

(Metoprolol

Tartrate)

C

Acetylsalicylic Acid

(Acetylsalicylic

Acid)

C

Magnesium

(Magnesium)

C

Calcium (Calcium)

C

Thiamine (Thiamine)

C

Insulin (Insulin)

C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/17/03ISR Number: 4214628-5Report Type:Expedited (15-DaCompany Report #2003030919

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Hospitalisation	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/20/03ISR Number: 4214580-2Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20030904720

Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest Depressed Level Of Consciousness	Literature Health Professional	Risperdal (Risperidone) Unspecified	PS		ORAL
ORAL		Hypotension Multi-Organ Failure Tachycardia	Distributor	Valproic Acid (Valproic Acid) Clonazepam (Clonazepam) Atorvastatin (Atorvastatin) Gabapentin (Gabapentin) Clozapine (Clozapine) Nizatidine (Nizatidine) Trihexyphenidyl (Trihexyphenidyl)	SS SS SS SS		

Date:10/20/03ISR Number: 4214681-9Report Type:Expedited (15-DaCompany Report #2003039135

Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100 MG (DAILY), ORAL		Abdominal Pain Upper Coma Cough	Consumer Health Professional	Zoloft (Sertraline) Neurontin	PS		ORAL

(TID), ORAL

Depression

(Gabapentin)

SS

ORAL

Diarrhoea
Drug Withdrawal Syndrome
Fatigue
Feeling Jittery
Flushing
Grand Mal Convulsion
Hyperhidrosis
Hypoaesthesia
Menopausal Symptoms
Retching
Tremor

Date:10/20/03ISR Number: 4214691-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Actinic Keratosis
Disability	Alanine Aminotransferase
Other	Increased
	Alcoholism

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (BID),		Anxiety Arthropathy Aspartate	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Aminotransferase Increased	Professional				
		Astrocytoma	Company Representative	Valproate Sodium (Valproate Sodium)	SS		
		Blood Urea Nitrogen/Creatinine Ratio Increased		Buspirone Hydrochloride (Buspirone Hydrochloride)	C		
		Brain Neoplasm Coma		Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		
		Conversion Disorder Convulsion		Omeprazole (Omeprazole)	C		
		Cyst Depressed Level Of Consciousness		Capsaicin (Capsaicin)	C		
		Ependymoma Folliculitis					
		Gamma-Glutamyltransferase Increased					
		Hypoglycaemia Meniscus Lesion					
		Neoplasm Nervous System Disorder					
		Neurodegenerative Disorder					
		Osteopenia Patellofemoral Pain Syndrome					
		Pigmented Naevus Seborrhoeic Keratosis					
		Sinusitis Sleep Disorder					

Date:10/21/03ISR Number: 4215698-0Report Type:Expedited (15-DaCompany Report #2003111407

Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Gastric Haemorrhage					
900 MG (EVERY							

8 HOURS),

ORAL

Dilantin (Phenytoin Sodium)	SS
Atenolol	C
Lipitor (Atorvastatin)	C
All Other Therapeutic Products	C
Benazepril Hydrochloride	C
Warfarin Sodium	C
Nasal Preparations	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/21/03ISR Number: 4215795-XReport Type:Expedited (15-DaCompany Report #HQWYE099009OCT03

Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG 1X PER		Ankle Fracture Cerebral Atrophy	Health Professional	Cordarone (Amiodarone, Tablet)	PS		ORAL
1 DAY ORAL		Confusional State	Other				
TRANSDERMAL	20 MG 1X PER	Disorientation Disturbance In Attention		Buprenorphine (Buprenorphine)	SS		
1 DAY		Drug Interaction					
TRANSDERMAL	11 DAY	Encephalopathy					
900 MG 1X PER		Fall Myoclonus		Neurontin (Gabapentin,)	SS		ORAL
1 DAY ORAL							
900 MG 1X PER				Neurontin (Gabapentin,)	SS		ORAL
1 DAY ORAL							
				Antra (Omeprazole)	C		
				Selipran (Pravastatin Sodium)	C		
				Lasix (Furosemide)	C		
				Nitroderm Tts (Glyceryl Trinitrate)	C		
				Aspirin (Acetylsalicylic Acid)	C		
				Sintrom (Acenocoumarol)	C		
				Tramadol - Slow Release (Tramadol Hydrochloride)	C		

Date:10/22/03ISR Number: 4213556-9Report Type:Expedited (15-DaCompany Report #PHBS2003CH11427

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 37440MIN		Coagulopathy Condition Aggravated		Tegretol	PS	Novartis Sector: Pharma	ORAL
Initial or Prolonged UNKNOWN		Eosinophilia 27360MIN		Neurontin	SS		
UNKNOWN		Erythema		Huminsulin Basal	C		
UNKNOWN		Exanthem		Insulin Lispro	C		
UNKNOWN		Haematoma		Dafalgan	C		
		Hepatocellular Damage					
		Hypersensitivity					
		Hyponatraemia					
		Leukocytosis					
		Liver Function Test Abnormal					
		Petechiae					
		Polycythaemia					
		Pyrexia					
		Rash Maculo-Papular					
		Shift To The Left					
		Sinus Tachycardia					
		Skin Oedema					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/22/03ISR Number: 4215412-9Report Type:Direct
Age:61 YR Gender:Male I/FU:I

Company Report #CTU 204279

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	300 MG 3	Activities Of Daily		Neurontin 300 Mg	PS		ORAL
		Living Impaired					
	TIMES ORAL						
	600 MG 3	Agitation		Neurontin 600 Mg	SS		ORAL
		Belligerence					
	TIMES ORAL						
		Confusional State		Tizanidine	C		
		Dementia		Ms Contin	C		
		Diplopia		Lopressor	C		
		Disturbance In Attention		Zestril	C		
		Memory Impairment		Norvasc	C		
		Vision Blurred		Actos	C		
				Zocor	C		
				Hydrothiazide	C		
				Aspirin	C		
				Vancomycin	C		
				Soma	C		

Date:10/22/03ISR Number: 4216637-9Report Type:Expedited (15-DaCompany Report #2003111309
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	2700 MG ORAL	Clumsiness	Consumer	Neurontin			
		Drug Effect Decreased		(Gabapentin)	PS		ORAL
		Headache		Baclofen	C		
		Irritability		Amantadine	C		
		Tremor		Pantoprazole	C		
		Weight Increased		Gemfibrozil	C		
				Atorvastatin	C		
				Atenolol	C		
				Zolpidem Tartrate	C		
				Paroxetine			
				Hydrochloride	C		
				Oxycocet			
				(Paracetamol,			
				Oxycodone			

Hydrochloride) C
Axotal (Old Form)
(Caffeine,
Butalbital,
Paracetamol) C
Promethazine
Hydrochloride C
Paracetamol C
Ibuprofen C
Triamcinolone
Acetonide C

Date:10/22/03ISR Number: 4217067-6Report Type:Expedited (15-DaCompany Report #KII-2003-0003985
Age:45 YR Gender:Female I/FU:I

Outcome PT
Hospitalization - Blood Glucose Abnormal
Initial or Prolonged Body Temperature
Decreased
Body Temperature
Increased
Cellulitis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Screen Positive Dyspnoea Lethargy	Report Source	Product	Role	Manufacturer	Route
80 MG		Mental Status Changes Somnolence White Blood Cell Count Increased	Health Professional	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		
100 MG				Neurontin (Gabapentin)	SS		
300 MG				Effexor (Venlafaxine Hydrochloride)	SS		
10 MG, TID				Vicodin (Paracetamol, Hydrocodone Bitartrate)	SS SS		
100 MG, HS				Valium (Diazepam)	SS		
				Trazodone (Trazodone)	SS		
20 MG				Flexeril (Cyclobenzaprine Hydrochloride)	SS		
75 MG, BID				Diclofenac (Diclofenac) Topamax (Topiramate)	SS SS		

Date:10/22/03ISR Number: 4217100-1Report Type:Expedited (15-DaCompany Report #2002068598
Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (THREE TIMES DAILY), ORAL		Asthma Bronchitis Acute Condition Aggravated Drug Interaction	Foreign Study Health Professional	Gabapentin (Gabapentin)	PS		ORAL

Nasopharyngitis

Carbamazepine

(Carbamazepine)	C
Clobazam	
(Clobazam)	C
Theophylline	
(Theophylline)	C
Pranlukast	
(Pranlukast)	C
Salbutamol	
(Salbutamol)	C
Budesonide	
(Budesonide)	C
Tulobuterol	
Hydrochloride	C

Date:10/22/03ISR Number: 4217598-9Report Type:Expedited (15-DaCompany Report #2003016843

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Consumer	Dilantin (Phenytoin			
		Diplopia	Health	Sodium)	PS		ORAL
300 MG (BID),		Dizziness	Professional				
ORAL		Headache		Neurontin			
		Nervous System Disorder		(Gabapentin)	SS		
1200 MG (BID)		Overdose					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/23/03ISR Number: 4215734-1Report Type:Direct
Age:76 YR Gender:Male I/FU:I

Company Report #CTU 204397

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 300 MG BID Initial or Prolonged PRN BY MOUTH	Dysarthria Gait Disturbance Somnolence		Gabapentin	PS		ORAL
			Felodipine	C		
			Celecoxib	C		
			Calcitroil	C		
			Terazosin Hcl	C		
			Metolazone	C		
			Calcium Carbonate	C		
			Sertraline Hcl	C		
			Amoxicillin/Clavulan ate K	C		
			Furosemide	C		
			Clorazepate	C		
			Calcium Acetate	C		
			Triamcinolone Acet Oint	C		
			Carvedilol	C		
			Aspirin Low Dose	C		

Date:10/23/03ISR Number: 4217289-4Report Type:Direct
Age:42 YR Gender:Female I/FU:I

Company Report #CTU 204520

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 300 MG TID BY MOUTH	Anxiety Depression Hyperreflexia Mood Swings Multiple Sclerosis Muscle Spasms Nausea Nervous System Disorder Panic Attack Paraesthesia Photophobia Restlessness		300mg Neurontin T.I.D. Parke-Davis	PS	Parke-Davis	ORAL

Supraventricular
Extrasystoles
Ventricular Extrasystoles
Weight Decreased

Date:10/24/03ISR Number: 4217148-7Report Type:Expedited (15-DaCompany Report #PHFR2003GB03959

Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death		Trileptal	PS	Novartis Sector: Pharma	
UNKNOWN				Gabapentin	SS		
UNKNOWN				Vigabatrin	SS		
UNKNOWN				Alcohol	SS		ORAL
				Lamotrigine	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/24/03ISR Number: 4219613-5Report Type:Expedited (15-DaCompany Report #2003111244

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Abnormal Behaviour Aggression Agitation Nervousness	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/24/03ISR Number: 4219615-9Report Type:Expedited (15-DaCompany Report #2003111978

Age:85 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Autoimmune Disorder Blood Electrolytes	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
	Abnormal Dehydration Hypotension Infection Mood Disorder Due To A General Medical Condition Pyrexia Renal Failure Shock	Professional	Pramipexole Dihydrochloride (Pramipexole Dihydrochloride) Levodopa (Levodopa)	C C		

Date:10/24/03ISR Number: 4219672-XReport Type:Expedited (15-DaCompany Report #K200301649

Age:62 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other SEE IMAGE	Flushing Throat Tightness	Consumer	Levoxyl(Levothyroxin e) Tablet, 75 Mcg	PS		ORAL
300 MG, PRN, ORAL			Neurontin(Gabapentin)300 Mg	SS		ORAL
			Albuterol			

(Salbutamol) C
Premarin (Estrogens
Conjugated) C

Date:10/24/03ISR Number: 4219821-3Report Type:Expedited (15-DaCompany Report #USA-2002-0002855
Age:51 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Anxiety
Initial or Prolonged	Asthenia
	Blood Creatine
	Phosphokinase Mb
	Increased
	Cardiac Failure
	Congestive
	Coagulopathy
	Coma
	Confusional State
	Decreased Appetite
	Depression
	Disturbance In Attention
	Drug Abuser
	Drug Dependence

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Drug Screen Positive Drug Withdrawal Syndrome Dyspnoea				
		Dysuria Encephalopathy Fall Head Injury	Consumer Health Professional Other			
SEE IMAGE			Oxycontin Tablets(Oxycodone Hydrochloride)Cr Tablet	PS		ORAL
		Headache Hepatic Necrosis Hyperhidrosis Hypokalaemia				
SEE IMAGE			Oxycontin Tablets 20 Mg (Oxycodone Hydrochloride) Cr Tablet	SS		ORAL
		Hypotension Insomnia Klebsiella Sepsis Mood Swings	Hydrocodone Cp(Phenylephrine, Chlorpehnamine Maleate)Syrup	SS		ORAL
ORAL						
		Multi-Organ Failure Multiple Drug Overdose	Darvocet-N (Paracetamol)Tablet	SS		ORAL
ORAL	7	DAY				
		Nightmare Oedema Peripheral	Cyclobenzaprine(Cycl obenzaprine) Tablet	SS		ORAL
ORAL						
		Pain Paranoia Pneumonia	Hydroxyzine Pamoeate(Hydroxyzine Embonate)	SS		ORAL
ORAL						
		Pneumonia Aspiration Prothrombin Level	Valium(Diazepam) Tablet	SS		ORAL
ORAL						
		Psychomotor Retardation Pyrexia	Neurontin(Gabapentin)Tablet	SS		ORAL
ORAL	7	DAY				
		Renal Failure Rhabdomyolysis	Tegretal(Carbamazepi ne)Tablet	SS		ORAL
ORAL	7	DAY				
		Tendonitis Tremor Vision Blurred	Celexa Tablet (Citalopram Hydrobromide) Tablet	SS		ORAL
ORAL	7	DAY				
			Sonata(Zeleplon)Tabl et	SS		ORAL
ORAL	7	DAY				
			Trazodone(Trazodone)			

ORAL	7	DAY	Tablet	SS	ORAL
			Lorcet (Paracetamol, Hydrocodone Bitartrate)	SS	ORAL
ORA	7	DAY	Promethazine (Promethazine) Tablet	C	
			Clindamycin (Clindamycin) Tablet	C	
			Prednisone (Prednisone)	C	
			Premarin Tablet (Estrogens Conjugated)	C	
			Xenical (Orlistat)	C	
			Flurazepam (Flurazepa m)	C	
			Prevacid (Lansoprazol e)	C	

Date: 10/24/03 ISR Number: 4219924-3 Report Type: Expedited (15-DaCompany Report #2003111309
Age: 46 YR Gender: Male I/FU: F

Outcome PT
Other Clumsiness
Drug Effect Decreased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Headache Skin Irritation Tremor Weight Increased	Report Source	Product	Role	Manufacturer	Route
2700 MG, ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Baclofen (Baclofen)	C		
				Amantadine (Amantadine)	C		
				Pantoprazole (Pantoprazole)	C		
				Gemfibrozil (Gemfibrozil)	C		
				Atorvastatin (Atorvastatin)	C		
				Atenolol (Atenolol)	C		
				Zolpidem Tartrate (Zolpidem Tartrate)	C		
				Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
				Oxycocet (Paracetaol, Oxycodone Hyrochloride)	C		
				Axotal (Old Form) (Caffeine, Butalbital, Paracetamol)	C		
				Promethazine Hydrochloride (Promethazine Hydrochloride)	C		
				Paracetamol (Paracetamol)	C		
				Ibuprofen (Ibuprofen)	C		
				Triamcinolone Acetonide (Triamcinolone Acetonide)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG Other (TID), ORAL	Blood Pressure Increased Diverticulitis Gastrointestinal	Consumer	Neurontin (Gabapentin)	PS		ORAL
20 MG (DAILY), ORAL	Haemorrhage Headache Mental Disorder Syncope Vasovagal		Valdecoxib (Valdecoxib)	SS		ORAL
			Tramadol Hydrochloride (Tramadol Hydrochloride) Calcium (Calcium) Calcitonin, Salmon (Cacitonin, Salmon) Multivitamins	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Ergocalciferol,
Ascorbic Acid, Folic
Acid, Thiamine
Hydrochloride, C

Date:10/24/03ISR Number: 4220072-7Report Type:Expedited (15-DaCompany Report #2003112058

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Life Support	Consumer	Neurontin (Gabapentin)	PS		

Date:10/24/03ISR Number: 4220082-XReport Type:Expedited (15-DaCompany Report #2003036352

Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Anxiety Burning Sensation	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Condition Aggravated Depression Diarrhoea Hyperreflexia Hypertension Multiple Sclerosis Muscle Spasms Nausea Oesophageal Spasm Pain Panic Attack Photophobia Stress Tooth Infection Weight Decreased Wheelchair User	Professional	Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Baclofen (Baclofen) Omeprazole (Omeprazole) Glyceryl Trinitrate (Glyceryl Trinitrate) Carbamazepine (Carbamazepine) Obetrol (Dexamfetamine Sulfate, Amfetamine Sulfate, Dexamfetamine	C C C C C		

Date:10/24/03ISR Number: 4220085-5Report Type:Expedited (15-DaCompany Report #2003034184

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Hypersensitivity Neurosis	Consumer Health	Neurontin (Gabapentin)	PS		
THREE TIMES A DAY		Pharmaceutical Product	Professional				
		Complaint Stress		Gabapentin (Gabapentin) All Other Therapeutic Products Antidepressants	SS C C		

Date:10/24/03ISR Number: 4220088-0Report Type:Expedited (15-DaCompany Report #2003111834
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Gastrointestinal Haemorrhage	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/24/03 ISR Number: 4220163-0 Report Type:Expedited (15-DaCompany Report #2003110107
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Glucose Increased Breast Cancer	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other (TID), ORAL		Hyperlipidaemia Renal Impairment		Hydrocodone (Hydrocodone)	C		

Date:10/24/03 ISR Number: 4220277-5 Report Type:Expedited (15-DaCompany Report #KII-2003-0001967
 Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abdominal Pain Abdominal Tenderness Blood Pressure Systolic Decreased Bradycardia	Health Professional	Oxycodone Hydrochloride (Similar To Nda 20-553) (Oxycodone Hydrochloride)	PS		ORAL
ORAL		Depressed Level Of Consciousness		Soma(Carisoprodol)	SS		ORAL
ORAL		Drug Screen Positive		Neurontin(Gabapentin)	SS		ORAL
ORAL		Hyperhidrosis Hypotension Hypothermia Mydriasis Oliguria Rhonchi		Benzodiazepine Derivatives Opioids	SS SS		

Date:10/27/03 ISR Number: 4218969-7 Report Type:Expedited (15-DaCompany Report #WAES 0310FRA00071
 Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Purpura		Singulair Clonazepam	PS SS	Merck & Co., Inc	ORAL

Fluindione SS
Morphine Sulfate SS
Fluticasone
Propionate And
Salmeterol Xinafoate SS

RESPIRATORY

(INHALATION)

Gabapentin SS

Date:10/27/03ISR Number: 4220805-XReport Type:Expedited (15-DaCompany Report #2003112478

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 3600 MG (1200 MG TID)	Hyperkinesia Tremor	Foreign Health Professional	Neurontin (Gabapentin)	PS		
			Clonazepam	C		

Date:10/27/03ISR Number: 4220808-5Report Type:Expedited (15-DaCompany Report #2003112479

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

Outcome	PT
Hospitalization - Initial or Prolonged	Anorexia Disorientation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Somnolence Urinary Incontinence	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Foreign Health Professional	Neurontin (Gabapentin)	PS		
400 MG			Company Representative	Ergenyl Chrono (Valproic Acid)	C		
(DAILY)							

Date:10/27/03ISR Number: 4220958-3Report Type:Expedited (15-DaCompany Report #2003112557
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Disorder	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
Other							
300 MG							
(DAILY), ORAL							

Date:10/27/03ISR Number: 4221032-2Report Type:Expedited (15-DaCompany Report #2003111953
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Balance Disorder					
200 MG		Blood Pressure Increased					
(DAILY), ORAL							
		Depressed Mood		Paracetamol (Paracetamol)	SS		
		Electrocardiogram Abnormal		Fexofenadine Hydrochloride (Fexofenadine Hydrochloride)	C		
		Eye Pain		Rabeprazole Sodium (Rabeprazole Sodium)	C		
		Fatigue		Fluticasone Propionate			
		Increased Appetite					
		Insomnia					
		Intraocular Pressure Increased					
		Logorrhoea					

Stress
Vision Blurred
Visual Disturbance

(Fluticasone
Propionate) C
Centrum Silver
(Ascorbic Acid,
Tocopheryl Acetate,
Retinol, Zinc,
Calcium, Vitamins C
Cimicifuga Racemosa
Root (Cimicifuga
Racemosa Root) C
All Other
Therapeutic Products C
Calcium Carbonate
(Calcium Carbonate) C

Date:10/27/03ISR Number: 4221042-5Report Type:Expedited (15-DaCompany Report #2003112062
Age:70 YR Gender:Female I/FU:I

Outcome PT
Hospitalization - Acute Myocardial
Initial or Prolonged Infarction
Other Asthenia
Blood Pressure
Inadequately Controlled

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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
ORAL		Drug Ineffective Feeling Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Groin Pain Nasopharyngitis					
		Pain In Extremity		Antihypertensives Atorvastatin (Atorvastatin)	C C		

Date:10/27/03ISR Number: 4221197-2Report Type:Expedited (15-DaCompany Report #KII-2003-0002577
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Alanine Aminotransferase Increased	Health Professional	Oxycodone Hydrochloride (Similar To Nda 20-553) (Oxycodne Hydrochloride)	PS		ORAL
SEE TEXT,		Convulsion					
ORAL		Hypertension					
SEE TEXT,		Hypotension Multiple Drug Overdose		Neurontin (Gabapentin)	SS		ORAL
ORAL		Phantom Pain					
SEE TEXT,		Sinus Tachycardia Somnolence		Skelaxin (Metaxalone)	SS		ORAL
ORAL		Urine Output Decreased					
		Vomiting		Acetaminophen (Paracetamol) Capsule	SS		

Date:10/27/03ISR Number: 4221277-1Report Type:Expedited (15-DaCompany Report #KII-2003-0002343
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization - Initial or Prolonged Other	Blood Chloride Blood Sodium Blood Urea Hypothermia Loss Of Consciousness	Health Professional	Oxycodone Hydrochloride (Similar To Nda 20-553)(Oxycodone Hydrochloride)	PS	ORAL
ORAL					
	Multiple Drug Overdose Mydriasis Myoglobin Blood Pupil Fixed Toxicologic Test Abnormal		Morphine Sulfate (Similar To Nda 19-516) (Morphine Sulfate) Opioids() Valium(Diazepam)	SS SS SS	ORAL
ORAL					
			Percocet (Paracetamol , Oxycodone Hydrochloride)	SS	ORAL
ORAL					
			Barbiturates() Gabapentin(Gabapenti n)	SS SS	ORAL
ORAL					
			Hydrochlorothiazide(Hydrochlorothiazide)	SS	ORAL
ORAL					
			Benzodiazepine Derivatives() Acetaminophen (Paracetamol) Acetylsalicylic Acid (Acetylsalicylic Acid)	SS SS SS	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/28/03 ISR Number: 4220473-7 Report Type:Direct
 Age:17 YR Gender:Male I/FU:I

Company Report #CTU 204699

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Suicidal Ideation		Neurontin 150			
Other				Parke-Davis	PS	Parke-Davis	
150 2XDAY							

Date:10/28/03 ISR Number: 4222735-6 Report Type:Expedited (15-Da
 Age:46 YR Gender:Female I/FU:F Company Report #001-0945-M0200657

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Angioneurotic Oedema	Consumer	Neurontin			
Other		Antinuclear Antibody	Health	(Gabapentin)	PS		ORAL
1800 MG							
(DAILY), ORAL		Positive	Professional				
		Biopsy Tongue Abnormal		Lamotrigine			
		Blood Albumin Decreased		(Lamotrigine)	C		
		Blood Glucose Increased		Lithium Carbonate			
		Brain Damage		(Lithium Carbonate)	C		
		Coordination Abnormal		Clonazepam			
		Crying		(Clonazepam)	C		
		Depression		Methylphenidate			
		Diplopia		Hydrochloride			
		Drug Hypersensitivity		(Methylphenidate)	C		
		Dysarthria		Levothyroxine Sodium			
		Dysphagia		(Levothyroxine			
		Foreign Body Trauma		Sodium)	C		
		Gingival Pain		Liothyronine Sodium			
		Glossodynia		(Liothyronine			
		Malnutrition		Sodium)	C		
		Mania		Sertraline			
		Medication Error		Hydrochloride			
		Nervous System Disorder		(Sertraline			
		Nystagmus		Hydrochloride)	C		
		Oral Pain		Pilocarpine			
		Overdose		Hydrochloride			
		Sensory Disturbance		(Pilocarpine			
		Speech Disorder		Hydrochloride)	C		
		Stress		Metoprolol Succinate			
		Suicidal Ideation		(Metoprolol			
		Swollen Tongue		Succinate)	C		

Tardive Dyskinesia
Tongue Disorder
Tooth Disorder

Lansoprazole
(Lansoprazole) C
Hyoscyamine Sulfate
(Hyoscyamine
Sulfate) C
Diltiazem
Hydrochloride
(Diltiazem
Hydrochloride) C
Lamotrigine
(Lamotrigine) C

Date:10/28/03ISR Number: 4222771-XReport Type:Expedited (15-DaCompany Report #USA-2003-0007807

Age:55 YR Gender:Female I/FU:F

Outcome PT
Death Analgesic Drug Level
Above Therapeutic
Drug Level Above
Therapeutic

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Myocardial Ischaemia Toxicologic Test Abnormal	Report Source	Product	Role	Manufacturer	Route
			Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)(Oxycodone Hydrochloride)	PS		
				Acetaminophen(Parace tamol)	SS		
				Alprazolam(Alprazola m)	SS		
				Amitriptyline(Amitri ptyline)	SS		
				Codeine(Codeine)	SS		
				Gabapentin(Gabapenti n)	SS		
				Tramadol(Tramadol)	SS		
				Ibuprofen(Ibuprofen)	SS		
				Lorazepam(Lorazepam)	SS		

Date:10/29/03ISR Number: 4223111-2Report Type:Expedited (15-DaCompany Report #USA-2003-0007441
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death		Atherosclerosis Bacterial Infection Bronchopneumonia Coma Drug Screen Positive Lung Abscess Pulmonary Congestion Pulmonary Embolism Pulmonary Oedema Sepsis	Health Professional Other	Oxycodone Hydrochloride (Similar Nda 20-553) (Oxycodone Hydrochloride)	PS		
				Hydrocodone Bitartrate (Similar To Ind 59,175) (Naltrexone, Paracetamol, Alprazolam)	SS		
				(Alprazolam) Citalopram (Citalopram)	SS		
				Procainamide (Procainamide) Acetaminophen (Paracetamol)	SS		

Buspirone	
(Buspirone)	SS
Diltiazem	
(Diltiazem)	SS
Gabapentin	
(Gabapentin)	SS
Trazodone	
(Trazodone)	SS
Zolpidem (Zolpidem)	SS

Date:10/29/03ISR Number: 4223127-6Report Type:Expedited (15-DaCompany Report #USA-2003-0007385
Age:50 YR Gender:Female I/FU:F

Outcome PT
Death Coronary Artery
Atherosclerosis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Abuser Drug Screen Positive Pulmonary Congestion					
		Pulmonary Oedema	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) (Oxycodone Hydrochloride)	PS		
				Diazepam (Diazepam)	SS		
				Gabapentin (Gabapentin)	SS		
				Oxazepam (Oxazepam)	SS		
				Temazepam (Temazepam)	SS		

Date:10/29/03ISR Number: 4223130-6Report Type:Expedited (15-DaCompany Report #USA-2003-0010426
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death		Carcinoid Tumour Of The Appendix Coma Coronary Artery Atherosclerosis Drug Level Increased Drug Screen Positive Genital Disorder Female Hepatic Congestion Hepatic Necrosis Myocardial Ischaemia Pulmonary Congestion Snoring	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) (Oxycodone Hydrochloride)	PS		
				Acetaminophen (Paracetamol)	SS		
				Alparzolam (Alprazolam)	SS		
				Gabapentin (Gabapentin)	SS		
				Fluoxetine (Fluoxetine)	SS		
				Methadone (Methadone)	SS		
				Oxazepam (Oxazepam)	SS		
				Trazodone (Trazodone)	SS		
				Lorazepam (Lorazepam)	SS		

Outcome	PT
Life-Threatening	Agranulocytosis
Hospitalization -	Blood Lactate
Initial or Prolonged	Dehydrogenase Increased
Other	Body Temperature Increased Electrocardiogram St-T Segment Elevation Erythema Infectiosum Impaired Work Ability Leukopenia Myocarditis Parvovirus B19 Serology Positive Pericardial Effusion Splenomegaly

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Staphylococcal Infection
Troponin Increased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
900 MG (TID), ORAL		Foreign Consumer Health Professional Other	Neurontin (Gabapentin)	PS		ORAL
			Cefotaxime Sodium (Cefotaxime Sodium)	C		

Date:10/29/03
Age: ISR Number: 4223240-3
Gender:Male I/FU:I Report Type:Expedited (15-DaCompany Report #2003112724

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 400 MG (DAILY), ORAL		Cardiovascular Disorder Respiratory Arrest	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:10/29/03
Age:1 DY ISR Number: 4223480-3
Gender:Female I/FU:F Report Type:Expedited (15-DaCompany Report #2003037078

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Disability 1600 MG Congenital Anomaly (BID), ORAL Other		Anorectal Disorder Cardiac Murmur Congenital Tracheomalacia Feeding Problem In Newborn Maternal Drugs Affecting Foetus Neonatal Disorder Respiratory Arrest Respiratory Disorder	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Complex Regional Pain Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (DAILY), ORAL		Condition Aggravated					
		Fatigue Headache		Zyrtec (Tablets) (Cetirizine)	SS		ORAL
10 MG (DAILY), ORAL		Musculoskeletal Stiffness					
		Neuralgia Pyrexia		Celecoxib (Celecoxib)	SS		ORAL
200 MG (DAILY), ORAL		Urticaria					
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochlorid,	C		
				Calcium (Calcium)	C		
				Hydrocodone (Hydrocodone)	C		
				Diazepam (Diazepam)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/30/03ISR Number: 4222197-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0312562A
Age:36 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - RESPIRATORY Initial or Prolonged (INHALATION)	Activated Partial Thromboplastin Time Prolonged Infection International Normalised Ratio Increased Purpura		Seretide Rivotril Skenan Singulair Neurontin Previscan	PS SS SS SS C	Glaxosmithkline Glaxosmithkline	 ORAL ORAL ORAL ORAL ORAL

Date:10/30/03ISR Number: 4223220-8Report Type:Direct Company Report #CTU 204953
Age:71 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose 300 MG PO HS	Dyskinesia Fall Mental Status Changes Tremor		Neurontin Zocor Fosamax Aspirin Serzone Atenolol Oxycontin	PS C C C C C		ORAL

Date:10/30/03ISR Number: 4224155-7Report Type:Expedited (15-DaCompany Report #KII-2003-0003995
Age:41 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Acidosis Coma Depressed Level Of Consciousness Drug Screen Positive Drug Withdrawal Syndrome Haemoptysis Multiple Drug Overdose Pco2	Health Professional	Oxycontin Cr Hydrocodone Bitartrate (Similar To Id 59,175) (Hydrocodone Bitartrate, Neurontin (Gabapentin) Celexa (Citalopram	PS SS SS		

Po2
Tachycardia
Vomiting

Hydrobromide) SS
Soma
(Carisoprodol) SS
Tricyclic
Antidepressant SS
Acetaminophen
(Paracetamol) SS
Marijuana (Cannabis) SS
Benzodiazepine
Derivatives () SS
Alcohol (Ethanol) SS

Date:10/30/03ISR Number: 4224263-0Report Type:Expedited (15-DaCompany Report #KII-2003-0004101
Age:57 YR Gender:Female I/FU:I

Outcome PT
Hospitalization - Alanine Aminotransferase
Initial or Prolonged Aspartate
Other Aminotransferase
Blood Creatinine
Blood Glucose
Blood Potassium

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
	Blood Pressure Increased Blood Urea Coma					
	Depressed Level Of Consciousness Drug Screen Positive Electrocardiogram Qrs	Health Professional	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		
SEE TEXT,	Complex Prolonged Feeling Abnormal International Normalised Ratio Increased Moaning Myocardial Infarction Nasopharyngitis Nodal Rhythm Pco2 Prothrombin Time Respiratory Rate Increased Ventricular Extrasystoles		Potassium (Potassium) Ibuprofen (Ibuprofen) Prozac (Fluoxetine Hydrochloride) Hydrochlorothiazide (Hydrochlorothiazide) Prevacid (Lansoprazole) Estratest (Estrogens Esterified, Methyltestosterone) Norvasc (Amlodipine Besilate) Midrin (Isometheptene, Dichloralphenazone) Clonazepam (Clonazepam) Soma (Carisoprodol) Amitriptyline (Amitriptyline) Neurontin (Gabapentin) Tramadol (Tramadol)	SS SS SS SS SS SS SS SS SS SS		

Date:10/30/03ISR Number: 4224411-2Report Type:Expedited (15-DaCompany Report #2003113340

Age:73 YR Gender:Male I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death Other 1600 MG	Completed Suicide Disinhibition	Foreign Health	Neurontin (Gabapentin)	PS		

(DAILY)

Drug Effect Decreased

Professional

Pain

Company
Representative

Dyazide
(Hydrochlorothiazide
, Triamterene) C
Simvastatin
(Simvastatin) C
Metformin
(Metformin) C
Antidepressants C

Date:10/31/03ISR Number: 4223931-4Report Type:Direct
Age:34 YR Gender:Male I/FU:I

Company Report #CTU 204974

Outcome PT
Death Completed Suicide
Confusional State
Disorientation
Feeling 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
1 PO 300 MG 1		Gun Shot Wound Hallucination Head Injury		Neurontin 300 Mg Parke-Davis	PS	Parke-Davis	ORAL
Q6H ORAL							

Date:10/31/03ISR Number: 4224063-1Report Type:Expedited (15-DaCompany Report #2003113331
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blindness Quadriplegia	Consumer	Neurontin (Gabapentin)	PS		

Date:10/31/03ISR Number: 4224688-3Report Type:Direct Company Report #CTU 204972
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Upper Anxiety		Neurontin (Gabapentin)	PS		ORAL
300 MG PO		Condition Aggravated					
TID, 100 MG		Depression					
PO TID		Diarrhoea		Baclofen	C		
		Multiple Sclerosis		Fluoxetine	C		
		Muscle Spasms		Lovastatin	C		
		Palpitations		Maxzide	C		
		Paraesthesia		Nortriptyline	C		
		Stomatitis		Adderal	C		
		Stress					
		Weight Decreased					

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
	Heart Rate Increased
	Miosis
	Multiple Drug Overdose
	Muscle Twitching
	Mydriasis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Posturing Pupil Fixed Respiratory Rate	Report Source	Product	Role	Manufacturer	Route
		Decreased Respiratory Rate Increased	Health Professional Other	Oxycontin Cr Methadone (Methadone) Trazodone(Trazodone) Gabapentin(Gabapenti n)	PS SS SS SS		
400 MG				Quetiapine(Quetiapin e)	SS		
200 MG				Citalopram(Citalopra m)	SS		
20 MG				Clonazepam (Clonazepam)	SS		
1 MG				Zanaflex(Tizanidine Hydrochloride)	SS		ORAL
4 MG, ORAL				Bupropion(Amfebutamo ne)	SS		
150 MG				Cyclobenzaprine(Cycl obenzaprine)	SS		
10 MG				Amitriptyline(Amitri ptyline)	SS		
50 MG				Estalopram()	SS		

Date:10/31/03ISR Number: 4224892-4Report Type:Expedited (15-DaCompany Report #2003036045
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other (BID), ORAL		Burning Sensation Cardiac Disorder Coronary Artery Occlusion Epistaxis Hypoaesthesia	Consumer Health Professional	Neurontin (Gabapentin) Carbamazepine (Carbamazepine) All Other	PS C		ORAL

Insomnia

Therapeutic Products C

Date:10/31/03ISR Number: 4224894-8Report Type:Expedited (15-DaCompany Report #2003113619
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Neoplasm Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
5200 MG							
(DAILY), ORAL							

Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C
Celecoxib (Celecoxib)	C
Lansoprazole (Lansoprazole)	C

Date:10/31/03ISR Number: 4224925-5Report Type:Expedited (15-DaCompany Report #2003113597
Age:11 YR Gender:Female I/FU:I

Outcome	PT
Other	Anger Homicidal Ideation

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Freedom Of Information (FOI) Report

Suicidal Ideation

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
600 MG (BID), ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:11/03/03ISR Number: 4225981-0Report Type:Expedited (15-DaCompany Report #2003113616
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 400 MG (BID), ORAL		Abnormal Behaviour Anxiety Back Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Condition Aggravated Decreased Appetite Feeling Abnormal Gastrointestinal Haemorrhage Gastrointestinal Motility Disorder Insomnia Irritability Limb Injury Somnolence Tremor		Mirtazapine (Mirtazapine)	SS		

Date:11/03/03ISR Number: 4226014-2Report Type:Expedited (15-DaCompany Report #2003113595
Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (TID), ORAL		Chest Pain Condition Aggravated Post Procedural Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL

Sciatica
Stress

Amlodipine Besilate
(Amlodipine
Besilate) C
Metformin
Hydrochloride
(Metformin
Hydrochloride) C
Pioglitazone
(Pioglitazone) C
Esomeprazole
(Esomeprazole) C
Fluoxetine
Hydrochloride
(Fluoxetine
Hydrochloride) C
Lorazepam
(Lorazepam) C

Date:11/03/03ISR Number: 4226024-5Report Type:Expedited (15-DaCompany Report #2003113461
Age:39 YR Gender:Male I/FU:I

Outcome PT
Other Anxiety
Drug Effect Decreased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (TID), ORAL		Eye Rolling Muscle Twitching Nervousness Sleep Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Salbutamol (Salbutamol)	C		
				Triamcinolone Acetonide (Triamcinolone Acetonide)	C		
				Peginterferon Alfa-2b (Peginterferon Alfa-2b)	C		
				Ribavirin (Ribavirin)	C		
				Fentanyl (Fentanyl)	C		
				Lisinopril (Lisinopril)	C		
				Atenolol (Atenolol)	C		
				Rabeprazole Sodium (Rabeprazole Sodium)	C		
				Metoclopramide (Metoclopramide)	C		
				Ondansetron Hydrochloride (Ondansetron Hydrochloride)	C		
				Lidocaine (Lidocaine)	C		

Date:11/03/03ISR Number: 4226026-9Report Type:Expedited (15-DaCompany Report #2003113041
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2700 MG		Apathy Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL

(TID), ORAL	Drug Ineffective			
100 MG	Impaired Work Ability	Zoloft (Sertraline)	SS	ORAL
(DAILY), ORAL	Marital Problem			
	Suicidal Ideation	Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS	
		Hydrocodone (Hydrocodone)	C	
		Zolpidem Tartrate (Zolpidem Tartrate)	C	
		Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride)	C	
		Baclofen (Baclofen)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/03ISR Number: 4226198-6Report Type:Expedited (15-DaCompany Report #2003032942

Age:24 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged OPHTHALMIC 900 MG ORAL	Bipolar Disorder Fatigue (TID), Somnolence Treatment Noncompliance Weight Increased	Consumer Health Professional	Neurontin (Gabapentin)	PS		
			Methylphenidate Hydrochloride	C		
			Valproate Semisodium	C		
			Fluoxetine			
			Hydrochloride	C		
			Rantidine			
			Hydrochloride	C		

Date:11/03/03ISR Number: 4226555-8Report Type:Expedited (15-DaCompany Report #2003113622

Age:19 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 1200 MG Initial or Prolonged (TID), ORAL Disability ORAL	Cholestasis Coagulopathy Dermatitis Exfoliative Diabetic Neuropathy Eosinophilia Erythema Haematoma Hepatocellular Damage Hypersensitivity Hyponatraemia Leukocytosis Liver Function Test Abnormal Pain Petechiae Platelet Count Increased Pyrexia Rash Maculo-Papular Sinus Tachycardia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Carbamazepine (Carbamazepine)	SS		ORAL
			Insulin Human (Insulin Human)	C		
			Insulin Lispro (Insulin Lispro)	C		
			Paracetamol			
			(Paracetamol)	C		

Date:11/03/03ISR Number: 4226576-5Report Type:Expedited (15-DaCompany Report #2003113340
 Age:73 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Foreign	Neurontin			
Other		Condition Aggravated	Health	(Gabapentin)	PS		
1600 MG		Death	Professional				
(DAILY)		Pain	Company	Dyazide			
			Representative	(Hydrochlorothiazide			
				, Triamterene)	C		
				Simvastatin			
				(Simvastatin)	C		
				Metformin			
				(Metformin)	C		
				Antidepressants	C		
				Oxcarbazepine			
				(Oxcarbazepine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/03ISR Number: 4226871-XReport Type:Expedited (15-DaCompany Report #2003-DE-05108GD

Age:50 YR Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Cardio-Respiratory Arrest Completed Suicide	Literature	Methadone (Methadone)	PS		ORAL
PO			Multiple Drug Overdose		Olanzapine	SS		ORAL
PO					Gabapentin (Gabapentin)	SS		ORAL

Date:11/04/03ISR Number: 4227354-3Report Type:Expedited (15-DaCompany Report #2003113768

Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Cachexia Somnolence	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Morphine (Morphine)	PS C		

Date:11/04/03ISR Number: 4227356-7Report Type:Expedited (15-DaCompany Report #2003036800

Age:61 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (DAILY), ORAL			Carpal Tunnel Decompression Oedema Peripheral Post Procedural Complication	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL

Date:11/05/03ISR Number: 4228488-XReport Type:Expedited (15-DaCompany Report #2003114194

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1500 MG (5 Other DOSES PER DAY), ORAL	Somnolence Thirst	Consumer	Neurontin (Gabapentin)	PS		ORAL

Zoloft (Sertraline)	C
Zyrtec (Cetirizine)	C
All Other Therapeutic Products	C
Levothyroxine Sodium (Levothyroxine Sodium)	C
Rabeprazole Sodium (Rabeprazole Sodium)	C
Nadolol (Nadolol)	C
Lactulose (Lactulose)	C
Becosym Forte (Pyridoxine Hydrochloride, Thiamine Hydrochloride,	C
Folic Acid (Folic Acid)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/05/03ISR Number: 4228492-1Report Type:Expedited (15-DaCompany Report #2003114316
Age:67 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 200 MG (BID), Other ORAL	Hepatic Haematoma Hypotension Intra-Abdominal Haemorrhage Liver Function Test Abnormal Post Procedural Complication	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Alprazolam (Alprazolam)	C		
			Phenelzine Sulfate (Phenelzine Sulfate)	C		
			Olanzapine (Olanzapine)	C		
			Estradiol (Estradiol)	C		
			Medroxyprogesterone Acetate (Medroxyprogesterone Acetate)	C		
			Estrogens Conjugated (Estrogens Conjugated)	C		
			Fluticasone PrOpionate (Fluticasone Propionate)	C		
			Guaifenesin (Guaifenesin)	C		
			Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)	C		
			Ipratropium Bromide (Ipratropium Bromide)	C		
			Nystatin (Nystatin)	C		
			Macrogol (Macrogol)	C		

Date:11/05/03ISR Number: 4228561-6Report Type:Expedited (15-DaCompany Report #2003111834
Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Gastrointestinal Haemorrhage	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:11/05/03ISR Number: 4228563-XReport Type:Expedited (15-DaCompany Report #2003114191
Age:55 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Anxiety
Initial or Prolonged	Asthenia
Other	Brain Abscess
	Crying
	Dizziness Postural
	Drug Ineffective
	Fear
	Headache

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Somnolence Speech Disorder	Report Source	Product	Role	Manufacturer	Route
2400 MG			Consumer	Neurontin (Gabapentin)	PS		ORAL
(TID), ORAL				Escitalopram (Escitalopram)	C		
				Methadone (Methadone)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Omeprazole (Omeprazole)	C		
				Nadolol (Nadolol)	C		
				Verapamil Hydrochloride (Verapamil Hydrochloride)	C		
				Hydrochlorothiazide (Hydrochlorothiazide)	C		

Date:11/05/03ISR Number: 4228564-1Report Type:Expedited (15-DaCompany Report #2003034004
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Agitation	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Body Temperature	Health				
(TID), ORAL		Increased Difficulty In Walking	Professional	Triazolam (Triazolam)	SS		
0.25 MG		Drug Ineffective					
(DAILY),		Dyspepsia		Oxycocet (Paracetamol, Oxycodone Hydrochloride)	C		
		Feeling Cold		Oxycodone			
		Insomnia					
		Nausea					
		Tremor					

(Oxycodone) C
 Ibuprofen C
 (Ibuprofen) C
 Diazepam (Diazepam) C
 Colchicine C
 (Colchicine) C
 Allopurinol C
 (Allopurinol) C

Date:11/05/03ISR Number: 4228566-5Report Type:Expedited (15-DaCompany Report #2003114674
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Filariasis	Consumer	Neurontin			
		Lymphoedema		(Gabapentin)	PS		ORAL
900 MG (TID),		Oedema Peripheral					
ORAL				Tamoxifen			
				(Tamoxifen)	C		
				Esomeprazole			
				(Esomeprazole)	C		
				Losartan Potassium			

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Freedom Of Information (FOI) Report

(Losartan Potassium) C
 Amitriptyline
 Hydrochloride
 (Amitriptyline
 Hydrochloride) C

Date:11/05/03ISR Number: 4228838-4Report Type:Expedited (15-DaCompany Report #2003114127
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Anorexia Gastritis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Hyponatraemia Oral Fungal Infection Psychological Factor Affecting Medical Condition Weight Decreased	Professional	Levothyroxine Sodium (Levothyroxine Sodium)	C		

Date:11/05/03ISR Number: 4228840-2Report Type:Expedited (15-DaCompany Report #2003040746
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Loss Of Consciousness Overdose	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Pulmonary Function Test Decreased Skin Laceration	Professional Company Representative	Ethanol (Ethanol)	C		

Date:11/05/03ISR Number: 4229002-5Report Type:Expedited (15-DaCompany Report #2003114317
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 100 MG		Asthenia Difficulty In Walking	Consumer	Neurontin (Gabapentin)	PS		ORAL

(DAILY), ORAL

Muscle Spasms

Diazepam (Diazepam) C
Warfarin Sodium
(Warfarin Sodium) C

Date:11/06/03ISR Number: 4228239-9Report Type:Direct
Age:77 YR Gender:Male I/FU:I

Company Report #CTU 205406

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG PO TID		Chills Myoclonus Pruritus Pyrexia Tremor		Gabapentin (Neurontin)	PS		ORAL
				Cyclosporin (Neoral)	C		
				Doxazosin	C		
				Folic Acid	C		
				Sodium Bicarbonate	C		
				Prednisone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/06/03ISR Number: 4229825-2Report Type:Expedited (15-DaCompany Report #2003111978
Age:85 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Autoimmune Disorder	Foreign	Neurontin			
Hospitalization - ORAL		Blood Electrolytes	Health	(Gabapentin)	PS		ORAL
Initial or Prolonged		Abnormal Dehydration Hypotension Laboratory Test Abnormal Mood Disorder Due To A General Medical Condition Pulmonary Embolism Pyrexia Renal Failure Shock	Professional	Pramipexole Dihydrochloride (Pramipexole Dihydrochloride) Levodopa (Levodopa)	C C		

Date:11/06/03ISR Number: 4230681-7Report Type:Expedited (15-DaCompany Report #2003114675
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (DAILY), ORAL		Balance Disorder Chest Discomfort Chest Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Diarrhoea Dizziness Dyspepsia Dyspnoea		Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride)	SS		ORAL
		Fatigue Feeling Abnormal Increased Appetite Skin Disorder Somnolence Speech Disorder Vision Blurred		Paroxetine Hydrochloride (Paroxetine Hydrochloride) Methylphenidate Hydrochloride (Methylphenidate Hydrochloride) Zolpidem Tartrate (Zolpidem Tartrate) Meloxicam	C C C		

(Meloxicam)

C

Date:11/07/03ISR Number: 4230497-1Report Type:Expedited (15-DaCompany Report #WAES 0310FRA00071
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Purpura	Health Professional	Singulair Clonazepam Fluindione Morphine Sulfate Fluticasone Propionate And Salmeterol Xinafoate	PS SS SS SS	Merck & Co., Inc	ORAL
RESPIRATORY							
(INHALATION)							
				Gabapentin	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4231612-6Report Type:Expedited (15-DaCompany Report #2003115039
 Age:42 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG Other (DAILY), ORAL	Dyskinesia Mental Disorder Myoclonus	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
			Rofecoxib (Rofecoxib)	C		
			Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		
			Tramadol (Tramadol)	C		

Date:11/07/03ISR Number: 4231764-8Report Type:Expedited (15-DaCompany Report #2003UW14116
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent 50 MG DAILY Impairment/Damage PO 0.3	Alopecia Angina Pectoris Arrhythmia Blood Pressure Increased Cardiac Disorder Cardiac Failure Cardiac Valve Disease Chest Pain Condition Aggravated Disturbance In Attention Dizziness Electrolyte Imbalance Heart Rate Increased Hypotension Memory Impairment Myocardial Infarction Pulmonary Hypertension Restlessness Somnolence	Foreign Health Professional Other	Xylocaine Baclofen Neurontin Nitrostat Atenolol Avapro K-Lyte No Match Oxygen Taurine	PS SS SS SS C C C C C C		ORAL

Weight Decreased
Weight Increased

Date:11/07/03ISR Number: 4231771-5Report Type:Periodic
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0417864A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness		Wellbutrin	PS	Glaxosmithkline	ORAL
100MG Per day	WK	Vertigo		Neurontin	SS		
UNKNOWN				Tagamet	C	Glaxosmithkline	

Date:11/07/03ISR Number: 4232052-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0285837A
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Confusional State Encephalopathy Hypokalaemia		Abacavir Sulfate + Lamivudine + Zidovudine	PS	Glaxosmithkline	ORAL
1UNIT Twice per day		Hypothermia					
UNKNOWN		Respiratory Alkalosis		Gabapentin	SS		
UNKNOWN		Somnolence		Nicardipine	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/07/03ISR Number: 4232273-2Report Type:Expedited (15-DaCompany Report #2003115043

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cartilage Injury	Consumer	Neurontin			
3600 MG		Joint Effusion		(Gabapentin)	PS		ORAL
(TID), ORAL		Joint Sprain					
		Joint Swelling		Metoprolol Succinate			
				(Metoprolol Succinate)	C		
				Cibadrex			
				"Ciba-Geigy"			
				(Hydrochlorothiazide			
				, Benazepril			
				Hydrochloride)	C		
				Celecoxib			
				(Celecoxib)Q	C		
				Ascorbic Acid			
				(Ascorbic Acid)	C		
				Calcium D3 "Stada"			
				(Colecalciferol,			
				Calcium)	C		
				Estrogens			
				Conjugated(Estrogens			
				Conjugated)	C		
				Acetylsalicylic Acid			
				(Acetylsalicylic			
				Acid)	C		
				Sertraline			
				Hydrochloride			
				(Sertraline			
				Hydrochloride)	C		
				Cyclobenzaprine			
				Hydrochloride			
				(Cyclobenzaprine			
				Hydrochloride)	C		
				Furosemide			
				(Furosemide)	C		
				Clonazepam			
				(Clonazepam)	C		
				Esomeprazole			
				(Esomeprazole)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Colostomy	Consumer	Neurontin (Solution)			
Initial or Prolonged	Drug Effect Decreased		(Gabapentin)	PS		ORAL
ORAL						
Other	Hip Fracture		Methylphenidate			
	Pharmaceutical Product		Hydrochloride			
	Complaint		(Methylphenidate			
	Somnolence		Hydrochloride)	C		
			Famotidine			
			(Famotidine)	C		
			Heparin-Fraction,			
			Sodium Salt			
			(Heparin-Fraction,			
			Sodium Salt)	C		
			All Other			
			Therapeutic Products	C		

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Freedom Of Information (FOI) Report

Paracetamol
(Paracetamol) C

Date:11/10/03ISR Number: 4233763-9Report Type:Expedited (15-DaCompany Report #2003115340
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Emotional Disorder Mental Disorder	Consumer	Neurontin (Gabapentin)	PS		

Date:11/10/03ISR Number: 4233786-XReport Type:Expedited (15-DaCompany Report #2003114753
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		
				Omeprazole (Omeprazole)	C		
				Clonazepam (Clonazepam)	C		

Date:11/10/03ISR Number: 4234291-7Report Type:Expedited (15-DaCompany Report #2003115045
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 600 MG (TID), Other ORAL		Cataract Drug Ineffective Impaired Healing Neuropathic Pain Retinal Vascular Disorder Swelling	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Insulin Lispro (Insulin Lispro)	C		
				Inslulin Glargine (Insulin Glargine)	C		
				Metformin Hydrochloride (Metformin Hydrochloride)	C		

Glibenclamide	C
(Glibenclamide)	
Metolazone	
(Metolazone)	C
Simvastatin	
(Simvastatin)	C
Oxycocet	
(Paracetamol,	
Oxycodone	
Hydrochloride)	C
Citalopram	
Hydrobromide	
(Citalopram	
Hydrobromide)	C
Lisinopril	
(Lisinopril)	C
Alprazolam	
(Alprazolam)	C
Potassium Chloride	
(Potassium Chloride)	C
Fenofibrate	
(Fenofibrate)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/10/03ISR Number: 4234336-4Report Type:Expedited (15-DaCompany Report #2003115707
Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (DAILY); ORAL		Dyskinesia Dystonia	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
				Acetylsalicylic Acid	C		
				Lisinopril	C		
				Isosorbide Mononitrate	C		
				Amiodarone	C		
				Furosemide	C		
				Levothyroxine	C		

Date:11/10/03ISR Number: 4234337-6Report Type:Expedited (15-DaCompany Report #2003115344
Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 400 MG (BID); ORAL		Activities Of Daily Living Impaired Condition Aggravated	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Depression		Digoxin	C		
		Dyskinesia		Warfarin Sodium	C		
		Dysphasia		Candesartan			
		Tremor		Cilexetil	C		
				Hydrochlorothiazide	C		
				Ferrous Sulfate	C		
				Allopurinol	C		
				Folic Acid	C		
				Initard (Insulin, Insulin Injection, Isophane)	C		
				Fentanyl	C		

Date:11/10/03ISR Number: 4234381-9Report Type:Expedited (15-DaCompany Report #2003115568
Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (TID) Other		Blood Pressure Decreased Blood Pressure Fluctuation Dialysis Hypertension Renal Failure	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:11/10/03ISR Number: 4234390-XReport Type:Expedited (15-DaCompany Report #2003008068
Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability Other	Aplastic Anaemia Arthralgia Asthenia Coma Convulsion Disturbance In Attention Dizziness Dry Mouth 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
		Gait Disturbance					
		Headache					
		Heart Rate Increased	Report Source				
		Mental Impairment	Consumer	Neurontin			
		Muscle Spasms		(Gabapentin)	PS		
		Nerve Injury		Dilantin Suspension			
		Nervousness		(Phenytoin Sodium)	SS		
		Pain		Acetylsalicylic Acid			
		Pain In Extremity		(Acetylsalicylic			
		Panic Attack		Acid)	SS		
		Paralysis		Vitamins With			
		Speech Disorder		Minerals	SS		
		Thermal Burn					

Date:11/10/03ISR Number: 4234420-5Report Type:Expedited (15-DaCompany Report #2003115415
 Age:86 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300 MG		Respiratory Arrest	Foreign Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
								(DAILY), ORAL

Date:11/10/03ISR Number: 4234444-8Report Type:Expedited (15-DaCompany Report #2003114957
 Age:40 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Angioneurotic Oedema Food Allergy	Foreign Health Professional	Amlodipine (Amlodipine) Neurontin (Gabapentin) Bisoprolol (Bisoprolol)	PS SS SS		

Date:11/11/03ISR Number: 4233003-0Report Type:Direct Company Report #CTU 205795
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pancytopenia		Gabapentin Neurontin	PS C		

Date:11/12/03ISR Number: 4233984-5Report Type:Direct Company Report #CTU 205823
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Disability		Abdominal Pain Anger		Neurontin 400 Mg Pfizer	PS	Pfizer	ORAL
Other		Cold Sweat					
TIMES OARL		Condition Aggravated Diarrhoea Drug Dependence Drug Withdrawal Syndrome Hostility Influenza Like Illness Irritable Bowel Syndrome Piloerection Urticaria					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/12/03ISR Number: 4234866-5Report Type:Expedited (15-DaCompany Report #2003115700

Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chills	Consumer	Neurontin			
1200 MG		Cold Sweat		(Gabapentin)	PS		ORAL
(DAILY), ORAL		Drug Withdrawal Syndrome					
		Feeling Abnormal		Vicodin			
		Feeling Hot		(Paracetamol,			
		Medication Error		Hydrocodone			
		Nausea		Bitartrate)	SS		
		Pain		All Other			
		Pyrexia		Therapeutic Products	C		
		Vomiting					

Date:11/12/03ISR Number: 4234942-7Report Type:Expedited (15-DaCompany Report #HQWYE542027OCT03

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Haemoglobin	Study	Serax (Oxazepam,			
Initial or Prolonged		Intentional Misuse		Unspec)	PS		ORAL
"EXCESSIVE		Suicide Attempt					
INGESTION"							
(DOSE							
UNKNOWN) ORAL				Ethanol (Ethanol,)	SS		ORAL
ORAL				Lexapro			
10 MG 1 X PER				(Escitalopram,)	SS		ORAL
1 DAY ORAL							
20 MG 1X PER				Lexapro			
1 DAY				(Escitalopram,)	SS		
				Neurontin			

300 MG 3X PER

(Gabapentin,) SS

1 DAY

Neurontin
(Gabapentin,) SS

'EXCESSIVE

INGESTION"

(DOSE

UNKNOWN)

Corgard (Nadolol) C

Date:11/12/03ISR Number: 4234983-XReport Type:Expedited (15-DaCompany Report #2003115873

Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Alanine Aminotransferase	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
Other		Increased	Professional				
600 MG BID		Aspartate					
ORAL		Aminotransferase	Other	Immunoglobulin Human Normal	SS		
INTRA VENOUS	30 GRAM	Increased					
(DAILY),		Gamma-Glutamyltransferase					
INTRA VENOUS		Increased					
		Hepatocellular Damage		Paracetamol	C		
				Tramadol	C		
				Diazepam	C		
				Heparin-Fraction, Sodium Salt	C		
				Nulytely (Sodium Bicarbonate, Potassium Cl, Sodium			

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Freedom Of Information (FOI) Report

Cl Macrogol) C
 Nystatin C
 Morphine Sulfate C

Date:11/12/03ISR Number: 4234988-9Report Type:Expedited (15-DaCompany Report #2003115112
 Age:49 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 400 MG ORAL	Cerebral Disorder Convulsion	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL	Hepatitis	Professional	Lysine (Lysine)	SS		ORAL
	Ischaemia Rash Erythematous		Insulin Human (Insulin Human)	SS		
SUBCUTANEOUS	SUBCUTANEOUS		Enoxaparin (Heparin)	SS		
SUBCUTANEOUS	SUBCUTANEOUS		Clopidogrel (Clopidogrel)	SS		ORAL
ORAL			Phenobarbital Clonazepam Thiopental	C C C		

Date:11/12/03ISR Number: 4235005-7Report Type:Expedited (15-DaCompany Report #2003115629
 Age:47 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1 GRAM (TID)	Headache Psychotic Disorder	Foreign Health	Neurontin (Gabapentin)	PS		
Other	Visual Disturbance	Professional Company Representative				

Date:11/13/03ISR Number: 4235743-6Report Type:Expedited (15-DaCompany Report #2003116234
 Age:8 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Drug Level Decreased	Consumer	Neurontin (Solution) (Gabapentin)	PS		ORAL
15 CC (TID), ORAL				Valproic Acid (Valproic Acid)	C		
				Levetiracetam (Levetiracetam)	C		
				All Other Therapeutic Products	C		

Date:11/13/03ISR Number: 4235836-3Report Type:Expedited (15-DaCompany Report #2003116539
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Asthenopia Dependence	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Dizziness Drug Abuser Dry Mouth Dysarthria Keratoconjunctivitis Sicca Overdose Vision Blurred		All Other Therapeutic Products	C		
				Aporex (Paracetamol, Dextropropoxyphene Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/13/03ISR Number: 4235837-5Report Type:Expedited (15-DaCompany Report #2003116209

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Apathy	Consumer	Neurontin			
2700 MG		Cold Sweat		(Gabapentin)	PS		ORAL
(TID), ORAL		Diabetes Mellitus					
		Social Avoidant Behaviour		Hydrocodone			
		Suicidal Ideation		(Hydrocodone)	C		

Date:11/13/03ISR Number: 4235855-7Report Type:Expedited (15-DaCompany Report #2003116111

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Cholesterol	Consumer	Lipitor			
Initial or Prolonged		Increased		(Atorvastatin)	PS		ORAL
10 MG							
Other		Confusional State					
(DAILY), ORAL		Convulsion		Neurontin			
		Drug Ineffective		(Gabapentin)			
(TID), ORAL		Dyskinesia		(Gabapentin)	SS		ORAL
		Pain		Clopidogrel Sulfate			
		Pain Of Skin		(Clopidrogel			
		Visual Acuity Reduced		Sulfate)	C		
				Losartan Potassium			
				(Losartan Potassium)	C		
				Valdecoxib			
				(Valdecoxib)	C		
				Cetirizine			
				Hydrochloride			
				(Cetirizine			
				Hydrochloride)	C		
				Gliclazide			
				(Gliclazide)	C		
				Insulin Lispro			
				(Insulin Lispro)	C		

Date:11/13/03ISR Number: 4235897-1Report Type:Expedited (15-DaCompany Report #KII-2001-0001303
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional Company Representative	Morphine Sulfate (Similar To Nda 19-516)(Morphine Sulfate) Other Neurontin (Gabapentin) Unknown	PS		SS

Date:11/14/03ISR Number: 4235512-7Report Type:Direct Company Report #CTU 206088
Age:58 YR Gender:Female I/FU:I

Outcome	PT
Life-Threatening	Anger
Hospitalization -	Crying
Initial or Prolonged	Facial Pain
Disability	Formication
	Headache
	Hemiplegia
	Impaired Work Ability
	Speech Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Trigeminal Neuralgia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
INTRACERVICAL	ONE AT NIGHT		Neurontin 300 Mg Pfizer	PS	Pfizer	
INTRACERVICAL						

Date:11/14/03ISR Number: 4236196-4Report Type:Expedited (15-DaCompany Report #2003116915
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other 300 MG (TID),		Attention-Seeking Behaviour	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Blood Pressure Increased					
ORAL		Drug Effect Decreased Drug Level Decreased Fluid Retention Intentional Self-Injury Suicide Attempt		Lithium (Lithium) (Lithium) Valproate Semisodium (Valproate Semisodium)	SS SS		ORAL

Date:11/14/03ISR Number: 4236203-9Report Type:Expedited (15-DaCompany Report #2003040766
Age:5 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Congenital Anomaly Other		Autism Maternal Drugs Affecting Foetus	Health Professional	Neurontin (Gabapentin) Carbamazepine (Carbamazepine)	PS SS		

Date:11/14/03ISR Number: 4236208-8Report Type:Expedited (15-DaCompany Report #2003116212
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (TID), Other ORAL		Dissociation Feeling Abnormal Overdose Respiratory Arrest Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Quetiapine Fumarate (Quetiapine Fumarate)	SS		
				Clonidine (Clonidine)	SS		
				Sertraline Hydrochloride (Sertraline Hydrochloride)	C		
				Bupropion Hydrochloride (Bupropion Hydrochloride)	C		
				Risperidone (Risperidone)	C		
				Ibuprofen (Ibuprofen)	C		
				All Other Therapeutic Products	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/14/03ISR Number: 4236210-6Report Type:Expedited (15-DaCompany Report #2003116213
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Antinuclear Antibody Positive	Consumer	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL		Family Stress		Hydrochlorothiazide (Hydrochlorothiazide)	C		

Date:11/14/03ISR Number: 4236212-XReport Type:Expedited (15-DaCompany Report #2003116214
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Carpal Tunnel Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Dizziness					
Other		Drug Ineffective		Atorvastatin (Atorvastatin)	C		
		Fatigue		Levothyroxine Sodium (Levothyroxine Sodium)	C		
		Hypoaesthesia		Dyazide (Hydrochlorothiazide , Triamterene)	C		
		Hypokalaemia		Anitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
		Mood Swings		Chlordiazepoxide (Chlordiazepoxide)	C		
		Pain In Extremity					
		Viral Infection					

Date:11/14/03ISR Number: 4236213-1Report Type:Expedited (15-DaCompany Report #2003116561
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Hospitalisation	Consumer	Neurontin (Gabapentin)	PS		
Other							

Date:11/14/03ISR Number: 4236218-0Report Type:Expedited (15-DaCompany Report #2003030526

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Aggression	Consumer	Neurontin			
Initial or Prolonged	Bipolar Disorder	Health	(Gabapentin)	PS		
4800 MG						
Other	Condition Aggravated	Professional				
(DAILY)	Psychotic Disorder					
	Suicide Attempt					
	Weight Decreased					

Date:11/14/03ISR Number: 4236427-0Report Type:Expedited (15-DaCompany Report #USA-2002-0002855

Age:51 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Accidental Overdose
Initial or Prolonged	Anxiety
	Asthenia
	Cardiac Failure

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Congestive Chronic Obstructive Pulmonary Disease Coagulopathy				
SEE IMAGE		Coma Condition Aggravated Confusional State Decreased Appetite	Consumer Health Professional Other		Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS ORAL
SEE IMAGE		Depressed Level Of Consciousness Depression Disturbance In Attention			Oxycontin Tablets 20 Mg (Oxycodone Hydrochloride) Cr Tablet	SS ORAL
ORAL		Drug Dependence Drug Toxicity Drug Withdrawal Syndrome Dysuria			Hydrocodone Cp (Phenylephrine, Chlorphenamine Maleate) Syrup	SS ORAL
ORAL	7	DAY	Encephalopathy Fall		Darvocet-N (Paracetamol) Tablet	SS ORAL
ORAL		Head Injury Headache Hepatic Necrosis			Cyclobenzaprine (Cyclobenzaprine) Tablet	SS ORAL
ORAL		Hyperhidrosis Hypokalaemia Hypotension			Hydroxyzine Pamoate (Hydroxyzine Embonate)	SS ORAL
ORAL		Insomnia Loss Of Consciousness			Valium (Diazepam) Tablet	SS ORAL
ORAL	7	DAY	Mood Swings Multi-Organ Failure		Neurontin (Gabapentin) Tablet	SS ORAL
ORAL	7	DAY	Multiple Drug Overdose Nervousness Nightmare		Tegretal (Carbamazepine) Tablet	SS ORAL
ORAL	7	DAY	Oedema Peripheral Pain		Celexa (Citalopram Hydrobromide) Tablet	SS ORAL
ORAL	7	DAY	Paranoia Pneumonia Aspiration Pneumonia Klebsiella		Sonata (Zaleplon) Tablet Trazodone	SS ORAL

ORAL	7	DAY	Psychomotor Retardation	(Trazodone) Tablet	SS	ORAL
			Renal Failure	Lorcet (Paracetamol,		
			Respiratory Failure	Hydrocodone		
			Rhabdomyolysis	Bitartrate)	SS	
7	DAY		Sepsis	Promethazine		
			Speech Disorder	(Promethazine)	C	
			Tendonitis	Clindamycin		
			Vision Blurred	(Clindamycin)	C	
				Prednisone		
				(Prednisone)	C	
				Premarin Tablet		
				(Estrogens		
				Conjugated)	C	
				Xenical (Orlistat)	C	
				Flurazepam		
				(Flurazepam)	C	
				Prevacid		
				(Lansoprazole)	C	

Date:11/14/03ISR Number: 4236432-4Report Type:Expedited (15-DaCompany Report #2003117027
Age: Gender:Female I/FU:I

Outcome PT
Other Abnormal Behaviour
Burns Second Degree

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Somnolence Vertigo	Report Source	Product	Role	Manufacturer	Route
			Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:11/14/03ISR Number: 4236434-8Report Type:Expedited (15-DaCompany Report #2003116555
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Agitation Increased Appetite	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
		Nausea Nervousness Weight Increased		Venlafaxine (Venlafaxine) Zolpidem (Zolpidem) Metoprolol (Metoprolol) Clonazepam (Clonazepam)	C C C C		

Date:11/14/03ISR Number: 4236443-9Report Type:Expedited (15-DaCompany Report #2003116858
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other (TID), ORAL		Confusional State	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:11/14/03ISR Number: 4236445-2Report Type:Expedited (15-DaCompany Report #2003116859
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death Other (TID), ORAL	Anaemia Neoplasm Malignant	Foreign Health Professional	Neurontin (Gabapentin) Valproate Sodium (Valproate Sodium)	PS C	ORAL
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Date:11/17/03ISR Number: 4236255-6Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0313585A
 Age:39 YR Gender:Male I/FU:I

Outcome Dose Other 300MG per day 800MG Twice per day	Duration	PT Alcohol Withdrawal Syndrome Epilepsy Optic Neuropathy Visual Acuity Reduced	Report Source	Product Lamictal Neurontin Sectral Mogadon	Role PS SS C C	Manufacturer Glaxosmithkline	Route ORAL ORAL ORAL ORAL
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/17/03ISR Number: 4236261-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0314212A
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchial Carcinoma	Consumer	Combivir	PS	Glaxosmithkline	ORAL
Hospitalization - SUBCUTANEOUS		Cholestasis 23 MON		Macrolin (France)	SS	Glaxosmithkline	
Initial or Prolonged 1UNIT Per day		Cytolytic Hepatitis		Bactrim	SS	Glaxosmithkline	ORAL
		General Physical Health Deterioration		Malocide Neurontin	SS SS	Glaxosmithkline	ORAL ORAL
1200MG Per day		Hepatic Cancer Metastatic					
2250MG Per day		Pericarditis		Viracept	SS		ORAL
		Thrombocytopenia					
		Weight Decreased		Lioresal	C		

Date:11/17/03ISR Number: 4237026-7Report Type:Expedited (15-DaCompany Report #2003116648
Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchial Carcinoma	Foreign Health	Viracept (Tablet) (Nelfinavir Mesilate)	PS		ORAL
Hospitalization - Initial or Prolonged 2250 MG Disability (DAILY), ORAL Other		Cerebral Toxoplasmosis General Physical Health Deterioration	Professional				
		Hemiplegia Metastases To Liver Pericardial Disease		Neurontin (Tablets) (Gabapentin) (Gabapentin)	SS		ORAL
ORAL		Pneumonia Pulmonary Oedema		Aldesleukin (Aldesleukin)	SS		
SUBCUTANEOUS	SUBCUTANEOUS	Thrombocytopenia Weight Decreased		Pyrimethamine (Pyrimethamine)	SS		ORAL
ORAL				Zidovudine W/Lamivudine			

ORAL				(Zidovudine, Lamivudine)	SS		ORAL
ORAL				Sulfamethoxazole (Sulfamethoxazole)	SS		ORAL
				Baclofen (Baclofen)	C		

Date:11/17/03ISR Number: 4237029-2Report Type:Expedited (15-DaCompany Report #DEU-2003-0000487
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Drug Interaction Oedema Peripheral	Foreign Health Professional	Oxygesic (Oxycodone Hydrochloride) Cr Tablet	PS		ORAL
ORAL			Other	Gabapentin (Gabapentin)	SS		

Date:11/17/03ISR Number: 4237078-4Report Type:Expedited (15-DaCompany Report #2003115629
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Abnormal Behaviour Diplopia	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other (TID), ORAL		Hallucination, Auditory Headache Psychotic Disorder Visual Disturbance	Professional Company Representative	Paroxetine Hydrochloride (Paroxetine Hydrochloride) Oxybutynin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Oxybutynin) C
 Clonazepam
 (Clonazepam) C
 Baclofen (Baclofen) C

Date:11/17/03ISR Number: 4237086-3Report Type:Expedited (15-DaCompany Report #2003116215
 Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG (TID), ORAL		Accident Condition Aggravated Fall Lower Limb Fracture Neuropathic Pain	Foreign Consumer	Gabapentin (Gabapentin) All Other Therapeutic Products Carbamazepine (Carbamazepine) Ranitidine (Ranitidine)	PS C C C		ORAL

Date:11/17/03ISR Number: 4237138-8Report Type:Expedited (15-DaCompany Report #2003116246
 Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 1200 MG (EVERY DAY), ORAL		Bronchial Carcinoma Cholestasis Cytolytic Hepatitis General Physical Condition Abnormal Hepatic Lesion Pericardial Disease Pulmonary Oedema Thrombocytopenia Weight Decreased	Foreign Health Professional	Neurontin (Gabapentin) Viracept (Tablet) (Nelfinavir Mesilate)	PS SS		ORAL ORAL

SUBCUTANEOUS	SUBCUTANEOUS	(Interleukin-2)	SS	
		Bactrim (Sulfamethoxazole, Trimethoprim)	SS	ORAL
ORAL				
		Pyrimethamine (Pyrimethamine)	SS	ORAL
ORAL				
		Zidovudine W/Lamivudine (Zidovudine, Lamivudine)	SS	ORAL
ORAL				
		Baclofen (Baclofen)	C	

Date:11/18/03ISR Number: 4237623-9Report Type:Expedited (15-DaCompany Report #2003116914
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged	Hospitalisation	Consumer	Neurontin (Gabapentin)	PS		

Date:11/18/03ISR Number: 4237627-6Report Type:Expedited (15-DaCompany Report #2003116912
Age:37 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Multiple Sclerosis	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL

3200 MG (800,

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Freedom Of Information (FOI) Report

QID), ORAL

Date: 11/18/03
 ISR Number: 4237837-8
 Report Type: Expedited (15-DaCompany Report #2002068598)
 Age: 30 YR Gender: Female I/FU: F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (THREE TIMES DAILY), ORAL	Arthralgia Asthma Bronchitis Acute Condition Aggravated Cough Dizziness Drug Interaction Dyspnoea Inflammation Nasopharyngitis Pain Pyrexia	Foreign Study Health Professional	Gabapentin (Tablets) (Gabapentin) Carbamazepine (Carbamazepine) Clobazam (Clobazam) Theophylline (Theophylline) Pranlukast (Pranlukast) Salbutamol (Salbutamol) Budesonide (Budesonide) Tulobuterol Hydrochloride (Tulobuterol Hydrochloride) Methylprednisolone Sodium Succinate (Methylprednisolone Sodium Succinate) Aminophylline (Aminophylline) Actit (Maltose, Sodium Acetate, Potassium Chloride, Sodium Chloride, Potassium Phosphate Solita T (Electrolytes Nos) Cefdinir (Cefdinir) Acemetacin (Acemetacin)	PS C		ORAL

Pentoxifyverine
(Pentoxifyverine) C
Sodium Chloride
(Sodium Chloride) C

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
Death	Areflexia
Hospitalization -	Brain Death
Initial or Prolonged	Cardio-Respiratory Arrest
	Completed Suicide
	Depressed Level Of
	Consciousness
	Hyporeflexia
	Intentional Misuse
	Pupillary Reflex Impaired
	Subarachnoid Haemorrhage

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tachycardia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
UNKNOWN			Bupropion	PS	Glaxosmithkline	ORAL
UNKNOWN			Clonazepam	SS		
UNKNOWN			Trazodone	SS		
UNKNOWN			Gabapentin	SS		

Date:11/19/03ISR Number: 4237465-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379987A
 Age:35 YR Gender:Female I/FU:F

Outcome	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: 4238561-8Report Type:Expedited (15-DaCompany Report #2003037701
 Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1800 MG	Toxic Epidermal Necrolysis	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other (TID), ORAL			Professional				
			Company Representative	Amitriptyline (Amitriptyline)	C		
				Analgesics	C		
				Atenolol (Atenolol)	C		

Enalapril Maleate (Enalapril Maleate) C
 Betahistine Hydrochloride (Betahistine Hydrochloride) C
 Lormetazepam (Lormetazepam) C
 Piascledine (Soya Oil, Perseae Oleum) C
 Caffeine/Opium/Paracetamol (Caffeine, Paracetamol, Opium Tincture) C

Date:11/19/03ISR Number: 4238582-5Report Type:Expedited (15-DaCompany Report #2003117025
 Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG ORAL	Blood Creatine Increased Coordination Abnormal	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
2.5 MG ORAL		Dysstasia Tremor	Professional	Ramipril (Ramipril)	SS		ORAL
				Terbutaline Sulfate	C		
				Zolpidem Tartrate	C		

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Freedom Of Information (FOI) Report

Insulin Human
 Injection, Isophane C
 All Other
 Therapeutic Products C
 Glyceryl Trinitrate C
 Folic Acid, Calcium Phosphate) C
 Oxazepam C
 Cromoglicic Acid Sodium C
 Glipizide C
 Alfacalcidol C
 Carbamazepine C
 Furosemide C
 Budesonide C
 Acetylcysteine C
 Simvastatin C
 Acetylsalicylic Acid C
 Insulin Injection, Isophane) C
 Isosorbide
 Mononitrate C
 Salmeterol Xinafoate C
 Orlistat C
 Bisoprolol C

Date:11/19/03ISR Number: 4238914-8Report Type:Expedited (15-DaCompany Report #2003117320
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Loss Of Consciousness Somnolence	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 MG (TID), ORAL				Propacet (Paracetamol, Dextropropoxyphene Napsilate) Oxycocet (Paracetamol, Oxycodone Hydrochloride) Valdecoxib	C		

(Valdecoxib)	C
Nitrofurantoin	
(Nitrofurantoin)	C
Cyclobenzaprine	
Hydrochloride	
(Cyclobenzaprine	
Hydrochloride)	C
Alprazolam	
(Alprazolam)	C
Librax	
(Chlordiazepoxide	
Hydrochloride,	
Clidinium Bromide)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/19/03ISR Number: 4238938-0Report Type:Expedited (15-DaCompany Report #2003117242

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Optic Neuritis	Consumer	Neurontin (Gabapentin)	PS		

Date:11/19/03ISR Number: 4238979-3Report Type:Expedited (15-DaCompany Report #2003041305

Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 4800 MG		Acute Respiratory Failure Pneumonia Aspiration	Health Professional	Neurontin (Gabapentin)	PS		ORAL

(QID), ORAL

				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
--	--	--	--	--	---	--	--

Date:11/19/03ISR Number: 4239229-4Report Type:Expedited (15-DaCompany Report #2003117241

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Hospitalization - 800 MG (TID), Initial or Prolonged ORAL		Condition Aggravated Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Drug Ineffective Loss Of Consciousness Memory Impairment Suicidal Ideation					

Date:11/20/03ISR Number: 4238034-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0379990A

Age:46 YR Gender: I/FU:F

Outcome	Duration	PT	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: 4239380-9Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Acrochordon
Disability	Actinic Keratosis
Other	Adjustment Disorder With Depressed Mood

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Freedom Of Information (FOI) Report

Alanine Aminotransferase
Increased
Alcoholism
Anaphylactic Shock
Anxiety
Aspartate
Aminotransferase
Increased
Back Injury
Bipolar Disorder
Blood Urea
Nitrogen/Creatinine Ratio
Increased
Brain Neoplasm
Coma
Contusion
Conversion Disorder
Convulsion
Dependence
Depressed Level Of
Consciousness
Depression
Dermal Cyst
Disturbance In Attention
Dizziness
Drug Abuser
Drug Hypersensitivity
Dysarthria
Dysphemia
Electroencephalogram
Abnormal
Expressive Language
Disorder
Fall
Fatigue
Fibromyalgia
Folliculitis
Gamma-Glutamyltransferase
Increased
Hallucination
Head Injury
Hypersomnia
Hypoaesthesia
Hypoglycaemia
Impaired Work Ability
Intentional Misuse
Limb Injury

Loss Of Consciousness
Memory Impairment
Meniscus Lesion
Mental Impairment
Nervous System Disorder
Neurodegenerative
Disorder
Nuclear Magnetic
Resonance Imaging Brain
Abnormal
Osteopenia
Pain
Paraesthesia
Paraneoplastic Syndrome

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Patellofemoral Pain Syndrome Pigmented Naevus	Report Source	Product	Role	Manufacturer	Route
400 MG (BID),		Road Traffic Accident Seborrheic Keratosis	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Sinusitis	Professional				
		Skin Irritation Sleep Disorder Speech Disorder Suicidal Ideation Suicide Attempt Tinnitus Tremor Vision Blurred Weight Increased	Company Representative	Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin)	SS C C C C		

Date:11/20/03ISR Number: 4239508-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 1800 MG		Antinuclear Antibody Positive	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL		Blood Albumin Decreased	Professional				
		Blood Glucose Increased Brain Damage Coordination Abnormal Depression Drug Hypersensitivity Dysarthria Dyskinesia Dysphagia Emotional Distress Gingival Disorder Intentional Misuse		Lamotrigine (Lamotrigine) Lithium Carbonate (Lithium Carbonate) Clonazepam (Clonazepam) Methylphenidate Hydrochloride Levothyroxine Sodium (Levothyroxine Sodium)	C C C C C C		

Malnutrition	Liothyronine Sodium	
Medication Error	(Liothyronine	
Nervous System Disorder	Sodium)	C
Suicidal Ideation	Sertraline	
Swollen Tongue	Hydrochloride(Sertra	
Tardive Dyskinesia	line Hydrochloride)	C
Temperature Intolerance	Pilocarpine	
Tongue Injury	Hydrochloride	
	(Polocarpine	
	Hydrochloride)	C
	Metoprolol Succinate	
	(Metoprolol	
	Succinate)	C
	Lansoprazole	
	(Lansoprazole)	C
	Hyoscyamine Sulfate	
	(Hyoscyamine	
	Sulfate)	C
	Diltiazem	
	Hydrochloride	
	(Diltiazem	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C
 Lamotrigine
 (Lamotrigine) C

Date:11/20/03ISR Number: 4239672-3Report Type:Expedited (15-DaCompany Report #2003117657
 Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Fall	Foreign	Neurontin			
Hospitalization -		Orthostatic Hypotension	Health	(Gabapentin)	PS		
Initial or Prolonged		Vertigo	Professional				
Other		Wrist Fracture	Company				
			Representative				

Date:11/20/03ISR Number: 4239703-0Report Type:Expedited (15-DaCompany Report #2003117707
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Diabetic Complication	Foreign	Neurontin			
Initial or Prolonged		Hypotonia	Health	(Gabapentin)	PS		
300 MG (TID)		Hypovolaemia	Professional	Cilazapril			
		Metabolic Disorder		(Cilazapril)	C		
		Orthostatic Hypotension		Acetylsalicylic Acid			
				(Acetylsalicylic	C		
				Acid)			
				Clindamycin			
				Hydrochloride			
				(Clindamycin	C		
				Hydrochloride)			
				Ciprofloxacin			
				(Ciprofloxacin)	C		
				Insulin Glargine			
				(Insulin Glargine)	C		

Date:11/20/03ISR Number: 4239706-6Report Type:Expedited (15-DaCompany Report #2003117658
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death
Other

Face Oedema

Foreign
Health
Professional
Company
Representative

Neurontin
(Gabapentin)

PS

Date:11/21/03ISR Number: 4240124-5Report Type:Expedited (15-DaCompany Report #US056015
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - SUBCUTANEOUS	25 MG, TWICE	Renal Failure Acute	Study	Enbrel	PS		
Initial or Prolonged WEEKLY, SC			Health				
SUBCUTANEOUS	20 MG, DAILY,		Professional	Teriparatide	SS		
SC				Gabapentin	SS		ORAL
300 MG, 3							
TIMES/DAY, PO				Quinapril Hydrochloride	C		
				Metoprolol	C		
				Panadeine Co	C		

Freedom Of Information (FOI) Report

Baclofen C

Date:11/21/03ISR Number: 4240171-3Report Type:Expedited (15-DaCompany Report #2003117891
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Consumer	Neurontin			
800 MG (BID),		Agitation		(Gabapentin)	PS		ORAL
ORAL		Delusion					
		Drug Withdrawal Syndrome		Diphenhydramine			
		Mood Swings		Hydrochloride			
		Paranoia		(Diphenhydramine			
				Hydrochloride)	C		

Date:11/21/03ISR Number: 4240451-1Report Type:Expedited (15-DaCompany Report #2003036800
Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Carpal Tunnel	Foreign	Gabapentin			
Initial or Prolonged		Decompression	Consumer	(Gabapentin)	PS		ORAL
400 MG		Oedema Peripheral					
(DAILY), ORAL		Post Procedural		All Other			
		Complication		Therapeutic Products	C		

Date:11/21/03ISR Number: 4240921-6Report Type:Expedited (15-DaCompany Report #DE-SHR-03-015636
Age:41 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Alanine Aminotransferase	Foreign	Betaferon			
Initial or Prolonged		Increased	Consumer	(Interferon Beta-1b)	PS		
SUBCUTANEOUS	SUBCUTANEOUS	Ascites	Health	Neurontin(Gabapentin			
Other		Asthenia	Professional)	SS		ORAL
600 MG,							

3X/DAY, ORAL

Blood Bilirubin Increased Other
Fatigue
Gallbladder Disorder
Hepatomegaly
Jaundice
Liver Function Test
Abnormal

Date:11/24/03ISR Number: 4239995-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0314709A

Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -	4 DAY	Dysaesthesia		Clamoxyl	PS	Glaxosmithkline	ORAL
Initial or Prolonged	5DROP Per day	Hypochloraemia		Laroxyl Drops	SS	Glaxosmithkline	ORAL
	9UNIT Per day	Hyponatraemia		Lioresal	SS		ORAL
	2UNIT Per day	Pyrexia		Neurontin	SS		ORAL
	7DROP per day			Rivotril Drops	SS		
	3UNIT Per day			Dantrium	SS	Glaxosmithkline	ORAL
	3TAB Per day			Heptamyl	C		ORAL
				Diantalvic	C		
				Eductyl	C		
				Forlax	C		
				Oroken	C		
	2 DAY			Rocephine	C		
INTRAVENOUS				Gentalline	C	Glaxosmithkline	
INTRAVENOUS							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/24/03ISR Number: 4240152-XReport Type:Expedited (15-DaCompany Report #ES-BRISTOL-MYERS SQUIBB COMPANY-12335402
 Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Karvea Tabs 300 Mg	PS	Bristol-Myers Squibb Company	ORAL
9 DAY		Drug Interaction					
				Adiro	C		
				Zocor	C		
				Dilutol	C		
				Buscapina	C		
				Atrovent	C		
RESPIRATORY							
(INHALATION)							
				Almax	C		
				Urolosin	C		
				Beglan	C		
RESPIRATORY							
(INHALATION)							
RESPIRATORY				Pulmicort	C		
(INHALATION)							
				Polaramine	C		
				Neurontin	I		
				Phenytoin +			
				Phenobarbital	I		

Date:11/24/03ISR Number: 4240527-9Report Type:Direct Company Report #CTU 206734
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Tremor		Neurontin 300 Mg	PS		ORAL
300MG AT							
BEDTIME ORAL							

Date:11/24/03ISR Number: 4240794-1Report Type:Direct Company Report #CTU 206766
 Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Activities Of Daily		Neurontin	PS		
1 MON							
Initial or Prolonged		Living Impaired					
Required		Depression					
Intervention to		Hallucination					
Prevent Permanent		Nervousness					
Impairment/Damage		Psychotic Disorder					
		Suicidal Ideation					

Date:11/24/03ISR Number: 4241206-4Report Type:Expedited (15-DaCompany Report #2003110107
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Blood Glucose Increased	Health	Neurontin			
Initial or Prolonged		Breast Cancer	Professional	(Gabapentin)	PS		ORAL
3600 MG							
Other		Cholecystitis Acute					
(TID), ORAL							
		Hyperlipidaemia		Hydrocodone			
		Renal Impairment		(Hydrocodone)	C		

Date:11/24/03ISR Number: 4241689-XReport Type:Expedited (15-DaCompany Report #2003117608
Age:42 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

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG		Abdominal Pain Cholecystitis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
(TID), ORAL		Flatulence					
		Intervertebral Disc Disorder		Clonazepam (Clonazepam)	C		
		Pancreatitis		Atenolol (Atenolol)	C		
		Ulcer Haemorrhage		Lansoprazole (Lansoprazole)	C		
		Vomiting		Oxycocet (Paracetamol, Oxycodone Hydrochloride)	C		

Date:11/24/03ISR Number: 4241691-8Report Type:Expedited (15-DaCompany Report #2003117610

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	600 MG (BID),	Blood Pressure Abnormal Cardiac Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other ORAL		Nausea					
		Renal Failure Acute Vomiting		Isosorbide (Isosorbide)	C		
				Heparin-Fraction, Sodium Salt (Heparin-Fraction, Sodium Salt)	C		
				Metoprolol Tartrate (Metoprolol Tartrate)	C		
				Furosemide (Furosemide)	C		
				Paracetamol (Paracetamol)	C		
				Ondansetron Hydrochloride (Ondansetron			

Hydrochloride) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C
Glyceryl Trinitrate
(Glyceryl
Trinitrate) C

Date:11/24/03ISR Number: 4241737-7Report Type:Expedited (15-DaCompany Report #2003118182
Age:63 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Hypoaesthesia
Initial or Prolonged	Intervertebral Disc
Other	Protrusion
	Lung Neoplasm Malignant
	Middle Insomnia
	Muscle Atrophy
	Paraesthesia
	Spinal Cord Compression

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100 MG	(DAILY), ORAL	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Captopril (Captopril)	C		
			Vicodin (Paracetamol, Hydrocodone)	C		
			Metolazone (Metolazone)	C		
			Furosemide (Furosemide)	C		
			Potassium (Potassium)	C		
			Metformin Hydrochloride (Metformin Hydrochlorde)	C		
			Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)	C		
			Insulin (Insulin)	C		
			All Other Therapeutic Products	C		

Date:11/24/03ISR Number: 4243526-6Report Type:Expedited (15-DaCompany Report #SAGL/03/25/UNK
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alanine Aminotransferase Increased Aspartate	Foreign Health Professional	Sandoglobulin (Immunoglobulin Human Normal)	PS		
INTRAVENOUS	30 G, IV	Aminotransferase	Other	Gabapentin	SS		
300 MG, BID		Increased		Paracetamol	C		

Blood Bilirubin Increased
Gamma-Glutamyltransferase
Increased
Hepatocellular Damage

Tramadol C
Diazepam C
Tinzaparin Sodium
(Heparin-Fraction,
Sodium Salt) C
Movicol (Nulytely) C
Nystatin C
Mst Continus "Asta
Medica" (Morphine
Sulfate) C

Date:11/25/03ISR Number: 4240941-1Report Type:Expedited (15-DaCompany Report #CA-ROCHE-351823
Age:79 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 5 DAY	Lethargy	Health	Tamiflu	PS	Roche	ORAL
Initial or Prolonged	Thrombosis Vomiting	Professional	Levothyroxine Asa Calcium Vitamin D	SS SS SS SS	Roche	ORAL ORAL ORAL

UNKNOWN

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

UNKNOWN				Atenolol	SS	Roche
UNKNOWN				Etidronate	SS	
UNKNOWN				Gabapentin	SS	
UNKNOWN				Carbamazepine	SS	

Date:11/25/03ISR Number: 4242153-4Report Type:Expedited (15-DaCompany Report #2003118177
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin			
Other		Facial Palsy Medication Error		(Gabapentin)	PS		

Date:11/25/03ISR Number: 4242357-0Report Type:Expedited (15-DaCompany Report #2003-03900
 Age:34 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Literature Health Professional	Hydroxyzine (Watson Laboratories) (Hydroxyzine Hydrochloride) Tablet	PS		
				Perphenazine/Amitrip tyline Hydrochloride Unknown Strength (Watson) (Amitriptyline Gabapentin (Gabapentin)	SS SS		

Date:11/25/03ISR Number: 4242414-9Report Type:Expedited (15-DaCompany Report #03P-163-0240786-00
 Age:54 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Death	Completed Suicide	Literature	Hydrocodone/Acetaminophen (Vicodin)		
		Health	(Hydrocodone/Acetaminophen)	PS	ORAL
ORAL		Professional			
			Amitriptyline	SS	ORAL
ORAL					
			Gabapentin	SS	ORAL
ORAL					

Date:11/25/03ISR Number: 4242702-6Report Type:Expedited (15-DaCompany Report #2003117889
Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 800 MG, ORAL	Back Pain Blood Bicarbonate	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
	Abnormal Blood Uric Acid Decreased Dysaesthesia Heat Stroke	Professional	Amoxicillin Trihydrate (Amoxicillin Trihydrate)	SS		ORAL
ORAL						
	Hypochloraemia Inappropriate Antidiuretic Hormone Secretion		Heptaminol Hydrochloride (Heptaminol Hydrochloride)	SS		ORAL
563.4 MG, ORAL	Pyrexia					
			Amitriptyline			

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ORAL	Hydrochloride (Amitriptyline Hydrochloride)	SS	ORAL
90 MG, ORAL	Baclofen (Baclofen)	SS	ORAL
75 MG, ORAL	Dantrolene Sodium (Dantrolene Sodium)	SS	ORAL
	Clonazepam (Clonazepam)	C	
	Dextropropoxyphene (Dextropropoxyphene)	C	
	Paracetamol (Paracetamol)	C	
	Eductyl (Sodium Bicarbonate, Potassium Bitartrate)	C	
	Macrogol (Macrogol)	C	
	Cefixime (Cefixime)	C	
	Ceftriaxone (Ceftriaxone)	C	

Date:11/25/03ISR Number: 4242704-XReport Type:Expedited (15-DaCompany Report #2003118180
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG (ONCE), ORAL	Medication Error Sedation Tremor	Foreign Health Professional	Neurontin (Gabapentin) Fentanyl (Fentanyl)	PS C		ORAL

Date:11/25/03ISR Number: 4242905-0Report Type:Periodic Company Report #2003001895
Age:79 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 10 MG	Blood Glucose Increased Hypertension	Consumer	Lipitor (Atorvastatin)	PS		ORAL

(DAILY), ORAL	Migraine				
	Muscle Spasms		Neurontin (Gabapentin)	SS	ORAL
100 MG					
(DAILY), ORAL			Levothyroxine Sodium (Levothyroxine Sodium)	C	
			Diazepam (Diazepam)	C	

Date:11/26/03ISR Number: 4243394-2Report Type:Expedited (15-DaCompany Report #2003118576
Age:67 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Pyrexia	Foreign	Neurontin			
Initial or Prolonged	Viral Infection	Health	(Gabapentin)	PS		
900 MG (TID)						
	Vomiting	Professional	Glimepiride			
		Company	(Glimepiride)	C		
		Representative	Zofenopril Calcium			
			(Zofenopril Calcium)	C		
			Clopidogrel	C		
			Fenofibrate			
			(Fenofibrate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/26/03ISR Number: 4243403-0Report Type:Expedited (15-DaCompany Report #2003118605
Age:36 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 800 MG (BID), ORAL	Generalised Oedema Hepatitis Hepatosplenomegaly Infectious Mononucleosis Pyrexia Swelling	Foreign Consumer	Gabapentin (Gabapentin) Carbamazepine (Carbamazepine)	PS C		ORAL

Date:11/26/03ISR Number: 4243532-1Report Type:Expedited (15-DaCompany Report #2003118984
Age:71 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID), ORAL	Abdominal Pain Arthritis Blood Glucose Decreased Change Of Bowel Habit Chondropathy Fatigue Food Allergy Irritable Bowel Syndrome Neuropathy Peripheral Pancreatic Enzymes Abnormal Vision Blurred	Consumer	Neurontin (Gabapentin) Glucosamine Sulfate/Minerals/Mul tivitamins (Glucosamine Sulfate, Vitamins Hyoscyamine Sulfate (Hyoscyamine Sulfate) Pancrelipase (Pancrelipase)	PS SS SS C		ORAL

Date:11/26/03ISR Number: 4244168-9Report Type:Expedited (15-DaCompany Report #2003037701
Age:56 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG	Neuropathic Pain Toxic Epidermal	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL

(TID), ORAL

Necrolysis

Professional

Company
Representative

Caffeine/Opium/Paracetamol (Caffeine, Paracetamol, Opium Tincture)	C
Amitriptyline (Amitriptyline)	C
Analgesics	C
Atenolol (Atenolol)	C
Enalapril Maleate (Enalapril Maleate)	C
Betahistine Hydrochloride (Betahistine Hydrochloride)	C
Lormetazepam (Lormetazepam)	C
Piascledine (Soya Oil, Perseae Oleum)	C
Enalapril (Enalapril)	C
Betahistine (Betahistine)	C
Lormetazepam (Lormetazepam)	C

Freedom Of Information (FOI) Report

All Other
Therapeutic Products C

Date:11/26/03ISR Number: 4244441-4Report Type:Expedited (15-DaCompany Report #2003038069

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL Other	Blood Pressure Increased Haemorrhage Malaise Pain In Extremity Weight Decreased	Consumer Health Professional	Neurontin (Gabapentin) Clopidogrel Sulfate (Clopidogrel Sulfate) Glyceryl Trinitrate (Glyceryl Trinitrate) Risedronate Sodium (Risedronate Sodium) Isosorbide Mononitrate (Isosorbide Mononitrate) Calcium Carbonate (Calcium Carbonate) Atenolol (Atenolol) Lorazepam (Lorazepam) Simvastatin (Simvastatin) Zolpidem Tartrate (Zolpidem Tartrate)	PS SS C C C C C C C C		ORAL

Date:11/26/03ISR Number: 4244668-1Report Type:Expedited (15-DaCompany Report #2003034005

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL Other	Arterial Occlusive Disease Cardiac Failure Congestive	Consumer	Neurontin (Gabapentin) Risedronate Sodium (Risedronate Sodium)	PS C		ORAL

Dehydration	Esomeprazole	C
Drug Ineffective	(Esomeprazole)	
Fatigue	Levothyroxine Sodium	
Feeling Abnormal	(Levothyroxine	
Pain	Sodium)	C
Spinal Disorder	Clopidogrel Sulfate	
Urinary Incontinence	(Clopidogrel	
	Sulfate)	C
	Metoprolol Succinate	
	(Metoprolol	
	Succinate)	C
	Vicodin	
	(Paracetamol,	
	Hydrocodone	
	Bitartrate)	C
	Amlodipine Besilate	
	(Amlodipine	
	Besilate)	C
	Simvastatin	
	(Simvastatin)	C

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Meclozine
 (Meclozine) C
 Alprazolam
 (Alprazolam) C
 Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C

Date:11/26/03ISR Number: 4244740-6Report Type:Expedited (15-DaCompany Report #2003037043
 Age:8 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dysphagia	Health	Neurontin			
Other		Neoplasm Malignant	Professional	(Gabapentin)	PS		ORAL
600 MG (BID),							
ORAL							

Date:11/28/03ISR Number: 4244109-4Report Type:Expedited (15-DaCompany Report #2003118832
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Cholesterol Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1600 MG							
(UNKNOWN),							
ORAL							
Blood Creatine							
Professional							
Phosphokinase Increased							
Blood Creatinine Increased							
Levetiracetam (Levetiracetam)							
SS							
4000 MG, ORAL							
Blood Triglycerides Increased							
Lamotrigine (Lamotrigine)							
SS							
400 MG, ORAL							
Liver Function Test Abnormal							
Mirtazapine (Mirtazapine)							
SS							
45 MG							
Myalgia							
(UNKNOWN),							

ORAL

Bisoprolol Fumarate
(Bisoprolol
Fumarate) C

Date:11/28/03ISR Number: 4244658-9Report Type:Expedited (15-DaCompany Report #FRP03000847

Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 25 MG, 3/DAY,		Back Pain Blood Osmolarity Decreased	Foreign Health Professional	Dantrium (Dantrolene Sodium)Capsule, 25mg	PS		ORAL
ORAL		Blood Uric Acid Decreased Dysaesthesia	Other	Clamoxyl (Amoxicillin Trihydrate)	SS		ORAL
ORAL		Heat Stroke Hypochloraemia Hyponatraemia		Hept-A-Myl (Heptamino l Hydrochloride)	SS		ORAL
563.4 MG, DAILY , ORAL		Inappropriate Antidiuretic Hormone Secretion		Laroxyl (Amitriptylin e Hydrochloride)	SS		ORAL
ORAL		Pyrexia Urine Sodium Abnormal		Lioresal "Ciba-Geigy" (Baclofe n)	SS		ORAL
90 MG DAILY				Neurontin (Gabapentin)	SS		ORAL
ORAL							
800 MG, DAILY , ORAL							

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<p>0.7 MG , DAILY ,ORAL</p> <p>2 DF, DAILY, INJECTION NOS</p>	<p>Rivotril(Clonazepam) SS</p> <p>Di-Antalvic(Dextropropoxyphene Hydrochloride) SS</p> <p>Eductyl(Potassium Bitartrate, Sodium Bicarbonate) SS</p> <p>Forlax(Macrogol) SS</p> <p>Oroken(Cefixime) SS</p> <p>Rocephin(Ceftriaxone Sodium) SS</p> <p>Gentamycin-Mp(Gentamicin Sulfate) SS</p>	<p>ORAL</p>
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Date:11/28/03ISR Number: 4244925-9Report Type:Expedited (15-DaCompany Report #2003-04122
Age:27 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature Health Professional	Doxepin(Watson Laboratories)(Doxepin Hydrochloride) Capsule	PS		
				Gabapentin (Gabapentin)	SS		
				Olanzapine (Olanzapine)	SS		

Date:11/28/03ISR Number: 4245075-8Report Type:Expedited (15-DaCompany Report #2003116648
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged 2250 MG		Brain Abscess Cerebral Toxoplasmosis Cholestasis	Foreign Health Professional	Viracept (Tablet) (Nelfinavir Mesilate)	PS		ORAL

Disability (DAILY), ORAL Other	Cytolytic Hepatitis			
	Hemiplegia	Neurontin (Tablets)		
	Lung Carcinoma Cell Type	(Gabapentin)		
1200 MG	Unspecified Stage Iv	(Gabapentin)	SS	ORAL
(DAILY), ORAL	Malignant Neoplasm			
	Progression	Aldesleukin		
	Metastases To Liver	(Aldesleukin)	SS	
SUBCUTANEOUS	SUBCUTANEOUS			
	Pericardial Disease	Pyrimethamine		
ORAL	Pneumonia	(Pyrimethamine)	SS	ORAL
	Pulmonary Oedema	Zidovudine W/ Lamivudine		
	Thrombocytopenia	(Zidovudine, Lamivudine)	SS	ORAL
ORAL		Sulfamethoxazole		
		(Sulfamethoxazole)	SS	ORAL
ORAL		Baclofen (Baclofen)	C	

Date:11/28/03ISR Number: 4245077-1Report Type:Expedited (15-DaCompany Report #2003119402

Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source
Hospitalization -	Dizziness	Foreign
Initial or Prolonged	Dyskinesia	Health

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Professional

Dose	Duration	Product	Role	Manufacturer	Route
300 MG TID		Neurontin (Gabapentin)	PS		ORAL
ORAL					

Date:11/28/03ISR Number: 4245078-3Report Type:Expedited (15-DaCompany Report #2003119508
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Mood Altered Overdose	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL							
Other			Professional Company Representative	All Other Therapeutic Products Diamorphine	SS C		

Date:11/28/03ISR Number: 4245161-2Report Type:Expedited (15-DaCompany Report #2003119358
Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Chronic Obstructive Airways Disease	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
300 MG TID		Exacerbated	Professional				
ORAL				Furosemide	C		
				Ramipril	C		
				Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)	C		
				Salbutamol	C		
				Aminophylline	C		

Date:11/28/03ISR Number: 4245165-XReport Type:Expedited (15-DaCompany Report #2003119407
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Mood Altered Overdose	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:11/28/03ISR Number: 4245360-XReport Type:Expedited (15-DaCompany Report #2003110069
Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG (TID), ORAL		Bladder Spasm Feeling Abnormal Nervousness Vaginitis Bacterial White Blood Cell Count Decreased	Health Professional	Neurontin (Gabapentin) Zonisamide (Zonisamide) Ropinirole Hydrochloride (Ropinirole Hydrochloride)	PS SS SS		ORAL

Freedom Of Information (FOI) Report

Tizanidine
 Hydrochloride
 (Tizanidine
 Hydrochloride) SS
 Montelukast Sodium
 (Montelukast Sodium) C
 All Other
 Therapeutic Products C
 Multivitamins
 (Ergocalciferol,
 Ascorbic Acid, Folic
 Acid, Thiamine
 Hydrochloride, C
 Magnesium
 (Magnesium) C
 Calcium (Calcium) C
 Ascorbic Acid
 (Ascorbic Acid) C
 Zinc (Zinc) C
 Chromium (Chromium) C
 Policosanol
 (Policosanol) C
 Lactobacillus
 Acidophilus
 (Lactobacillus
 Acidophilus) C

Date:11/28/03ISR Number: 4245388-XReport Type:Expedited (15-DaCompany Report #2012272
 Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Death	Accidental Overdose	Consumer	Oxycodone			
Hospitalization - Initial or Prolonged	Back Pain	Health Professional	Hydrochloride (Similar To Nda 20-553) (Oxycodone Hydrochloride)	PS		ORAL
Other	Haematemesis Headache	Other				
ORAL			Diazepam (Diazepam)	SS		
			Oxazepam (Oxazepam)	SS		
			Temazepam (Temazepam)	SS		
			Lorazepam (Lorazepam)	SS		
			Cannabnoids			

(Cannabis)	SS
Diphenhydramine	
Hydrochloride	
(Diphenhydramine	
Hydrochloride)	SS
Gabapentin	
(Gabapentin)	SS
Claritin	
(Loratadine)	C
Zoloft (Sertraline	
Hydrochloride)	C
Altace (Ramipril)	C
Allopurinol	
(Allopurinol)	C
Depakote (Valproate	
Semisodium)	C
Soma (Carisoprodol)	C

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Relafen (Nabumetone) C

Date:11/28/03ISR Number: 4245416-1Report Type:Expedited (15-DaCompany Report #2003114317
 Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Consumer	Neurontin			
		Difficulty In Walking	Health	(Gabapentin)	PS		ORAL
100 MG (AS							
NEEDED), ORAL		Muscle Spasms	Professional				
				Diazepam (Diazepam)	C		
				Warfarin Sodium			
				(Warfarin Sodium)	C		

Date:11/28/03ISR Number: 4245418-5Report Type:Expedited (15-DaCompany Report #2003001458
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arrhythmia	Consumer	Neurontin			
Other		Condition Aggravated		(Gabapentin)	PS		ORAL
3200 MG (FOUR							
TIMES DAILY),		Depression					
ORAL		Dysphonia					
		Fatigue		Vitamins	C		
		Multiple Chemical					
		Sensitivity					
		Nausea					
		Seasonal Affective					
		Disorder					

Date:11/28/03ISR Number: 4245451-3Report Type:Expedited (15-DaCompany Report #2003110156
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		No Adverse Drug Effect	Health	Neurontin			

Professional

(Gabapentin)

PS

Date:12/02/03ISR Number: 4244850-3Report Type:Expedited (15-DaCompany Report #US-SHR-03-002517

Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Burning Sensation Chest Pain Disease Recurrence	Consumer Health Professional	Betaseron (Interferon Beta-1b) Injection			
SUBCUTANEOUS	8 MIU, EVERY 2D,	Dizziness			PS		
SUBCUTANEOUS		Dysphagia					
ORAL		Electrocardiogram Abnormal		Neurontin (Gabapentin)	SS		ORAL
		Hyperaesthesia Oral Candidiasis Pruritus Generalised		Quinine Sulfate (Quinine Sulfate)	C		

Date:12/02/03ISR Number: 4246001-8Report Type:Expedited (15-DaCompany Report #CH-JNJFOC-20031104186

Age: Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Choreoathetosis Drug Interaction

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Somnolence

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
TRANSDERMAL	TRANSDERMAL	Foreign Health	Durogesic (Fentanyl) Patch	PS		
600 MG, 1 IN		Professional	Neurontin (Gabapentin)	SS		ORAL
1 DAY, ORAL			Mst Continus (Morphine Sulfate)	SS		ORAL
40 MG, 1 IN 1			Ciproxin (Ciprofloxacin)	C		
DAY, ORAL			Dalacin (Clindamycin Hydrochloride)	C		

Date:12/02/03ISR Number: 4246008-0Report Type:Expedited (15-DaCompany Report #2003119658
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign Health	Neurontin (Gabapentin)	PS		
Other		Confusional State Grand Mal Convulsion Hypertension Salivary Hypersecretion Tachycardia	Professional				

Date:12/02/03ISR Number: 4246059-6Report Type:Expedited (15-DaCompany Report #2003119630
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Cleft Palate Congenital Anomaly	Foreign Health	Gabapentin (Gabapentin)	PS		
PLACENTAL		Maternal Drugs Affecting Foetus	Professional				

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Life-Threatening	Drug Interaction	Health	Neurontin			
Hospitalization -	Feeling Abnormal	Professional	(Gabapentin)	PS		ORAL
ORAL						
Initial or Prolonged	Heart Injury		Metaxalone			
Other	Insomnia		(Metaxalone)	SS		
	Restlessness		Cyclobenzaprine			
	Suicide Attempt		Hydrochloride			
			(Cyclobenzaprine			
			Hydrochloride)	SS		
			Mirtazapine			
			(Mirtazapine)	SS		
			Quetiapine Fumarate			
			(Quetiapine			
			Fumarate)	SS		
			Amitriptyline			
			Hydrochloride			
			(Amitriptyline			
			Hydrochloride)	SS		
			Estrogens Conjugated			
			(Estrogens			

Freedom Of Information (FOI) Report

Conjugated) C
 Levothyroxine Sodium
 (Levothyroxine Sodium) C
 Oxycocet
 (Paracetamol, Oxycodone Hydrochloride) C
 Axotal (Old Form)
 (Caffeine, Butalbital, Paracetamol) C
 Tizanidine Hydrochloride
 (Tizanidine Hydrochloride) C
 Parafon Forte
 (Chlorzoxazone, Paracetamol) C
 Morphine (Morphine) C

Date:12/02/03ISR Number: 4246274-1Report Type:Expedited (15-DaCompany Report #2003119599
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG (BID), Other ORAL	Carotid Bruit Drug Ineffective Hypertension International Normalised Ratio Abnormal Neuropathy	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Heparin (Heparin)	C		
			Warfarin Sodium (Warfarin Sodium)	C		
			Ramipril (Ramipril)	C		
			Insulin Injection, Isophane (Insulin Injection, Isophane)	C		
			Metoprolol (Metoprolol)	C		
			Glipizide (Glipizide)	C		
			Furosemide (Furosemide)	C		
			Digoxin (Digoxin)	C		
			Clonidine			

(Clonidine) C
 Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C
 Dextropropoxyphene
 Hydrochloride
 (Dextropropoxyphene
 Hydrochloride) C

Date:12/02/03ISR Number: 4246278-9Report Type:Expedited (15-DaCompany Report #2003116234
 Age:8 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Drug Ineffective	Consumer	Neurontin (Solution) (Gabapentin)	PS		ORAL
15 CC (TID), ORAL		Pharmaceutical Product Complaint		Valproic Acid			

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(Valproic Acid) C
 Levetiracetam
 (Levetiracetam) C
 All Other
 Therapeutic Products C

Date:12/02/03ISR Number: 4246326-6Report Type:Expedited (15-DaCompany Report #2003033537
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Confusional State	Consumer	Neurontin			
Other		Memory Impairment	Health	(Gabapentin)	PS		
2400 MG (QID)		Pain	Professional	Paroxetine			
		Tremor		Hydrochloride			
				(Paroxetine			
				Hydrochloride)	C		
				Trazodone			
				(Trazodone)	C		
				Ibuprofen			
				(Ibuprofen)	C		

Date:12/03/03ISR Number: 4246524-1Report Type:Expedited (15-DaCompany Report #2003119620
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Conjunctival Haemorrhage	Health	Neurontin			
ORAL			Professional	(Gabapentin)	PS		ORAL
				Escitalopram			
				(Escitalopram)	C		
				Estrogens Conjugated			
				(Estrogens			
				Conjugated)	C		
				Famotidine			
				(Famotidine)	C		
				Multivitamins			
				(Ergocalciferol,Asco			
				rbic Acid, Folic			
				Acid, Thiamine			
				Hydrochloride,	C		
				Tocopherol			

(Tocopherol)

C

Date:12/03/03ISR Number: 4246525-3Report Type:Expedited (15-DaCompany Report #2003119747
Age:15 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Overdose	Consumer	Neurontin			
Initial or Prolonged	Suicide Attempt		(Gabapentin)	PS		ORAL
300 MG						
Other	Vomiting					
(DAILY), ORAL						
			Paracetamol			
			(Paracetamol)	SS		ORAL
			Acetylcysteine			
			(Acetylcysteine)	SS		
			Celexa (Citalopram			
			Hydrobromide)	C		

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Date:12/03/03ISR Number: 4246526-5Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20031001759

Age:30 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged SEEC IMAGE, ORAL	Aggression Psychotic Disorder Self Injurious Behaviour	Consumer Health Professional	Topamax (Topiramate)	PS		ORAL
			Topamax (Topiramate)	SS		
			Topamax (Topiramate)	SS		
			Neurontin (Gabapentin)	SS		ORAL
SEE IMAGE, ORAL						

Date:12/03/03ISR Number: 4246627-1Report Type:Expedited (15-DaCompany Report #2003008068

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Disability Other 1200 MG (TID), ORAL	Anaemia Aplastic Anaemia Arthralgia Asthenia Balance Disorder Burns Second Degree Cardiac Disorder Coma Convulsion Coordination Abnormal Difficulty In Walking Disturbance In Attention Dizziness Dry Mouth Fall General Physical Health Deterioration Headache Hypotonia Loss Of Consciousness	Consumer	Neurontin (Gabapentin) Dilantin (Phenytoin Sodium)	PS SS		ORAL
			Acetylsalicylic Acid (Acetylsalicylic Acid)	SS		
			Vitamins With Minerals	SS		
			Megestrol (Megestrol)	C		

Mental Disorder
Mental Impairment
Muscle Spasms
Nervous System Disorder
Nervousness
Pain
Panic Attack
Paresis
Speech Disorder

Date:12/03/03ISR Number: 4246632-5Report Type:Expedited (15-DaCompany Report #2003018755
Age:37 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abasia
Initial or Prolonged	Abdominal Distension
Disability	Abdominal Pain
Other	Agitation
	Anxiety
	Arthralgia
	Back Pain
	Blood Calcium Decreased
	Blood Creatinine

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Decreased
Blood Electrolytes
Decreased
Blood Glucose Decreased
Blood Lactate
Dehydrogenase Decreased
Blood Potassium Decreased
Blood Pressure Increased
Blood Triglycerides
Increased
Blood Urea Decreased
Brain Damage
Bursitis
Cardiovascular Disorder
Cerebral Artery Occlusion
Cerebral Atrophy
Cerebral Infarction
Cerebrovascular Accident
Constipation
Convulsion
Depression
Diarrhoea
Difficulty In Walking
Dizziness
Dysmenorrhoea
Electrocardiogram St-T
Change
Emotional Distress
Extrapyramidal Disorder
Eye Pain
Facial Palsy
Faecal Occult Blood
Positive
Flatulence
Gliosis
Grand Mal Convulsion
Haemorrhoids
Hallucination, Auditory
Hemiparesis
Herpes Simplex
Insomnia
Intracranial Aneurysm
Irritable Bowel Syndrome
Joint Dislocation
Loss Of Consciousness
Major Depression
Mean Cell Haemoglobin
Increased

Mean Cell Volume
Increased
Medication Error
Menorrhagia
Menstruation Irregular
Migraine
Mitral Valve Incompetence
Movement Disorder
Muscle Atrophy
Muscle Contracture
Muscle Rigidity
Muscle Spasms
Muscle Strain
Oral Pain

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
2400 MG (600, FOUR TIMES A DAY); ORAL (SEE IMAGE)		Osteopenia Pain Parkinsonism Pco2 Decreased Peroneal Nerve Palsy Photopsia Psychomotor Retardation Psychotic Disorder	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
UNKNOWN (UNKNOWN), UNKNOWN	UNKNOWN	Red Blood Cell Count Decreased Road Traffic Accident Schizophrenia, Disorganised Type Somatisation Disorder		Loestrin (Anovlar) (Norethindrone Acetate, Ethinyl Estradiol)	SS		
20 MG (Q12H); ORAL (SEE IMAGE)		Suicidal Ideation Urinary Incontinence Weight Increased		Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL

Date:12/03/03ISR Number: 4246668-4Report Type:Expedited (15-DaCompany Report #2003119691
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG, ORAL		Dermatitis Allergic Exanthem	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
5 MG			Professional	Risperidone (Risperidone)	SS		
				Amitriptyline Hydrochloride			

100 MG

(Amitriptyline Hydrochloride) SS

500 MG

Paracetamol (Paracetamol) SS

Lithium Acetate (Lithium Acetate) C
Lactulose (Lactlose) C

Date:12/03/03ISR Number: 4246796-3Report Type:Expedited (15-DaCompany Report #2003119648

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2400 MD	Balance Disorder Complex Regional Pain Syndrome	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Disability (QID), ORAL		Condition Aggravated		Oxaprozin (Oxaprozin)	C		
Other		Depression Dizziness Fall Gait Disturbance Injury Panic Attack		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride) Lorazepam (Lorazepam) Levothyroxine Sodium (Levothyroxine Sodium)	C C C C		

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Date:12/03/03ISR Number: 4246880-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 400 MG (BID), Disability ORAL Other	Anxiety Back Pain Bipolar Disorder Brain Neoplasm Cerebrovascular Disorder Cognitive Disorder Coma Conversion Disorder Convulsion Disease Recurrence Dizziness Drug Abuser Dysarthria Ependymoma Fatigue Hyperaemia Loss Of Consciousness Memory Impairment Nervous System Disorder Single Photon Emission Computerised Tomogram Abnormal Somnolence Speech Disorder Tremor	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin)	PS SS C C C C		ORAL

Date:12/03/03ISR Number: 4246882-8Report Type:Expedited (15-DaCompany Report #DEU-2003-0000665

Age:80 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 20 MG, BID, ORAL	Choreoathetosis Somnolence	Foreign Health Professional Other	Morphine Sulfate Cr Tablet (Morphine Sulfate) Cr Tablet Neurontin	PS		ORAL

600 MG,

(Gabapentin)Capsule SS

ORAL

DAILY, ORAL

Durogesic(Fentanyl)
Patch SS

TRANSDERMAL TRANSDERMAL

Ciproxin
(Ciprofloxacin) C
Dalacin (Clindamycin
Hydrochloride) C

Date:12/03/03ISR Number: 4246990-1Report Type:Expedited (15-DaCompany Report #2003119892
Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Alkaline	Foreign	Gabapentin			
Other		Phosphatase Increased	Health	(Gabapentin)	PS		ORAL
2.4 GRAM		Osteomalacia	Professional				
(DAILY), ORAL			Other				

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Date:12/03/03ISR Number: 4247071-3Report Type:Direct
Age:53 YR Gender:Male I/FU:I

Company Report #CTU 207406

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 30 MG Q8H						
Initial or Prolonged ORAL	Medication Error		Methadone 10 Mg	PS		ORAL
Required 600 MG Q8H	Mental Status Changes					
Intervention to ORAL	Overdose		Gabapentin 300 Mg	SS		ORAL
Prevent Permanent Impairment/Damage			Augmentin	C		
			Doss	C		
			Vicodin	C		
			Hydroxyzine	C		
			Aspirin	C		
			Ranitidine	C		

Date:12/03/03ISR Number: 4250116-8Report Type:Periodic
Age:48 YR Gender:Female I/FU:I

Company Report #WAES 0309USA00377

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose UNK/PO						
300 MG/TID/PO	Ageusia	Consumer	Tab Zetia 10 Mg	PS		ORAL
	Asthenia		Neurontin Unk	SS		
	Back Pain		Paracetamol Tablets			
	Dysgeusia		Bp	C		
	Glossodynia		Fosinopril Sodium	C		
	Headache		Sertraline			
	Neuropathy Peripheral		Hydrochloride	C		
	Pain In Extremity					
	Stress					

Date:12/04/03ISR Number: 4245723-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0441720A
Age: Gender:Male I/FU:F

Outcome	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: 4246695-7Report Type:Expedited (15-DaCompany Report #2003039760
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Osteitis Deformans	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:12/04/03ISR Number: 4246699-4Report Type:Expedited (15-DaCompany Report #2003116213
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Antinuclear Antibody Positive	Consumer	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL				Hydrochlorothiazide (Hydrochlorothiazide)	C		

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Date:12/04/03ISR Number: 4246700-8Report Type:Expedited (15-DaCompany Report #2003119899

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (BID, Other ORAL	Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:12/04/03ISR Number: 4246712-4Report Type:Expedited (15-DaCompany Report #ENZ-ABE-156

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged INTRAVENOUS 350 MG QD IV	Blood Creatinine Increased Cholestasis	Health Professional	Abelcet (Amphotericin B Lipid) Abelcet (Amphotericin B Lipid Complex Inj) Abelcet (Amphotericin B Lipid Complex Inj) Neurontin	PS SS SS SS		ORAL
300 MG BID ORAL			Augmentin Effexor Soma Lisinopril Singulair Allegra Mvi Advair Lasix Combivent Nph Insulin Celebex	SS C C C C C C C C C C		

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG (TID), Other ORAL	Cataract Diabetic Neuropathy Drug Ineffective Eye Swelling Impaired Healing	Consumer Health Professional	Neurontin (Gabapentin) Insulin Lispro (Insulin Lispro) Insulin Glargine (Insulin Glargine) Metformin Hydrochloride (Metformin Hydrochloride) Glibenclamide (Glibenclamide) Metolazone (Metolazone) Simvastatin (Simvastatin) Oxycocet (Paracetamol, Oxycodone	PS C C C C C C		ORAL

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Hydrochloride) C
 Citalopram
 Hydrobromide
 (Citalopram
 Hydrobromide) C
 Lisinopril
 (Lisinopril) C
 Alprazolam
 (Alprazolam) C
 Potassium Chloride
 (Potassium Chloride) C
 Fenofibrate
 (Fenofibrate) C

Date:12/04/03ISR Number: 4246722-7Report Type:Expedited (15-DaCompany Report #2003119698
 Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Body Height Decreased	Consumer	Neurontin			
600 MG, ORAL		Cataract		(Gabapentin)	PS		ORAL
		Drug Ineffective		Feldene (Piroxicam)	C		
		Eye Oedema		Lipitor			
		Somnolence		(Atorvastatin)	C		
		Visual Disturbance		Lisinopril			
				(Lisinopril)	C		
				Verapamil			
				(Verapamil)	C		
				Furosemide			
				(Furosemide)	C		
				Potassium			
				(Potassium)	C		

Date:12/04/03ISR Number: 4248507-4Report Type:Expedited (15-DaCompany Report #2003119973
 Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -							
Initial or Prolonged		Phlebothrombosis	Foreign	Neurontin			
1800 MG		Post Thrombotic Syndrome	Health	(Gabapentin)	PS		ORAL
Disability			Professional				
(TID), ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation	Consumer	Neurontin			
Hospitalization - 200 MG (BID), Initial or Prolonged ORAL		Blood Pressure Decreased	Health	(Gabapentin)	PS		ORAL
Other ORAL		Blood Pressure Diastolic Increased Condition Aggravated	Professional	Morphine Sulfate (Morphine Sulfate)	SS		ORAL
		Confusional State		Atenolol (Atenolol)	SS		
		Delirium		Calcium (Calcium)	C		
		Drug Ineffective		Centrum Silver			
		Drug Interaction		(Ascorbic Acid,			
		Gastrointestinal Cancer Metastatic		Tocopheryl Acetate, Retinol, Zinc,			
		Pain		Calcium, Vitamins	C		
		Thinking Abnormal		Docusate Sodium			
		Weight Decreased		(Docusate Sodium)	C		
				Limbitrol			

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(Chlordiazepoxide,
Amitriptyline
Hydrochloride) C

Date:12/05/03ISR Number: 4247286-4Report Type:Expedited (15-DaCompany Report #2003034280
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Convulsion	Consumer Health Professional	Neurontin (Gabapentin) Diazepam (Diazepam) Oxycocet (Paracetamol, Oxycodone Hydrochloride) Ibuprofen (Ibuprofen)	PS C C C		

Date:12/05/03ISR Number: 4247287-6Report Type:Expedited (15-DaCompany Report #2003119930
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1500 MG, ORAL Other		Burns Third Degree Intentional Misuse Intervertebral Disc Disorder Loss Of Consciousness Spinal Disorder Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin) Alprazolam (Alprazolam) Escitalopram (Escitalopram) Venlafaxine Hydrochloride (Venlafaxine Hydrochloride) Quetiapine Fumarate (Quetiapine Fumarate)	PS SS C C C		ORAL

Date:12/05/03ISR Number: 4247366-3Report Type:Expedited (15-DaCompany Report #2003116212
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dissociation Irritability	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other ORAL		Mental Disorder	Professional				
		Overdose Respiratory Arrest		Quetiapine Fumarate (Quetiapine Fumarate)	SS		
				Clonidine (Clonidine)	SS		
				Sertraline Hydrochloride (Sertraline Hydrochloride)	C		
				Bupropion Hydrochloride (Bupropion Hydrochloride)	C		
				Risperidone (Risperidone)	C		
				Ibuprofen			

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(Ibuprofen) C
 All Other
 Therapeutic C

Date:12/05/03ISR Number: 4247369-9Report Type:Expedited (15-DaCompany Report #2003116111
 Age:66 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 20 MG Other (DAILY), ORAL	Blood Cholesterol Increased	Consumer Health	Lipitor (Atorvastatin)	PS		ORAL
2700 MG (TID), ORAL	Confusional State Convulsion Movement Disorder Muscle Spasms Pain Visual Disturbance	Professional	Neurontin (Gabapentin) (Gabapentin)	SS		ORAL
			Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
			Losartan Potassium (Losartan Potassium)	C		
			Valdecoxib (Valdecoxib)	C		
			Cetirizine Hydrochloride (Cetirizine Hydrochloride)	C		
			Gliclazide (Gliclazide)	C		
			Insulin Lispro (Insulin Lispro)	C		
			Novolin 20/80 (Insulin Human, Insulin Isophane, Human Biosynthetic)	C		
			Pioglitazone (Pioglitazone)	C		
			Sertraline Hydrochloride (Sertraline Hydrochloride)	C		
			Ezetimibe			

(Ezetimibe)

C

Date:12/05/03ISR Number: 4248341-5Report Type:Expedited (15-DaCompany Report #2003120549
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Joint Stiffness Swelling	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
1200 MG (QID), ORAL			Professional				
			Other	Ibuprofen (Ibuprofen)	C		

Date:12/08/03ISR Number: 4248521-9Report Type:Direct Company Report #CTU 207615
Age: Gender:Female I/FU:I

Outcome
Life-Threatening
Disability

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Other Required Intervention to Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Prevent Permanent Impairment/Damage 1200 MG 3 TIMES ORAL	Anger Chills Diarrhoea Drug Abuser Drug Dependence Drug Withdrawal Syndrome Emotional Disorder Hostility Influenza Like Illness Musculoskeletal Pain Urticaria		Neurontin 400 Mg Pfizer	PS	Pfizer	ORAL

Date:12/08/03ISR Number: 4248539-6Report Type:Expedited (15-DaCompany Report #001-0719-M0100407

Age: Gender:Male I/FU:F

Outcome Dose Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1200 MG (600 MG, BID), Disability ORAL Other 1600 MG (400 MG, 2 CAPSULES IN AM AND PM ONE EVERY 5 MIN., MAX: 3 (PRN)	Anhedonia Anxiety Asthenia Atrial Tachycardia Blood Calcium Decreased Blood Cholesterol Increased Blood Pressure Increased Blood Triglycerides Increased Bronchitis Cardiomegaly	Consumer Health Professional	Lopid (Gemfibrozil) Neurontin (Gabapentin) Nitroglycerin (Nitroglycerin)	PS SS SS		ORAL

0.4 MG

(DAILY)

Chest Discomfort
Chest Pain
Cough

Cerivastatin Sodium
(Cerivasstatin
Sodium) SS

Decreased Activity

Drug Ineffective
Drug Interaction
Electrocardiogram St-T
Change
Gastritis
Headache
High Density Lipoprotein
Decreased
Muscle Spasms
Nasal Congestion
Neuropathy
Pain
Pain In Extremity
Palpitations
Paraesthesia
Peripheral Coldness
Respiratory Tract
Congestion
Rhabdomyolysis
Sinus Bradycardia
Sinusitis
Supraventricular
Extrasystoles
Ventricular Extrasystoles

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Date:12/08/03ISR Number: 4248548-7Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 400 MG (BID), Disability ORAL Other	Anxiety Back Pain Bipolar Disorder Bone Cyst Brain Neoplasm Cognitive Disorder Coma Convulsion Depression Dizziness Drug Dependence Electroencephalogram Abnormal Ependymoma Fall Fatigue Head Injury Intentional Misuse Loss Of Consciousness Memory Impairment Nervous System Disorder Scar Somnolence Speech Disorder Suicide Attempt Tremor	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Valproate Sodium Buspirone Hydrochloride Fluoxetine Hydrochloride Omeprazole Capsaicin	PS SS C C C C		ORAL

Date:12/08/03ISR Number: 4248662-6Report Type:Expedited (15-DaCompany Report #2003118576

Age:67 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID)	Febrile Infection Lymphopenia Pyrexia Viral Infection Vomiting	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Glimepiride (Glimepiride) Zofenopril Calcium	PS C		

(Zofenoril Calcium) C
 Clopidogrel
 (Clopidogrel) C
 Fenofibrate
 (Fenofibrate) C
 Fosinopril
 (Fosinopril) C

Date:12/08/03ISR Number: 4248682-1Report Type:Expedited (15-DaCompany Report #2003116858
 Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID), ORAL		Agitation Confusional State Lymphadenopathy	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Celecoxib (Celecoxib)	C		
				Morphine Sulfate (Morphine Sulfate)	C		

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Fentanyl
 (Fentanyl) C
 Clorazepate
 Dipotassium
 (Clorazepate
 Dipotassium) C
 All Other
 Therapeutic Products C

Date:12/08/03ISR Number: 4248749-8Report Type:Expedited (15-DaCompany Report #CH-JNJFOC-20031104186
 Age:80 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged TRANSDERMAL	Duration Choreoathetosis Somnolence TRANSDERMAL	Foreign Health Professional	Durogesic (Fentanyl) Patch Neurontin (Gabapentin)	PS SS		ORAL
600 MG, 1 IN 1 DAY, ORAL			Mst Continus (Morphine Sulfate)	SS		ORAL
40 MG, 1 IN 1 DAY, ORAL						

Date:12/08/03ISR Number: 4248792-9Report Type:Expedited (15-DaCompany Report #K200301816
 Age:77 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2.5 MG, QD, ORAL	Duration Blood Creatinine Increased Coordination Abnormal Dysstasia	Foreign Health Professional Other	Altace Capsules (Ramipril)Capsule, 2.5 Mg Neurontin (Gabapentin) Capsule, 100mg	PS SS		ORAL ORAL
100 MG, QD,	General Physical Health Deterioration Tremor					

Bricanyl	
(Terbutaline	
Sulfate)	C
Stilnoct (Zolpidem	
Tartrate)	C
Insulatard (Insulin	
Human Injection,	
Isophane) Injection	C
Nitromex (Glyceryl	
Trinitrate)	C
Folacin (Calcium	
Phosphate, Folic	
Acid)	C
Sobril (Oxazepam)	C
Lomudal	
(Cromoglicate	
Sodium)	C
Mindiab (Glipizide)	C
Etalpa	
(Alfacalcidol)	C
Tegretol	
(Carbamazepine)	C
Furix (Furosemide)	C
Pulmicort (Budesonide	
)	C

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Zocord	C
Trombyl (Acetylsalicylic Acid)	C
Mixtard(Insulin, Insulin Injection, Isophane) Injection	C
Imdur (Isosorbide Mononitrate)	C
Serevent Diskus (Salmeterol Xinafoate)	C
Xenical (Orlistat)	C
Emconcor (Bisoprolol)	C
Laktulose "Dak" (Lactulose)	C
Acetylcysteine	C

Date:12/08/03ISR Number: 4249081-9Report Type:Expedited (15-DaCompany Report #2003121041
Age:61 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG TID Other ORAL	Blood Glucose Increased Cholestasis Normochromic Normocytic Anaemia Thrombocytopenia Weight Decreased	Foreign Health Professional	Neurontin (Gabapentin) Piribedil (Piribedil)	PS SS		ORAL ORAL
50 MG DAILY ORAL			Omeprazole Propranolol Spironolactone	C C C		

Date:12/08/03ISR Number: 4249082-0Report Type:Expedited (15-DaCompany Report #2003116859
Age:73 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Hospitalization - Anaemia Foreign Neurontin (Tablets) PS ORAL
Initial or Prolonged Difficulty In Walking Health (Gabapentin)
900 MG TID

Professional

ORAL

Valproate Sodium C
Omeprazole C
All Other
Therapeutic Products C
Betahistine
Hydrochloride C
Paracetamol C
Lormetazepam C

Date:12/08/03ISR Number: 4249432-5Report Type:Expedited (15-DaCompany Report #2003116539
Age:43 YR Gender:Male I/FU:F

Outcome PT
Other Abnormal Behaviour
Disturbance In Attention
Dizziness
Drug Dependence
Dry Mouth

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dysarthria Euphoric Mood Eyelid Disorder					
600 MG (BID), ORAL		Intentional Misuse Keratoconjunctivitis Sicca	Consumer Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				All Other Therapeutic Products Aporex (Paracetamol, Dextropropoxyphene Hydrochloride) Paroxetine Hydrochloride (Paroxetine Hydrochloride) Enalapril Maleate (Enalapril Maleate) Dyazide (Hydrochlorothiazide , Triamterene) Didanosine (Didanosine) Lamivudine (Lamivudine) Ritonavir (Ritonavir) Amprenavir (Amprenavir) Pravastatin Sodium (Pravastatin Sodium) ..	C C C C C C C C C C C C C		

Date:12/08/03ISR Number: 4249437-4Report Type:Expedited (15-DaCompany Report #2003120443

Age:54 YR Gender:Male I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (TID), Other ORAL		Hospitalization - Initial or Prolonged Vision Blurred	Consumer	Neurontin (Gabapentin)	PS		ORAL

Visual Disturbance

Lisinopril
(Lisinopril) C
Buspirone
(Buspirone) C
Trazodone
(Trazodone) C
Bupropion
Hydrochloride
(Bupropion
Hydrochloride) C
Clonazepam
(Clonazepam) C

Date:12/09/03ISR Number: 4248970-9Report Type:Direct
Age:56 YR Gender:Male I/FU:I

Company Report #CTU 207752

Outcome PT
Hospitalization - Confusional State
Initial or Prolonged Dehydration
Disorientation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hypotension Mental Status Changes Renal Failure Acute	Report Source	Product	Role	Manufacturer	Route
				Gabapentin	PS		
				Aripiprazole	SS		
				Digoxin	C		
				Bacitracin 500/Polumyxin 10000u/Gm	C		
				Methadone	C		
				Zolmitriptan	C		
				Lisinopril	C		
				Clonazepam	C		
				Aspirin	C		
				Insulin Reg Human	C		
				Triamcinolone			
				Acetonide	C		
				Salsalate	C		
				Cpd Men .025% In			
				Euc:Tac 0.1%	C		
				Gemfibrizil	C		
				Flouxetine	C		
				Fluocinonide	C		
				Ammonium Lactate	C		
				Allopurinol	C		
				Hydroxyzine	C		
				Furosemide	C		
				Spirolactone	C		
				Metoprolol	C		

Date:12/09/03ISR Number: 4248993-XReport Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 207750

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Suicide Attempt		Neurontin 1200 Mg Tid Po	PS		ORAL

Date:12/10/03ISR Number: 4248881-9Report Type:Expedited (15-DaCompany Report #PHRM2003FR03170
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Back Pain		Lioresal	PS	Novartis Sector: Pharma	ORAL
90 mg/day		Blood Bicarbonate					
5760 MIN		Abnormal		Clamoxyl	SS		ORAL
563.4 mg/day		Blood Osmolarity		Hept-A-Myl	SS		ORAL
5 drops/day		Decreased		Laroxyl	SS		ORAL
75 mg/day		Blood Uric Acid Decreased		Dantrium	SS		ORAL
800 mg/day		Body Temperature		Neurontin	SS		ORAL
7 drops/day		Increased		Rivotril	SS		ORAL
		Dysaesthesia		Di-Antalvic	SS		ORAL
		Heat Stroke		Eductyl	SS		
RECTAL		Hypochloraemia		Forlax	SS		ORAL
		Hyponatraemia					
		Inappropriate					
		Antidiuretic Hormone					
		Secretion					
		Pyrexia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/10/03ISR Number: 4250486-0Report Type:Expedited (15-DaCompany Report #2003121195
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intracranial Aneurysm	Consumer	Zoloft (Sertraline)	PS		
Other				Neurontin (Gabapentin)	SS		
				Zolpidem Tartrate (Zolpidem Tartrate)	SS		

Date:12/11/03ISR Number: 4250362-3Report Type:Direct Company Report #CTU 207876
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Abnormal Behaviour Intentional Misuse		Neurontin 300 Mg Parke Davis	PS	Parke Davis	ORAL
300 MG PER DAY ORAL		Self Mutilation Suicide Attempt					

Date:12/11/03ISR Number: 4251022-5Report Type:Expedited (15-DaCompany Report #2003036714
Age:78 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anxiety Headache	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
100 MG (DAILY, ORAL		Insomnia Visual Acuity Reduced	Professional	Candesartan (Cilexetil (Candesartan)	C		
				Omeprazole (Imeprazole)	C		
				Dorzolamide Hydrochloride	C		
				Brimonidine Tartrate Tobradex (Dexamethasone,	C		

Tobramycin) C
 Tolterodine
 L-Tartrate C
 Estradiol C
 Risendronate Sodium C

Date:12/11/03ISR Number: 4251027-4Report Type:Expedited (15-DaCompany Report #2003117608
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG		Cholecystitis Convulsion	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other (TID), ORAL		Intervertebral Disc Disorder Pancreatitis Ulcer Haemorrhage		Clonazepam (Clonazepam) Atenolol (Atenolol) Lansoprazole Oxycocet (Paracetamol, Oxycodone Hydrochloride) Lansoprazole(Lansopr azole)	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/11/03ISR Number: 4251031-6Report Type:Expedited (15-DaCompany Report #2003030520
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema	Health	Neurontin			
		Oedema Peripheral	Professional	(Gabapentin)	PS		
900 MG (TID)							
		Pitting Oedema					
UNKNONWN							
				Testosterone			
				(Testosterone)	SS		
INTRAMUSCULAR	200 MG						
INTRAMUSCULAR							
				Indapamide			
				(Indapamide)	C		
				Levothyroxine			
				(Levothyroxine)	C		
				Esomeprazole	C		
				Losartan Potassium	C		
				Phyzyme-95	C		

Date:12/11/03ISR Number: 4251033-XReport Type:Expedited (15-DaCompany Report #2003113597
 Age:11 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		No Adverse Drug Effect	Consumer	Neurontin			
			Health	(Gabapentin)	PS		ORAL
600 MG,							
(BID), ORAL			Professional				

Date:12/11/03ISR Number: 4251034-1Report Type:Expedited (15-DaCompany Report #2003121167
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicide Attempt	Consumer	Neurontin			
				(Gabapentin)	PS		

Date:12/11/03ISR Number: 4251035-3Report Type:Expedited (15-DaCompany Report #2003121173
Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG DAILY		Impaired Work Ability					
ORAL		Liver Disorder Mental Disorder Pain Renal Disorder Suicide Attempt		Zoloft	SS		

Date:12/11/03ISR Number: 4251036-5Report Type:Expedited (15-DaCompany Report #2003121792
Age:44 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Alopecia Dizziness
Other	Feeling Abnormal Feeling Drunk Gait Disturbance Hypoaesthesia Liver Disorder Multiple Sclerosis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Poisoning Sedation Systemic Lupus Erythematosus	Report Source	Product	Role	Manufacturer	Route
900 MD (TID), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Setraline Hydrochloride)	C		

Date:12/11/03ISR Number: 4251303-5Report Type:Expedited (15-DaCompany Report #2003121081
Age: Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Paralysis	Report Source	Product	Role	Manufacturer	Route
800 MG (BID), ORAL			Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:12/11/03ISR Number: 4251304-7Report Type:Expedited (15-DaCompany Report #2003116555
Age: Gender:Female I/FU:F

Outcome Dose Hospitalization - Initial or Prolonged ORAL	Duration	PT Agitation Increased Appetite Nausea Nervousness Unevaluable Event Weight Increased	Report Source	Product	Role	Manufacturer	Route
			Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
				Venlafaxin E(Venlafaxine)	C		
				Zolpidem (Zolpidem)	C		
				Metoprolol (Metoprolol)	C		
				Clonazepam (Clonazepam)	C		

Date:12/11/03ISR Number: 4251318-7Report Type:Expedited (15-DaCompany Report #2003121737
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG		Carotid Artery Disease Epilepsy	Foreign Health	Neurontin (Gabapentin)	PS		
Other		Somnolence	Professional Company Representative	Tramadol (Tramadol) Clomipramine (Clomipramine) Oxybutynin (Oxybutynin) Baclofen (Baclofen) Acetylsalicylate Lysine (Acetylsalicylate Lysine) Lormetazepam (Lormetazepam) Nulytely (Sodium Bicarbonate, Potassium Chloride, Sodium Chloride, Macrogol) Risedronate Sodium (Risedronate Sodium)	C C C C C C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/12/03ISR Number: 4251693-3Report Type:Expedited (15-DaCompany Report #2003121162
Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG		General Physical Health Deterioration	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:12/12/03ISR Number: 4251697-0Report Type:Expedited (15-DaCompany Report #2003041498
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2100 MG (TID), ORAL		Haemorrhage	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Diane (Ethinylestradiol, Cyproterone Acetate) Budesonide (Budesonide) Almotriptan Malate (Almotriptan Malate) Zolmitriptan (Zolmitriptan)	SS C C C		

Date:12/12/03ISR Number: 4251738-0Report Type:Expedited (15-DaCompany Report #2003122905
Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (TID), ORAL		Convulsion Dizziness Nausea	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Ergenyl Chrono			

(Valproic Acid,
Valproate Sodium) C
Lamotrigine
(Lamotrigine) C

Date:12/12/03ISR Number: 4251740-9Report Type:Expedited (15-DaCompany Report #2003121165
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (TID), ORAL		Choreoathetosis Dyskinesia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Pethidine
(Pethidine) C

Date:12/12/03ISR Number: 4251757-4Report Type:Expedited (15-DaCompany Report #2003120951
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (BID), ORAL		Drug Ineffective Loss Of Consciousness Syncope	Consumer	Neurontin (Gabapentin)	PS		ORAL

Clonazepam

Freedom Of Information (FOI) Report

(Clonazepam)

SS

ORAL

0.5 MG ORAL

Date:12/12/03ISR Number: 4251794-XReport Type:Expedited (15-DaCompany Report #2003114316

Age:67 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 200 MG (BID), Other ORAL	Abdominal Discomfort Biliary Cirrhosis Diarrhoea	Consumer	Neurontin (Gabapentin)	PS		ORAL
	Diverticulum Intestinal Haemoglobin Decreased Heart Rate Increased Hepatic Haematoma Hypotension Ileus Intra-Abdominal Haemorrhage Liver Function Test Abnormal Pain Post Procedural Complication Sarcoidosis Sinus Rhythm Ventricular Extrasystoles		Fludrocortisone Acetate (Fludrocortisone Acetate) Alprazolam (Alprazolam) Phenelzine Sulfate (Phenelzine Sulfate) Olanzapine (Olanzapine) Estradiol (Estradiol) Medroxyprogesterone Acetate (Medroxyprogesterone Acetate) Estrogens Conjugated (Estrogens Conjugated) Fluticasone Propionate (Fluticasone Propionate) Guaifenesin (Guaifenesin) Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate) Ipratropium Bromide (Ipratropium Bromide)	C C		

Nystatin (Nystatin) C
 Macrogol (Macrogol) C
 Docusate Sodium
 (Docusate Sodium) C
 Multivitamins
 (Ergocalciferol,
 Ascorbic Acid, Folic
 Acid, Thiamine
 Hydrochloride, C

Date:12/12/03ISR Number: 4251879-8Report Type:Expedited (15-DaCompany Report #2003121316

Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Biliary Dilatation Hepatitis Cholestatic	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/12/03ISR Number: 4252172-XReport Type:Expedited (15-DaCompany Report #2003121159

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Endocardial Fibroelastosis	Consumer	Neurontin (Gabapentin)	PS		
TRANSPLACENTAL (UNKNOWN), PLACENTAL	3600 MG	Haemangioma					
		Maternal Drugs Affecting Foetus		Levetiracetam (Levetiracetam)	SS		
TRANSPLACENTAL (UNKNOWN), PLACENTAL	1500 MG						

Date:12/12/03ISR Number: 4252333-XReport Type:Expedited (15-DaCompany Report #2002068598

Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (THREE TIMES DAILY), ORAL		Arthralgia Asthma Bronchitis Acute Condition Aggravated Dizziness Nasopharyngitis	Foreign Study Health Professional	Gabapentin (Tablets) (Gabapentin) Carbamazepine (Carbamazepine) Clobazam (Clobazam) Theophylline (Theophylline) Pranlukast (Pranlukast) Salbutamol (Salbutamol) Budesonide (Budesonide) Tulobuterol Hydrochloride	PS C C C C C C		ORAL

(Tulobuterol Hydrochloride)	C
Methylprednisolone Sodium Succinate (Methylprednisolone Sodium Succinate)	C
Aminophylline (Aminophylline)	C
Actit (Maltose, Sodium Acetate, Potassium Chloride, Sodium Chloride, Potassium Phosphate Solita T (Electrolytes Nos)	C
Cefdinir (Cefdinir)	C
Acemetacin (Acemetacin)	C
Pentoxifyverine (Pentoxifyverine)	C
Sodium Chloride (Sodium Chloride)	C
Bromhexine Hydrochloride (Bromhexine Hydrochloride)	C
Procaterol	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride (Procaterol Hydrochloride)	C
Hydrocortisone Sodium Succinate (Hydrocortisone Sodium Succinate)	C
Soldem 3a (Sodium Lactate, Potassium Chloride, Sodium Chloride, Carbohydrates Nos)	C
Diazepam (Diazepam)	C
Magnesium (Magnesium)	C

Date:12/15/03ISR Number: 4252089-0Report Type:Direct
Age:42 YR Gender:Female I/FU:I

Company Report #CTU 208008

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 3 PILLS 2 X Initial or Prolonged DAY A ORAL		Abnormal Behaviour Anhedonia		Nuerontin 400 Mg Pfizer, Inc.	PS	Pfizer, Inc	ORAL
Required Intervention to 3 PILLS 2 X Prevent Permanent DAY A ORAL Impairment/Damage		Suicide Attempt		Gabapentin 400 Mg Pfizer, Inc	SS	Pfizer, Inc	ORAL

Date:12/15/03ISR Number: 4252104-4Report Type:Direct
Age:21 YR Gender:Female I/FU:I

Company Report #CTU 208016

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 600 MG 4 TIMES ORAL		Activities Of Daily Living Impaired Aggression Anger		Neurontin 600 Mg Pfizer Neurontin 800 Mg	PS	Pfizer	ORAL

800 MG 4	Educational Problem	Pfizer	SS	Pfizer	ORAL
TIMES ORAL	Hallucination				
	Headache	Ortho Evra	C		
	Loss Of Consciousness	Elmiron	C		
	Memory Impairment	Elavil	C		
	Nervous System Disorder	Dicyclomine	C		
	Personality Change				
	Suicidal Ideation				

Date:12/15/03ISR Number: 4252111-1Report Type:Direct Company Report #CTU 207999
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression		Neurontin 300 Mg 3			
300 MG 3		Confusional State		Times Daily Pfizer	PS	Pfizer	ORAL
TIMES ORAL		Mood Swings					
		Overdose		Morphine	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/15/03ISR Number: 4252634-5Report Type:Expedited (15-DaCompany Report #2003121603

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Dilantin Suspension (Phenytoin Sodium)	PS		ORAL
Other		Arthralgia Confusional State					
ORAL		Convulsion Drug Ineffective		Neurontin (Gabapentin)	SS		ORAL
1200 MG		Epilepsy					
(QID), ORAL		Myalgia Pain In Extremity Weight Increased					

Date:12/15/03ISR Number: 4253501-3Report Type:Expedited (15-DaCompany Report #2003121729

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		ORAL
Disability		Abnormal Behaviour Asthenia					
Other		Back Pain Dysarthria Feeling Abnormal Post Procedural Pain Speech Disorder Thinking Abnormal Treatment Noncompliance Vision Blurred Weight Increased		Diazepam (Diazepam) Vicodin (Paracetamol, Hydrocodone Bititrate)	C C		
ORAL							

Date:12/15/03ISR Number: 4253567-0Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		ORAL
Hospitalization - Initial or Prolonged		Back Pain Bipolar Disorder	Health				
400 MG (BID),							

Disability	Brain Neoplasm	Professional		
ORAL				
Other	Cerebral Perfusion Pressure Decreased	Company Representative	Valproate Sodium (Valproate Sodium)	SS
	Cognitive Disorder		Buspirone Hydrochloride	
	Conversion Disorder		(Buspirone Hydrochloride)	SS
	Convulsion		Fluoxetine Hydrochloride	
	Hyperaemia		(Fluoxetine Hydrochloride)	C
	Impaired Work Ability		Omeprazole (Omeprazole)	C
			Capsaicin (Capsaicin)	C

Date:12/16/03ISR Number: 4253860-1Report Type:Expedited (15-DaCompany Report #2003041305
Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Acute Respiratory Failure	Health Professional	Neurontin (Gabapentin)	PS		ORAL
4800 MG (QID)	Pneumonia Aspiration					
ORAL			Oxycodone			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride
(Oxycodone
Hydrochloride) C

Date:12/17/03ISR Number: 4253654-7Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 208249

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - INTRAVENOUS Initial or Prolonged IV	163 MG Q WEEK	Breast Cancer Metastatic Computerised Tomogram		Taxol 80 Mg/M2	PS		
600 MG TID		Abnormal Hemiparesis		Gabapentin 300 Mg Tab	SS		ORAL
ORAL		Malignant Neoplasm					
		Progression Nausea Vomiting		Multi Vitamin Calcium Supplement	C C		

Date:12/17/03ISR Number: 4254236-3Report Type:Direct
Age:36 YR Gender:Male I/FU:I

Company Report #CTU 208244

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other 400 3 ORAL Required Intervention to Prevent Permanent Impairment/Damage		Abnormal Behaviour Drug Withdrawal Syndrome		Neurontin 400 Mg Pd Gabapentin	PS		ORAL
		Hallucination Medication Error Stupor					

Date:12/17/03ISR Number: 4254273-9Report Type:Expedited (15-DaCompany Report #2003123698
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alcohol Use Cholestasis	Foreign Health	Neurontin (Gabapentin)	PS		

Hepatic Steatosis
Nausea

Professional
Company
Representative

Date:12/17/03ISR Number: 4254652-XReport Type:Expedited (15-DaCompany Report #2003122501
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Congenital Anomaly 1400 MG, PLACENTAL		Foetal Disorder Maternal Drugs Affecting Foetus Patent Ductus Arteriosus	Consumer	Neurontin (Gabapentin) Lithium Carbonate (Lithium Carbonate)	PS C		

Date:12/17/03ISR Number: 4254654-3Report Type:Expedited (15-DaCompany Report #2003113619
Age:48 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 5200 MG (DAILY), ORAL		Brain Neoplasm Convulsion	Health Professional	Neurontin (Gabapentin) Fluoxetine Hydrochloride	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Fluoxetine
Hydrochloride) C
Celecoxib
(Celecoxib) C
Lansoprazole
(Lansoprazole) C

Date:12/17/03ISR Number: 4255120-1Report Type:Expedited (15-DaCompany Report #US_031299200
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 20 MG/1 DAY Initial or Prolonged 25 MG/2 WEEK		Renal Failure Acute	Study	Teriparatide	PS		
			Health	Enbrel	SS		
			Professional	Gabapentin	SS		
				Quinapril Hydrochloride	C		
				Metoprolol	C		
				Panadeine Co	C		
				Baclofen	C		

Date:12/17/03ISR Number: 4267679-9Report Type:Periodic Company Report #USA-2003-0007509
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) (Oxycodone Hydrochloride)	PS		
				Acetaminophen (Paracetamol)	SS		
				Diphenhydramine (Diphenhydramine)	SS		
				Fluoxetine (Fluoxetine)	SS		
				Gabapentin (Gabapentin)	SS		
				Ibuprofen (Ibuprofen)	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose Multiple Drug Overdose Pain	Consumer Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride) Oxycodone Hydrochloride (Similar To Nda 20-553) (Oxycodone Hydrochloride) Neurontin (Gabapentin)	PS SS SS		
UNK MG,				Depakote (Valproate Semisodium)	SS		
UNK MG,				Celexa (Citalopram Hydrobromide)	SS		
UNK MG,				Diltiazem			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

UNK MG,

(Diltiazem) SS
 Toprol XL C
 Glucophage C
 Niferex C

Date:12/17/03ISR Number: 4268450-4Report Type:Periodic
 Age:46 YR Gender:Male I/FU:I

Company Report #USA-2003-0008860

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Abuser	Health Professional	Oxycontin Tablets (Oxycodone Hydrochloride)	PS		
Other		Overdose	Company Representative	Alcohol (Ethanol)	SS		ORAL
40 MG, Q12H,			Other	Neurontin (Gabapentin)	SS		
ORAL				Elavil (Amitriptyline Hydrochloride)	SS		
				Paxil (Paroxetine Hydrochloride)	SS		
				Vicodin (Paracetamol, Hydrocodone Bitartrate)	SS		
				Clonazepam (Clonazepam)	SS		
				Ultram (Tramadol Hydrochloride)	SS		
				Norco (Hydrocodone Bitartrate, Paracetamol)	SS		
				Trazodone (Trazodone)	SS		
				Hydroxyzine Hydrochloride (Hydroxyzine Hydrochloride)	SS		
				Quetiapine (Quetiapine)	SS		
				Metoprolol (Metoprolol)	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Accidental Overdose	Consumer	Oxycodone			
Other		Drug Abuser	Health Professional	Hydrochloride			
			Other	(Similar To Nda 20-553) (Oxycodone Hydrochloride)	PS		ORAL
ORAL				Diazepam (Diazepam)	SS		
				Oxazepam (Oxazepam)	SS		
				Temazepam (Temazepam)	SS		
				Lorazepam (Lorazepam)	SS		
				Cannabnoids	SS		
				Diphenhydramine Hydrochloride			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Diphenhydramine Hydrochloride)	SS
Gabapentin (Gabapentin)	SS
Claritin	C
Zoloft	C
Altace	C
Allopurinol	C
Depakote	C

Date:12/17/03ISR Number: 4271159-4Report Type:Periodic Company Report #KII-2003-003959
 Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other 80 MG, ORAL	Affect Lability Chills Lethargy	Health Professional	Oxycontin Tablets (Oxycodone Hydrochloride)	PS		ORAL
TID, ORAL	Multiple Drug Overdose Tearfulness		Dextromethorphan (Dextromethorphan)	SS		ORAL
6 UNK, ORAL	Tremor		Neurontin (Gabapentin)	SS		ORAL
			Naproxen (Naproxen)	SS		

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 UNKNOWN	Cleft Lip Drug Exposure During Pregnancy		Wellbutrin Neurontin	PS SS	Glaxosmithkline	
			Multivitamins	C		
			Alcohol	C		
			Allegra	C		

Date:12/18/03ISR Number: 4255551-XReport Type:Expedited (15-DaCompany Report #2003117889
 Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	800 MG, ORAL	Back Pain Blood Osmolarity	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Decreased Blood Uric Acid Decreased Dysaesthesia Heat Stroke	Professional	Amoxicillin Trihydrate (Amoxicillin Trihydrate)	SS		ORAL
	ORAL	Hypochloraemia Hyponatraemia Inappropriate Antidiuretic Hormone		Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		ORAL
	ORAL	Secretion Pyrexia		Dantrolene Sodium (Dantrolene Sodium)	SS		ORAL
	75 MG, ORAL			Baclofen (Baclofen)	SS		ORAL
	90 MG, ORAL			Heptaminol Hydrochloride (Heptaminol Hydrochloride)	C		ORAL
	563.4 MG, ORAL			Clonazepam (Clonazepam) Dextropropoxyphene (Dextropropoxyphene)	C C		

Freedom Of Information (FOI) Report

Paracetamol
 (Paracetamol) C
 Eductyl (Sodium
 Bicarbonate,
 Potassium
 Bitartrate) C
 Macrogol (Macrogol) C
 Cefixime (Cefixime) C
 Ceftriaxone
 (Ceftriaxone) C
 Gentamicin
 (Gentamicin) C

Date:12/18/03ISR Number: 4255772-6Report Type:Expedited (15-DaCompany Report #2003113461
 Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Consumer	Neurontin			
		Chest Wall Pain	Health	(Gabapentin)	PS		ORAL
1800 MG (600,		Convulsion	Professional				
TID) ORAL		Eye Rolling		Salbutamol	C		
		Muscle Twitching		Triamcinolone			
		Nervousness		Acetonide	C		
		Sleep Disorder		Peginterferon			
				Alfa-2b	C		
				Ribavirin	C		
				Fentanyl	C		
				Lisinopril	C		
				Atenolol	C		
				Rabeprazole Sodium	C		
				Metoclopramide	C		
				Ondansetron			
				Hydrochloride	C		
				Lidocaine	C		
				Alprazolam	C		
				Escitalopram	C		
				Vicodin			
				(Paracetamol,			
				Hydrocodone			
				Bitartrate)	C		
				Naproxen	C		
				Peginterferon			

Date:12/18/03ISR Number: 4255935-XReport Type:Expedited (15-DaCompany Report #03-000486
Age:37 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Anxiety
Disability	Arthralgia
	Back Pain
	Blood Pressure Increased
	Brain Damage
	Bursitis
	Cerebral Infarction
	Cerebrovascular Accident
	Decreased Appetite
	Depression

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Interaction Dysmenorrhoea Electrocardiogram St-T	Report Source				
		Change Eye Pain Grand Mal Convulsion Haematochezia	Other	Loestrin (Ethinylestradiol, Norethindron Acetate) Tablet	PS		ORAL
SEE IMAGE		Hallucination, Auditory Herpes Simplex		Neurontin Gabapentin)Tablet	SS		ORAL
600 MG, QID,		Insomnia					
ORAL		Irritable Bowel Syndrome Laboratory Test Abnormal Loss Of Consciousness Menstruation Irregular Mitral Valve Incompetence		Oxycodone Hydrochloride(Oxycod one Hydrochloride)Tablet , 20 Mg	SS		ORAL
20 MG, BID,		Oral Pain					
ORAL		Parkinsonism Peroneal Nerve Palsy Psychotic Disorder Road Traffic Accident Schizophrenia, Disorganised Type Urinary Incontinence Weight Increased		Ativan (Lorazepam) Stadol(Butorphanol Tartrate) Paxil (Paroxetine Hydrochloride)	C C C		

Date:12/18/03ISR Number: 4255946-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (BID), Disability		Bipolar Disorder Bone Cyst Cerebral Perfusion	Literature Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other		Pressure Decreased Cognitive Disorder Convulsion Impaired Work Ability	Company Representative	Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride	SS		

Scar
Single Photon Emission
Computerised Tomogram
Abnormal

(Buspirone
Hydrochloride) C
Fluoxetine
Hydrochloride
(Fluoxetine
Hydrochloride) C
Omeprazole
(Omeprazole) C
Capsaicin
(Capsaicin) C

Date:12/19/03ISR Number: 4256439-0Report Type:Expedited (15-DaCompany Report #2003124063
Age:50 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID), ORAL	Abdominal Pain Upper Gastric Disorder Vomiting	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/19/03ISR Number: 4256545-0Report Type:Expedited (15-DaCompany Report #2003123581

Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Breast Cancer	Health	Neurontin			
		Cognitive Disorder	Professional	(Gabapentin)	PS		ORAL
100 MG (DAILY							
AT BEDTIME),		Drug Dependence					
ORAL		Drug Effect Decreased					
		Drug Tolerance Increased		Fentanyl (Fentanyl)	C		
		Dysarthria		Clonazepam			
		Feeling Abnormal		(Clonazepam)	C		
		Food Craving		Celecoxib			
		Neuropathic Pain		(Celecoxib)	C		
		Somnolence					
		Weight Increased					

Date:12/19/03ISR Number: 4256549-8Report Type:Expedited (15-DaCompany Report #2003123204

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Non-Hodgkin'S Lymphoma	Health	Neurontin			
Initial or Prolonged		Pleural Effusion	Professional	(Gabapentin)	PS		ORAL
900 MG (TID),							
Other							
ORAL							
				Fluoxetine			
				Hydrochloride			
				(Fluoxetine			
				Hydrochloride)	C		
				Doxepin (Doxepin)	C		

Date:12/19/03ISR Number: 4256706-0Report Type:Expedited (15-DaCompany Report #2003040394

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Consumer	Neurontin			
		Chest Pain		(Gabapentin)	PS		

Constipation
 Dyspnoea
 Feeling Abnormal
 Hypertension
 Musculoskeletal Stiffness
 Panic Attack
 Treatment Noncompliance

All Other
 Therapeutic Products SS

Date:12/22/03ISR Number: 4257542-1Report Type:Expedited (15-DaCompany Report #2003116234

Age:8 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Consumer	Neurontin (Solution)			
Other		Drug Level Decreased	Health	(Gabapentin)	PS		ORAL
15 CC (TID),		Pharmaceutical Product	Professional				
ORAL		Complaint		Valproic Acid (Valproic Acid)	C		
				Levetiracetam (Levetiracetam)	C		
				All Other Therapeutic Products	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/22/03ISR Number: 4257546-9Report Type:Expedited (15-DaCompany Report #2003118984
Age:71 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID), ORAL	Abdominal Pain Arthritis Blood Glucose Decreased Body Height Decreased Cartilage Injury Drug Hypersensitivity Enzyme Abnormality Fatigue Irritable Bowel Syndrome Vision Blurred	Consumer Health Professional	Neurontin (Gabapentin) Glucosamine Sulfate/Minerals/Mul tivitamins (Glucosamine Sulfate, Vitamins Hyoscyamine Sulfate (Hyoscyamine Sulfate) Pancrelipase (Pancrelipase)	PS SS SS C		ORAL

Date:12/22/03ISR Number: 4257612-8Report Type:Expedited (15-DaCompany Report #2003122499
Age:67 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG Other (TID), ORAL	Alanine Aminotransferase Increased Blood Alkaline Phosphatase Increased Cholestasis Dysarthria Dysphagia Pancreatitis Visual Field Defect	Foreign Health Professional	Neurontin (Gabapentin) Riluzole (Riluzole)	PS C		ORAL

Date:12/23/03ISR Number: 4257168-XReport Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0443704A
Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Hospitalization -	Blood Creatinine	Augmentin	PS	Glaxosmithkline	ORAL
Initial or Prolonged	Increased	Abelcet	SS		
UNKNOWN					
	Jaundice	Neurontin	SS		
UNKNOWN					

Date:12/23/03ISR Number: 4258077-2Report Type:Expedited (15-DaCompany Report #2003124090
Age:81 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 700 MG (TID)	Aphasia Reading Disorder	Foreign Health	Neurontin (Gabapentin)	PS		
	Transient Ischaemic Attack	Professional Company Representative	Aporex (Paracetamol, Dextropropoxyphene Hydrochloride)	C		
			Diclofenac Sodium (Diclofenac Sodium)	C		
			Calciferdiol (Calcifediol)	C		
			Pantoprazole (Pantoprazole)	C		
			Dimeticone (Dimeticone)	C		
			Mebeverine (Mebeverine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/23/03ISR Number: 4258112-1Report Type:Direct
Age: Gender: I/FU:I

Company Report #USP 56215

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
SUSPENSION		Medication Error		Zarontin	PS	Parke-Davis	
SUSPENSION				Gabapentin	SS	Parke-Davis	

Date:12/23/03ISR Number: 4258156-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000697
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (BID), Disability ORAL Other		Brain Neoplasm Diarrhoea Ependymoma Intervertebral Disc Operation Pharyngitis Single Photon Emission Computerised Tomogram Abnormal	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin)	PS SS C C C		ORAL

Date:12/23/03ISR Number: 4258423-XReport Type:Direct
Age:82 YR Gender:Male I/FU:I

Company Report #CTU 208646

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Coordination Abnormal		Gabapentin Terazosin Hcl Clopidogrel	PS C		

Bisulfate	C
Propoxyphene N	
100/Apap	C
Hydrochlorothiazide	C
Irbesartan	C
Rabeprazole Na	C
Paroxetine Hcl	C
Pentoxifylline	C
Felodipine	C

Date:12/23/03ISR Number: 4258456-3Report Type:Expedited (15-DaCompany Report #2003124164

Age: Gender:Female I/FU:I

Outcome	PT
Other	Arthritis
	Asthma
	Blood Cholesterol
	Increased
	Chronic Fatigue Syndrome
	Convulsion
	Drug Tolerance Decreased
	Gingival Swelling
	Hypothyroidism

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Lipoprotein (A) Abnormal Migraine Mitral Valve Prolapse	Consumer	Dilantin (Phenytoin Sodium) Neurontin (Gabapentin)	PS SS		ORAL
900 MG (TID), ORAL		Nerve Injury Oral Discomfort Procedural Complication Restless Legs Syndrome		Clonazepam (Clonazepam) All Other Therapeutic Products Phenobarbital (Phenobarbital)	SS SS C		

Date:12/23/03ISR Number: 4258459-9Report Type:Expedited (15-DaCompany Report #2003114753
Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged Other		Bone Neoplasm Malignant Catheter Related Complication Lung Cancer Metastatic	Consumer	Neurontin (Gabapentin) Omeprazole (Omeprazole) Clonazepam (Clonazepam)	PS C C		

Date:12/24/03ISR Number: 4258983-9Report Type:Direct Company Report #CTU 208774
Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Ear Disorder Feeling Abnormal Inner Ear Disorder Tinnitus		Neurontin Mobic	PS SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Back Pain	Consumer	Humira 40 Mg/0.8 ML			
		Constipation		Pre-Filled Syringe	PS		
SUBCUTANEOUS	40 MG,	1 IN					
2 WK,		Fatigue					
SUBCUTANEOUS		Tremor					
				Gabapentin	SS		
				Prednisone	C		
				Dyazide	C		
				Glaucoma Med	C		
				Latanoprost	C		
				Voltaren	C		
				Gemfibrozil	C		

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
1500 MG (5		Gastrointestinal Haemorrhage	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
DOSES PER		Somnolence	Professional				
DAY), ORAL		Thirst					
				Zoloft (Sertraline)	C		
				Zyrtec (Cetirizine)	C		
				All Other			
				Therapeutic Products	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Rabeprazole Sodium (Rabeprazole Sodium)	C		
				Nadolol (Nadolol)	C		
				Lactulose (Lactulose)	C		
				Becosym Forte (Pyridoxine Hydrochloride, Thiamine Hydrochloride, Folic Acid (Folic Acid)	C		

Date:12/26/03ISR Number: 4259887-8Report Type:Expedited (15-DaCompany Report #2003125235

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiac Arrest	Health	Neurontin			
Other		Ventricular Tachycardia	Professional Company Representative	(Gabapentin) All Other Therapeutic Products	PS C		

Date:12/26/03ISR Number: 4259889-1Report Type:Expedited (15-DaCompany Report #2003119930

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2500 MG, ORAL		Burns Third Degree Intentional Misuse	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other		Loss Of Consciousness Spinal Column Stenosis Suicidal Ideation Suicide Attempt	Professional	Alprazolam (Alprazolam) Escitalopram (Escitalopram) Venlafaxine Hydrochloride (Venlafaxine Hydrochloride) Quetiapine Fumarate (Quetiapine Fumarate)	SS C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/26/03ISR Number: 4259891-XReport Type:Expedited (15-DaCompany Report #2003124353
Age:40 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Blood Creatinine Increased Cholestasis Jaundice	Health Professional	Neurontin (Gabapentin)	PS		
			Amphotericine B, Liposome (Amphotericine B, Liposome)	SS		

Date:12/26/03ISR Number: 4260979-8Report Type:Expedited (15-DaCompany Report #2003118576
Age:67 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID)	Lymphopenia Pyrexia	Foreign Health	Neurontin (Gabapentin)	PS		
	Viral Infection Vomiting	Professional Company Representative	Glimepiride (Glimepiride)	C		
			Zofenopril Calcium (Zofenopril Calcium)	C		
			Clopidogrel (Clopidogrel)	C		
			Fenofibrate (Fenofibrate)	C		
			Fosinopril (Fosinopril)	C		

Date:12/26/03ISR Number: 4260983-XReport Type:Expedited (15-DaCompany Report #2003125090
Age:60 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG (TID)	Hepatic Enzyme Increased Hepatotoxicity	Foreign Health	Neurontin (Gabapentin)	PS		
		Professional Company Representative	Antineoplastic Agents	SS		

Date:12/29/03ISR Number: 4261110-5Report Type:Expedited (15-DaCompany Report #2003122905
Age:19 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Convulsion	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL		Dizziness Nausea	Professional Company Representative	Ergenyl Chrono (Valproic Acid, Valproate Sodium) Lamotrigine (Lamotrigine)	C C		

Date:12/29/03ISR Number: 4261126-9Report Type:Expedited (15-DaCompany Report #2003124063
Age:50 YR Gender:Female I/FU:F

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Abdominal Mass Abdominal Pain Vomiting	Foreign Consumer Health

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
900 MG (TID), ORAL		Neurontin (Gabapentin)	PS		ORAL

Date:12/29/03ISR Number: 4261489-4Report Type:Expedited (15-DaCompany Report #2003120450
Age:49 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID), Other ORAL	Acute Psychosis Catatonia Electroencephalogram Abnormal Hallucinations, Mixed	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL			Quetiapine Fumarate (Quetiapine Fumarate)	SS		ORAL
			Zoloft (Sertraline) Nortriptyline (Nortriptyline) Carbamazepine (Carbamazepine) Morphine Sulfate (Morphine Sulfate) Oestrnorm (Estradiol, Norethisterone) Levothyroxine Sodium (Levothyroxine Sodium) Clonazepam (Clonazepam)	C C C C C C C C		

Date:12/29/03ISR Number: 4261491-2Report Type:Expedited (15-DaCompany Report #2003116907
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chronic Sinusitis Cough	Consumer	Neurontin (Gabapentin)	PS		
900 MG (TID)		Difficulty In Walking Nasopharyngitis Pain In Extremity		Lipitor (Atorvastatin) Rofecoxib (Rofecoxib)	SS SS		
25 MG (DAILY)							

Date:12/29/03ISR Number: 4261673-XReport Type:Direct Company Report #CTU 208908
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 300MG PO TID		Drug Hypersensitivity		Gabapentin 300mg	PS		ORAL
Initial or Prolonged 500MG PO TID		Drug Screen Positive		Etodolac 500mg	SS		ORAL
		Loss Of Consciousness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/29/03ISR Number: 4261680-7Report Type:Expedited (15-DaCompany Report #2003125091

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Eye Operation	Consumer	Neurontin (Gabapentin)	PS		

Date:12/29/03ISR Number: 4261682-0Report Type:Expedited (15-DaCompany Report #2003125240

Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 1800 MG Initial or Prolonged (TID), ORAL Disability Other		Cerebrovascular Accident Drug Ineffective Dysarthria Hyponatraemia Musculoskeletal Disorder Posture Abnormal Refusal Of Treatment By Patient Weight Decreased	Consumer	Neurontin (Tablets) (Gabapentin) Diltiazem Hydrochloride (Diltiazem Hydrochloride) Atenolol (Atenolol) Enalapril Maleate (Enalapril Maleate) Gemfibrozil (Gemfibrozil) Metformin Hydrochloride (Metformin Hydrochloride) Glibenclamide (Glibenclamide)	PS C C C C C C		ORAL

Date:12/30/03ISR Number: 4261775-8Report Type:Expedited (15-DaCompany Report #2003124087

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 400 MG		Infection Lung Disorder	Foreign Health	Neurontin (Gabapentin)	PS		

		Somnolence	Professional			
(DAILY),			Company Representative	All Other Therapeutic Products	C	
Date:12/30/03ISR Number: 4261885-5Report Type:Expedited (15-DaCompany Report #2003113041						
Age:46 YR Gender:Male I/FU:F						
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer
Dose						
Other		Apathy Depression	Consumer Health	Neurontin (Gabapentin)	PS	
2700 MG		Drug Ineffective	Professional			
(TID), ORAL		Drug Interaction		Zoloft (Sertraline)	SS	
2700 MG		Drug Toxicity				
(TID), ORAL		Suicidal Ideation		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS	
(DAILY)				Hydrocodone (Hydrocodone)	C	
				Zolpidem Tartrate (Zolpidem Tartrate)	C	
				Cyclobenzaprine Hydrochloride		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Cyclobenzaprine
Hydrochloride) C
Baclofen (Baclofen) C

Date:12/30/03ISR Number: 4261887-9Report Type:Expedited (15-DaCompany Report #2003125897
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2800 MG, ORAL Other		Condition Aggravated Loss Of Consciousness Medication Error Myocardial Infarction Rheumatoid Arthritis	Consumer	Neurontin (Gabapentin) Sertraline Hydrochloride (Sertraline Hydrochloride)	PS C		ORAL

Date:12/30/03ISR Number: 4261889-2Report Type:Expedited (15-DaCompany Report #2003125631
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other (BID), ORAL		Ageusia Anosmia Somnolence	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:12/30/03ISR Number: 4261892-2Report Type:Expedited (15-DaCompany Report #2003041236
Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 50 MG (PRN), Other ORAL		Anorgasmia Arthralgia Body Height Decreased Drug Effect Decreased	Consumer	Viagra (Sildenafil) Gabapentin (Gabapentin)	PS SS		ORAL
TOPICAL	TOPICAL	Ejaculation Delayed Erectile Dysfunction Exostosis		Atomoxetine Hydrochloride (Atomoxetine)			

Hypoaesthesia	Hydrochloride)	SS
Mental Disorder	Progesterone	
Pain	W/Estrogens/	
Pain In Extremity	(Progesterone,	
Penis Disorder	Estrogen Nos)	SS
Sciatic Nerve Injury	Rofecoxib	
Spinal Compression	(Rofecoxib)	C
Fracture	Esomeprazole	
Spinal Cord Injury	(Esomeprazole)	C
Spinal Deformity		
Spondylolisthesis		
Acquired		
Tendon Disorder		

Date:12/30/03ISR Number: 4262207-6Report Type:Expedited (15-DaCompany Report #2003125947
Age:61 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG	Convulsion Loss Of Consciousness	Foreign Health	Gabapentin (Tablets) (Gabapentin)	PS		ORAL
Other (TID), ORAL	Transient Ischaemic Attack	Professional	Estradiol (Estradiol)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/31/03ISR Number: 4261654-6Report Type:Expedited (15-DaCompany Report #OXCD20030022
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alcohol Use	Literature	Oxycodone	PS		ORAL
Life-Threatening		Cardiac Arrest	Health	Propranolol	SS		ORAL
Hospitalization -		Coma	Professional	Iron	SS		ORAL
Initial or Prolonged		Completed Suicide		Gabapentin	SS		ORAL
		Overdose		Ethanol	SS		ORAL
		Pneumonia Aspiration					
		Pupil Fixed					
		Sepsis					

Date:12/31/03ISR Number: 4262553-6Report Type:Expedited (15-DaCompany Report #2003121603
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Arthralgia	Consumer	Dilantin Suspension			ORAL
		Bone Pain		(Phenytoin Sodium)	PS		ORAL
		Condition Aggravated		Neurontin			ORAL
		Confusional State		(Gabapentin)	SS		ORAL
		Drug Ineffective					
		Epilepsy					
		Fear					
		Impaired Work Ability					
		Myalgia					
		Nervousness					
		Pain In Extremity					
		Personality Disorder					
		Weight Increased					

Date:12/31/03ISR Number: 4262566-4Report Type:Expedited (15-DaCompany Report #2003125945
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams	Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG (BID),		Anger					
ORAL		Psychiatric Symptom					
		Psychotic Disorder		Clonidine (Clonidine)	C		
		Therapeutic Response		Lorazepam (Lorazepam)	C		
		Decreased		Fosinopril Sodium (Fosinopril Sodium)	C		
		Thinking Abnormal					

Date:12/31/03ISR Number: 4262572-XReport Type:Expedited (15-DaCompany Report #2003119634
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Abnormal Behaviour	Health	Neurontin (Gabapentin)	PS		ORAL
Hospitalization -		Drug Interaction	Professional				
ORAL		Feeling Abnormal		Mirtazapine (Mirtazapine)	SS		
Initial or Prolonged		Insomnia		Amitriptyline Hydrochloride			
Other		Intentional Self-Injury		(Amitriptyline Hydrochloride)	SS		
		Mental Disorder		Metaxalone (Metaxalone)	C		
		Restlessness					
		Suicide Attempt					
		Vertigo					

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Freedom Of Information (FOI) Report

Levothyroxine Sodium
 (Levothyroxine Sodium) C
 Axotal (Old Form)
 (Caffeine, Butalbital, Paracetamol) C
 Parafon Forte
 (Chlorzoxazone, Paracetamol) C
 Morphine (Morphine) C
 Tizanidine Hydrochloride
 (Tizanidine Hydrochloride) C
 Oxycocet
 (Paracetamol, Oxycodone Hydrochloride) C
 Estrogens Conjugated
 (Estrogens Conjugated) C
 Quetiapine Fumarate
 (Quetiapine Fumarate) C
 Cyclobenzaprine Hydrochloride
 (Cyclobenzaprine Hydrochloride) C

Date:12/31/03ISR Number: 4262578-0Report Type:Expedited (15-DaCompany Report #2003126467
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Activities Of Daily	Consumer	Neurontin			
Other		Living Impaired		(Gabapentin)	PS		
		Suicide Attempt					

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:12/31/03ISR Number: 4262582-2Report Type:Expedited (15-DaCompany Report #2003113357
Age: Gender:Female I/FU:I

Outcome	PT
Other	Hallucination, Visual Postictal State

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Psychotic Disorder

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG	(ONCE), ORAL	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Company Representative	Phenytoin Sodium (Phenytoin Sodium)	C		
			Hydrochlorothizide (Hydrochlorothiazide)	C		

Date:01/02/04
 Age:43 YR
 Gender:Male
 I/FU:I

ISR Number: 4262951-0
 Report Type:Direct
 Company Report #CTU 209158

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	4200MG QD IN	Depressed Level Of Consciousness		Gabapentin 600mg	PS		ORAL
Intervention to Prevent Permanent Impairment/Damage	3 DO ORAL	Respiratory Depression		Methadone 5mg	SS		ORAL
				Gabapentin	C		
				Ranitidine	C		
				Nefazodone	C		
				Risperdone	C		
				Paroxetine	C		
				Diazepam	C		
				Tizanidine	C		
				Beta-Interferon	C		
				Modafinil	C		
				Methadone	C		
				Zolpidem	C		
				Ibuprofen	C		
				Baclofen	C		
				Zinc	C		
				Vit E	C		
				Vit C	C		
				Docusate	C		
				Milk Of Magnesia	C		

Date:01/02/04ISR Number: 4262975-3Report Type:Direct Company Report #CTU 209200
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Alopecia		Gabapentin	PS		ORAL
600-900MG PO							
HS + 300 MG							
Q8H PRN							

Date:01/02/04ISR Number: 4263636-7Report Type:Expedited (15-DaCompany Report #2003119508
Age:31 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Contusion
Initial or Prolonged	Depressed Mood
Other	Euphoric Mood
	Fall
	Judgement Impaired
	Mood Altered
	Multiple Drug Overdose
	Muscle Spasms

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pain

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
900 MG (TID),		Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Company Representative	Diazepam (Diazepam)	SS		
ORAL			Tramadol (Tramadol)	SS		ORAL
ORAL			Temazepam (Temazepam)	SS		ORAL
			Diamorphine (Diamorphine)	C		
			Alimemazine Tartrate (Alimemazine Tartrate)	C		
			Indometacin (Indometacin)	C		
			Panadeine Co (Codeine Phosphate, Paracetamol)	C		
			Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		
			Alimemazine Tartrate	C		

Date:01/02/04ISR Number: 4264515-1Report Type:Expedited (15-DaCompany Report #2003126141

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Diarrhoea	Consumer	Neurontin (Gabapentin)	PS		ORAL
3600 MG		Drug Dependence					
(TID), ORAL		Drug Withdrawal Syndrome					
		Medication Error		Diazepam (Diazepam)	C		
				Venlafaxine Hydrochloride (Venlafaxine			

Hydrochloride) C
 Fluoxetine
 Hydrochloride
 (Fluoxetine
 Hydrochloride) C

Date:01/02/04ISR Number: 4264520-5Report Type:Expedited (15-DaCompany Report #2003121603
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Consumer	Dilantin Suspension (Phenytoin Sodium)	PS		ORAL
ORAL		Bone Pain					
		Confusional State		Neurontin (Gabapentin)	SS		ORAL
1200 MG		Convulsion					
(QID),ORAL		Drug Ineffective					
		Myalgia					
		Pain In Extremity					
		Personality Disorder					
		Social Avoidant Behaviour					
		Unemployment					
		Weight Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/02/04ISR Number: 4264740-XReport Type:Expedited (15-DaCompany Report #2003123581
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Breast Cancer Stage I Cognitive Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
100 MG (DAILY AT BEDTIME) , ORAL		Drug Dependence Drug Tolerance Dysarthria Feeling Abnormal Food Craving Hyperphagia Neuropathic Pain Somnolence Weight Increased		Fentanyl (Fentanyl) Clonazepam (Clonazepam) Celecoxib (Celecoxib)	C C C		

Date:01/02/04ISR Number: 4264742-3Report Type:Expedited (15-DaCompany Report #2003125621
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Discomfort Drug Dependence	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL		Drug Ineffective Drug Level Decreased Dyspepsia Fear Hallucination Treatment Noncompliance		Bismuth Subsalicylate (Bismuth Subsalicylate) Morphine Sulfate (Morphine Sulfate) Citalopram (Citalopram) Quetiapine Fumarate (Quetiapine Fumarate) Topiramate (Topiramate) Vicodin (Paracetamol,	SS C C C		

Hydrocodone
Bitartrate) C
Oxycodone
Hydrochloride
(Oxycodone
Hydrochloride) C

Date:01/02/04ISR Number: 4264745-9Report Type:Expedited (15-DaCompany Report #2003126148

Age: Gender:Female I/FU:I

Outcome PT
Disability Aggression
Other Anger
Condition Aggravated
Diarrhoea
Drug Withdrawal Syndrome
Feeling Abnormal
Hallucination
Heart Rate Abnormal
Homicidal Ideation
Hostility
Irritable Bowel Syndrome

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3600 MG (TID)		Muscle Spasms Nausea Paraesthesia Physical Assault Suicidal Ideation Vomiting	Consumer	Neurontin (Gabapentin)	PS		
				Diazepam (Diazepam)	C		
				Alprazolam (Alprazolam)	C		
				Valproate Semisodium (Valproate Semisodium)	C		

Date:01/02/04ISR Number: 4264747-2Report Type:Expedited (15-DaCompany Report #2003126147
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anger Diarrhoea	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Drug Dependence Drug Withdrawal Syndrome Faeces Discoloured Feeling Abnormal Heart Rate Abnormal Homicidal Ideation Intentional Self-Injury Muscle Spasms Nausea Oesophageal Disorder Pain Paraesthesia Physical Assault Suicidal Ideation Vomiting					

Date:01/03/04ISR Number: 4544162-4Report Type:Direct Company Report #CTU 235442
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other			Anxiety	Neurontin 300 Mg		
			Drug Ineffective	(Generic)	PS	ORAL
PO 300 MG (2)						
			Hangover			
QHS	2	WK	Insomnia			
			Pharmaceutical Product			
			Complaint			

Date:01/05/04ISR Number: 4263821-4Report Type:Direct Company Report #CTU 209300
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Herpes Simplex		Gabapentin			
		Mood Altered		(Neurontin)	PS		ORAL
300 MG PO 7							
DAYS, THEN		Sexual Dysfunction					
300 MG, TID 7							
DAYS		Stomatitis					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/06/04ISR Number: 4264995-1Report Type:Expedited (15-DaCompany Report #WAES 0312FRA00090

Age:80 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Pneumonitis	Health	Fosamax	PS	Merck & Co., Inc	ORAL
Initial or Prolonged	Pulmonary Fibrosis	Professional	Ergoloid Mesylates	SS		ORAL
			Gabapentin	SS		ORAL

2 MON

Date:01/06/04ISR Number: 4265495-5Report Type:Expedited (15-DaCompany Report #2003126804

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Contusion	Consumer	Neurontin			
900 MG	Coordination Abnormal		(Gabapentin)	PS		ORAL
(DAILY), ORAL	Difficulty In Walking					
	Drug Toxicity		Fentanyl (Fentanyl)	C		
	Fall		Fluoxetine			
	Head Injury		Hydrochloride			
	Pain In Extremity		(Fluoxetine			
	Speech Disorder		Hydrochloride)	C		
	Treatment Noncompliance		Bupropion			
			Hydrochloride			
			(Bupropion			
			Hydrochloride)	C		
			Fexofenadine			
			Hydrochloride			
			(Fexofenadine			
			Hydrochloride)	C		
			Pantoprazole			
			(Pantoprazole)	C		
			Oxycodone			
			Hydrochloride			
			(Oxycodone			
			Hydrochloride)	C		
			Cyclobenzaprine			
			Hydrochloride			
			(Cyclobenzaprine			
			Hydrochloride)	C		
			Rofecoxib			
			(Rofecoxib)	C		

Trazodone
(Trazodone) C
Modafinil
(Modafinil) C

Date:01/06/04ISR Number: 4265505-5Report Type:Expedited (15-DaCompany Report #2003126805
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence	Health Professional	Neurontin (Gabapentin)	PS		

Date:01/06/04ISR Number: 4266022-9Report Type:Expedited (15-DaCompany Report #HCDA20030010
Age:41 YR Gender:Female I/FU:I

Outcome	PT
Death	Blood Glucose Increased Blood Magnesium Increased Blood Sodium 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
		Body Temperature Decreased					
		Brain Death					
2-4 TABS PO		Cardiac Failure	Literature	Hydrocodone/Apap	PS		ORAL
PO		Cardio-Respiratory Arrest	Health	Fentanyl Patch	SS		ORAL
2-4 TABS PO		Coma	Professional	Carisoprodol	SS		ORAL
2 TABS PO		Drug Ineffective		Clonazepam	SS		ORAL
PO		Electrocardiogram St Segment Elevation		Gabapentin	SS		ORAL
PO		Electroencephalogram		Cyclobenzaprine	SS		ORAL
PO		Abnormal		Tolterodine	SS		ORAL
PO		Haemoglobin Decreased		Buspirone	SS		ORAL
PO		Heart Rate Increased		Doxycycline	SS		ORAL
		Hepatic Failure					
		Livedo Reticularis					
		Medication Error					
		Multi-Organ Failure					
		Multiple Drug Overdose					
		Rectal Haemorrhage					
		Renal Failure					
		Suicide Attempt					

Date:01/06/04ISR Number: 4266084-9Report Type:Expedited (15-DaCompany Report #2002051354
 Age:77 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asterixis	Foreign	Gabapentin			
3600 MG		Blood Glucose Increased	Literature	(Gabapentin)	PS		
Other (DAILY)		Cognitive Disorder					
		Confusional State		Sertraline			
		Decreased Activity		(Sertraline)	C		
		Disturbance In Attention					

Encephalopathy
 Lacunar Infarction
 Memory Impairment
 Mental Disorder
 Personality Change
 Sensory Loss

Date:01/06/04ISR Number: 4266163-6Report Type:Expedited (15-DaCompany Report #KII-2003-0006601
 Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Disorientation Drug Interaction Mental Status Changes Pain Respiratory Rate Decreased Sinus Tachycardia Somnolence	Study Health Professional Other	Oxycodone Hydrochloride Fentanyl (Fentanyl) Amitriptyline (Amitriptyline) Ssri () Baclofen (Baclofen) Gabapentin (Gabapentin) Tramadol (Tramadol) Cr Tablet Tolterodine (Tolterodine)			PS SS SS SS SS SS SS

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/06/04ISR Number: 4266245-9Report Type:Expedited (15-DaCompany Report #2003126600

Age:31 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID), ORAL	Abnormal Behaviour Diarrhoea Euphoric Mood Feeling Abnormal Judgement Impaired Nausea Overdose Self-Medication	Foreign Health Professional	Neurontin (Gabapentin) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Tramadol (Tramadol) Temazepam (Temazepam) Alimemazine Tartrate (Alimemazine Tartrate) Indometacin (Indometacin) Panadeine Co (Codeine Phosphate, Paracetamol)	 C C C C C C		

Date:01/06/04ISR Number: 4266252-6Report Type:Expedited (15-DaCompany Report #2002068598

Age:30 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (THREE TIMES DAILY), ORAL	Arthralgia Asthma Bronchitis Acute Dizziness Drug Interaction Inflammation Nasopharyngitis Pain Pyrexia	Foreign Study Health Professional	Gabapentin (Gabapentin) Carbamazepine (Carbamazepine) Clobazam (Clobazam) Theophylline (Theophylline) Pranlukast (Pranlukast)	 PS C C C C		ORAL

Salbutamol	
(Salbutamol)	C
Budesonide	
(Budesonide)	C
Tulobuterol	
Hydrochloride	
(Tulobuterol	
Hydrochloride)	C
Methylprednisolone	
Sodium Succinate	
(Methylprednisolone	
Sodium Succinate)	C
Aminophylline	
(Aminophylline)	C
Actit (Maltose,	
Sodium Acetate,	
Potassium Chloride,	
Sodium Chloride,	
Potassium Phosphate	C
Solita T	
(Electrolytes Nos)	C
Cefdinir (Cefdinir)	C
Acemetacin	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Acemetacin)	C
Pentoxyverine	
(Pentoxyverine)	C
Sodium Chloride	
(Sodium Chloride)	C
Bromhexine	
Hydrochloride	
(Bromhexine	
Hydrochloride)	C
Procaterol	
Hydrochloride	
(Procaterol	
Hydrochloride)	C
Hydrocortisone	
Sodium Succinate	
(Hydrocortisone	
Sodium Succinate)	C
Soldem 3a (Sodium	
Lactate, Potassium	
Chloride, Sodium	
Chloride,	
Carbohydrates Nos)	C
Diazepam (Diazepam)	C
Magnesium Oxide	
(Magnesium Oxide)	C
Cefcapene Pivoxil	
Hydrochloride	
(Cefcapene Pivoxil	
Hydrochloride)	C

Date:01/06/04ISR Number: 4266286-1Report Type:Expedited (15-DaCompany Report #2002051352
 Age:76 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Asterixis	Foreign	Gabapentin			
Initial or Prolonged		Literature	(Gabapentin)	PS		
900 MG						
Other						
(DAILY),						
			Oxcarbazepine			
			(Oxcarbazepine)	SS		
			Lacidipine			
			(Lacidipine)	C		
			Acetylsalicylic Acid			
1200 MG (BID)						

(Acetylsalicylic
Acid) C

Date:01/07/04ISR Number: 4267043-2Report Type:Expedited (15-DaCompany Report #2003127209
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Body Height Decreased	Consumer	Neurontin (Tablets)			
		Loss Of Consciousness		(Gabapentin)	PS		ORAL
3200 MG							
(DAILY), ORAL		Pain					
				Carisoprodol			
				(Carisoprodol)	C		
				Diazepam (Diazepam)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/07/04ISR Number: 4267097-3Report Type:Expedited (15-DaCompany Report #2012272

Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Consumer	Oxycontin			
Hospitalization -		Back Pain	Health	Tablets(Oxycodone			
Initial or Prolonged		Drug Abuser	Professional	Oxycontin			
Other		Haematemesis	Other	Tablets(Oxycodone			
SEE IMAGE		Headache		Hydrochloride) Cr	PS		
		Respiratory Arrest		Diazepam (Diazepam)	SS		
				Oxazepam (Oxazepam)	SS		
				Temazepam			
				(Temazepam)	SS		
				Lorazepam			
				(Lorazepam)	SS		
				Cannabnoids			
				(Cannabis)	SS		
				Diphenhydramine			
				Hydrochloride			
				(Diphenhydramine			
				Hydrochloride)	SS		
				Gabapentin			
				(Gabapentin)	SS		
				Claritin			
				(Loratadine)	C		
				Zoloft (Sertraline			
				Hydrochloride)	C		
				Altace (Ramipril)	C		
				Allopurinol			
				(Allopurinol)	C		
				Depakote (Valproate			
				Semisodium)	C		
				Soma (Carisoprodol)	C		
				Relafen (Nabumetone)	C		

Date:01/07/04ISR Number: 4267359-XReport Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Back Pain	Consumer	Neurontin			
Initial or Prolonged		Brain Neoplasm	Health	(Gabapentin)	PS		ORAL
400 MG (BID),							

Disability	Cardiac Arrest	Professional		
ORAL				
Other	Conversion Disorder	Company	Valproate Sodium	
	Disease Recurrence	Representative	(Valproate Sodium)	SS
	Dizziness		Buspirone	
	Dysphemia		Hydrochloride	
	Ependymoma		(Buspirone	
	Fatigue		Hydrochloride)	C
	Hypoaesthesia		Fluoxetine	
	Intervertebral Disc		Hydrochloride	
	Operation		(Fluoxetine	
	Memory Impairment		Hydrochloride)	C
	Nervous System Disorder		Omeprazole	
	Post Procedural		(Omeprazole)	C
	Complication		Capsaicin	
	Somnolence		(Capsaicin)	C
	Speech Disorder			
	Syncope			
	Tremor			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/08/04ISR Number: 4268836-8Report Type:Expedited (15-DaCompany Report #2003117320

Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness	Health	Neurontin			
		Somnolence	Professional	(Gabapentin)	PS		ORAL
1200 MG							
(TID), ORAL							
				Propacet			
				(Paracetamol,			
				Dexpropoxyphene			
				Napsilate)	C		
				Oxycocet			
				(Paracetamol,			
				Oxycodone			
				Hydrochloride)	C		
				Valdecoxib			
				(Valdecoxib)	C		
				Nitrofurantoin			
				(Nitrofurantoin)	C		
				Cyclobenzaprine			
				Hydrochloride			
				(Cyclobenzaprine			
				Hydrochloride)	C		
				Alprazolam			
				(Alprazolam)	C		
				Librax			
				(Chlordiazepoxide			
				Hydrochloride,			
				Clidinium Bromide)	C		

Date:01/09/04ISR Number: 4268982-9Report Type:Expedited (15-DaCompany Report #2003121603

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Consumer	Dilantin Suspension			
		Bone Pain		(Phenytoin Sodium)	PS		ORAL
ORAL							
		Confusional State		Neurontin			
		Convulsion		(Gabapentin)	SS		ORAL
1200 MG							

(QID), ORAL

Drug Effect Decreased

Epilepsy
Myalgia
Pain In Extremity
Tremor
Weight Increased

Date:01/09/04ISR Number: 4269289-6Report Type:Expedited (15-DaCompany Report #2003127136

Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Disturbance In Attention Hallucination	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Musculoskeletal Stiffness Sedation Vertigo Positional Vomiting	Professional	All Other Non-Therapeutic Products	C		

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Freedom Of Information (FOI) Report

Date:01/09/04 ISR Number: 4269633-X Report Type:Expedited (15-Da Company Report #2004000039
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache	Consumer	Neurontin			
Other		Neoplasm		(Gabapentin)	PS		
		Photophobia					
		Tooth Disorder					

Date:01/12/04 ISR Number: 4269620-1 Report Type:Expedited (15-Da Company Report #2003127135
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3200 MG		Asthenia Diarrhoea	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other (QID), ORAL		Dizziness	Professional				
		Oral Intake Reduced		Allopurinol (Allopurinol)	C		
		Paraesthesia		Karvea Hct (Hydrochlorothiazide , Irbesartan)	C		
		Tremor		Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
		Vomiting		Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Fluvastatin Sodium (Fluvastatin Sodium)	C		
				Primidone (Primidone)	C		
				Novothyral (Levothyroxine Sodium, Liothyronine Sodium)	C		
				Ranitidine Hydrochloride (Ranitidine Hydrochloride)	C		

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acidosis	Literature	Glipizide	PS		ORAL
PO							
Life-Threatening		Bradycardia	Health	Metformin	SS		ORAL
PO							
Hospitalization -		Completed Suicide	Professional	Acarbose	SS		ORAL
PO							
Initial or Prolonged		Diarrhoea		Terazosin	SS		ORAL
PO							
		Hypoglycaemia		Lisinopril	SS		ORAL
PO							
		Hypotension		Gabapentin	SS		ORAL
PO							
		Hypothermia		Hydroxyzine	SS		ORAL
PO							
		Lactic Acidosis		Dipyridamole	SS		ORAL
PO							
		Lethargy		Lovastatin	SS		ORAL
PO							
		Overdose		Finasteride	SS		ORAL
PO							

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Freedom Of Information (FOI) Report

Date:01/12/04ISR Number: 4269813-3Report Type:Expedited (15-DaCompany Report #2003126500
Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG (BID) Other	Blood Insulin Increased Blood Parathyroid Hormone Increased Decreased Appetite Dizziness Fatigue Hypoglycaemia Hypotension	Literature Health Professional	Gabapentin (Gabapentin) Sevelamer (Sevelamer) Lansoprazole (Lansoprazole) Gemfibrozil (Gemfibrozil)	PS C C C		

Date:01/12/04ISR Number: 4269880-7Report Type:Expedited (15-DaCompany Report #US-SHR-03-012130
Age:53 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged SUBCUTANEOUS Other EVERY 2 D, SUBCUTANEOUS SEE IMAGE	Dizziness Liver Function Test 5.6 MIU, Abnormal Pain Tremor	Consumer	Betaseron(Interferon Beta - 1b) Injection Prednisone (Prednisone) Neurontin (Gabapentin)	PS SS SS		
			Prempro (Estrogens Conjugated) Flomax (Morniflumate)	C C		

Date:01/13/04ISR Number: 4269547-5Report Type:Expedited (15-DaCompany Report #PHEH2004US00689
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Other	Amnesia	Trileptal	PS	Novartis Sector:
	Balance Disorder			Pharma
600 mg, UNK				
	Crying	Neurontin	SS	
3000 mg, UNK				
	Depressed Mood			
	Fall			
	Nausea			
	Screaming			
	Visual Disturbance			

Date:01/13/04ISR Number: 4271666-4Report Type:Expedited (15-DaCompany Report #2004000582
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (TID), ORAL		Stevens-Johnson Syndrome Toxic Skin Eruption	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Omeprazole (Omeprazole)	C		
				Metoprolol Tartrate (Metoprolol Tartrate)	C		
				Alprazolam (Alprazolam)	C		
				Atenolol (Atenolol)	C		
				Fluvastatin			

Freedom Of Information (FOI) Report

(Fluvastatin) C
 Endotelon (Vitis
 Vinifera, Herbal
 Extracts Nos) C
 Methylprednisolone
 Sodium Succinate
 (Emthylprednisolone
 Sodium Succinate) C

Date:01/13/04ISR Number: 4271668-8Report Type:Expedited (15-DaCompany Report #2004001239
 Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myalgia Sudden Death	Foreign Health	Gabapentin(Gabapenti n)	PS		ORAL
ORAL			Professional				

Date:01/13/04ISR Number: 4271986-3Report Type:Expedited (15-DaCompany Report #2003124090
 Age:81 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (BID),		Acalculia Agraphia Alexia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Professional				
			Company Representative	Aporex (Paracetamol, Dextropropoxyphene Hydrochloride) Diclofenac Sodium (Diclofenac Sodium) Calcifeiol (Calcifediol) Pantoprazole (Pantoprazole) Dimeticone (Dimeticone) Mebeverine (Mebeverine)	C C C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cholestasis	Foreign	Neurontin			
		Hypertension	Health	(Gabapentin)	PS		ORAL
300 MG (QID),							
ORAL		Hypotension	Professional				
		Loss Of Consciousness		Amitriptyline			
		Vertigo		Hydrochloride			
				(Amitriptyline			
				Hydrochloride)	SS		
35 MG (BID)				Atenolol (Atenolol)	SS		ORAL
100 MG							
(DAILY), ORAL				Nifedipine			
				(Nifedipine)	SS		ORAL
30 MG							
(DAILY), ORAL				Lisinopril			
				(Lisinopril)	SS		ORAL
5 MG (DAILY),							
ORAL				Sodium Polystyrene			
				Sulfonate (Sodium			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Polystyrene
Sulfonate) C
Cyanocobalamin
(Cyanocobalamin) C
Thiamine
Hydrochloride
(Thiamine
Hydrochloride) C
Pyridoxine
(Pyridoxine) C

Date:01/14/04ISR Number: 4271266-6Report Type:Direct
Age:48 YR Gender:Male I/FU:I

Company Report #CTU 210058

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Deafness		Neurontin 300 Mg Tid	PS		ORAL
300 MG PO TID		Hearing Impaired		Diclofenac	C		

Date:01/14/04ISR Number: 4272259-5Report Type:Expedited (15-DaCompany Report #2003110336
Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Balance Disorder Cerebellar Syndrome Coordination Abnormal	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:01/14/04ISR Number: 4272261-3Report Type:Expedited (15-DaCompany Report #2003117658
Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Face Oedema	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
SEE IMAGE			Professional Company Representative	Magnesium Pidolate (Magnesium Pidolate) Pantoprazole Sodium	C		

(Pantoprazole Sodium) C
 Folic Acid (Folic Acid) C
 Cyanocobalamin (Cyanocobalamin) C

Date:01/14/04ISR Number: 4272263-7Report Type:Expedited (15-DaCompany Report #2003119973
 Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG Disability (TID), ORAL		Post Thrombotic Syndrome Venous Thrombosis Limb	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Nitrendipine (Nitrendipine)	C		
				Insulin Lispro (Insulin Lispro)	C		
				Ramipril (Ramipril)	C		
				Torasemide (Torasemide)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Heparin-Fraction,
 Calcium Salt
 (Heparin Fraction,
 Calcium Salt) C
 Clindamycin
 Hydrochloride
 (Clindamycin
 Hydrochloride) C
 Pentoxifylline
 (Pentixifylline) C

Date:01/14/04ISR Number: 4272821-XReport Type:Expedited (15-DaCompany Report #2003016415
 Age:21 YR Gender: I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL				Professional	Haloperidol (Haloperidol)	SS		ORAL
ORAL					Benzatropine Mesilate (Benzatropine Mesilate)	SS		ORAL

Date:01/14/04ISR Number: 4272843-9Report Type:Expedited (15-DaCompany Report #2003119747
 Age:15 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	300 MG		Intentional Misuse Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other (DAILY), ORAL			Vomiting		Paracetamol (Paracetamol)	SS		ORAL
ORAL					Acetylcysteine (Acetylcysteine)	SS		

Celexa (Citalopram
Hydrobromide) C

Date:01/14/04ISR Number: 4272844-0Report Type:Expedited (15-DaCompany Report #2004001234
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pruritus Generalised	Health	Neurontin			
ORAL		Rash	Professional	(Gabapentin)	PS		ORAL
		Skin Disorder		All Other			
				Therapeutic Products	C		
				Vitamins	C		
				Estrogens			
				Conjugated			
				(Estrogens			
				Conjugated)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/04ISR Number: 4272901-9Report Type:Expedited (15-DaCompany Report #2004001236
Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG Other (TID), ORAL		Asthenia Bacteria Sputum Identified	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Cardiomegaly Chlamydial Infection		Dihydroerogotoxine (Dihydroergotoxine)	SS		ORAL
ORAL		Diastolic Dysfunction Dyspnoea Exertional		Alendronate Sodium (Alendronate Sodium)	SS		ORAL
		Headache Inflammation Interstitial Lung Disease Pulmonary Embolism Pulmonary Fibrosis Serology Positive					

Date:01/15/04ISR Number: 4274262-8Report Type:Expedited (15-DaCompany Report #2004001095
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other UNKNOWN		Acute Coronary Syndrome Amputation	Consumer	Neurontin (Gabapentin)	PS		
		Angina Unstable Atrioventricular Block Cardiac Valve Sclerosis Cardio-Respiratory Arrest Cardiomegaly Dehydration Drug Toxicity Gangrene Gastroenteritis Myalgia Osteomyelitis Peripheral Vascular Disorder Pneumonia		Valsartan (Valsartan) Amlodipine (Amlodipine) Hydrochlorothiazide (Hydrochlorothiazide) Fentanyl (Fentanyl) Vicodin (Paracetamol, Hydrocodone Bitartrate) Cilostazol (Cilostazol)	C C C C C C		

Skin Disorder
Skin Ulcer

Acetylsalicylic Acid
(Acetylsalicylic
Acid) C
Metoclopramide
(Metoclopramide) C
Furosemide
(Furosemide) C
Esomeprazole
(Esomeprazole) C
Novolin 20/80
(Insulin Human,
Insulin Isophane,
Human Biosynthetic) C
Insulin (Insulin) C
Metoprolol
(Metoprolol) C
Docusate (Docusate) C
Chromagen (Ascorbic
Acid,
Cyanocobalamin,
Ferrous Fumarate,
Stomach Desiccated) C
Peri-Colace

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(Docusate Sodium,
Casanthranol) C

Date:01/15/04ISR Number: 4274298-7Report Type:Expedited (15-DaCompany Report #2003125621
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Daydreaming	Consumer	Neurontin			
		Drug Ineffective	Health	(Gabapentin)	PS		ORAL
900 MG (TID),							
ORAL		Drug Interaction	Professional				
		Drug Level Decreased		Bismuth Subalicylate			
		Drug Withdrawal Syndrome		(Bismuth			
		Dyspepsia		Subsalicylate)	SS		
		Hallucination, Visual		Morphine Sulfate			
		Medication Error		(Morphine Sulfate)	C		
		Mental Disorder		Citalopram			
				(Citalopram)	C		
				Quetiapine Fumarate			
				(Quetiapine			
				Fumarate)	C		
				Topiramate			
				(Topiramate)	C		
				Vicodin			
				(Paracetamol,			
				Hydrocodone			
				Bitartrate)	C		
				Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	C		

Date:01/16/04ISR Number: 4275310-1Report Type:Expedited (15-DaCompany Report #KII-2003-0006846
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -							
Initial or Prolonged		Confusional State	Study	Oxycodone			
		Drug Withdrawal Syndrome	Health	Hydrochloride			
		Flushing	Professional	(Similar To Nda			
		Hyperhidrosis	Other	20-553)(Oxycodone			

ORAL	Hypertension	Hydrochloride)	PS	ORAL
	Mental Status Changes	Morphine Sulfate		
	Overdose	(Similar No		
	Somnolence	Nda19-516) (Morphine		
	Tachycardia	Sulfate) Unknown	SS	ORAL
ORAL		Baclofen (Baclofen)	SS	ORAL
ORAL		Neurontin		
		(Gabapentin)	SS	
		Zinc (Zinc)	SS	
		Hydrochlorothiazide		
		(Hydrochlorothiazide		
) Tablet	SS	
		Paxil ((Paroxetine		
		Hydrochloride)	SS	
		Vitamin A (Retinol)	SS	
		Vitamin C (Ascorbic		
		Acid)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/16/04ISR Number: 4275402-7Report Type:Expedited (15-DaCompany Report #2004001231

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disturbance In Attention	Health	Neurontin			
		Suicidal Ideation	Professional	(Gabapentin)	PS		ORAL
2400 MG		Weight Increased					
(QID), ORAL							
				Prednisone			
				(Prednisone)	C		
				Clopidogrel Sulfate			
				(Clopidogrel			
				Sulfate)	C		
				Dipyridamole			
				(Dipyridamole)	C		
				Diltiazem (
				Diltiazem)	C		
				Omeprazole			
				(Omeprazole)	C		
				Hydrochlorothiazide			
				(Hydrochlorothiazide)	C		
				Hydroxyzine			
				Hydrochloride			
				(Hydroxyzine			
				Hydrochloride)	C		
				Progesterone			
				(Progesterone)	C		
				Metoprolol			
				(Metoprolol)	C		
				Fludrocortisone			
				Acetate			
				(Fludrocortisone			
				Acetate)	C		
				Zolpidem Tartrate			
				(Zolpidem Tartrate)	C		
				Fentanyl (Fentanyl)	C		

Date:01/16/04ISR Number: 4275412-XReport Type:Expedited (15-DaCompany Report #2003117242

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Optic Neuritis	Consumer	Neurontin			

Vision Blurred

(Gabapentin)

PS

Date:01/16/04ISR Number: 4297231-0Report Type:Direct
Age:48 YR Gender:Female I/FU:I

Company Report #USP 56317

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Medication Error		Augmentin Xr	PS	Glaxosmith Kline	
TABLETS				Neurontin 600 Mg	SS	Park Davis	
CAPSULES							

Date:01/20/04ISR Number: 4274471-8Report Type:Expedited (15-DaCompany Report #US-ROCHE-315823
Age:71 YR Gender:Female I/FU:F

Outcome	PT
Other	Cataract Creatinine Renal Clearance Decreased Dry Mouth Dysphonia

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Keratoconjunctivitis Sicca Laboratory Test Abnormal	Consumer	Klonopin	PS	Roche	ORAL
		Palpitations	Consumer	Neurontin	SS		ORAL
		Tardive Dyskinesia		Lamictal	SS		ORAL
		Weight Increased		Diavan	C		ORAL
				Vitamins	C		ORAL

Date:01/20/04ISR Number: 4276382-0Report Type:Expedited (15-DaCompany Report #2004001696
Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cataract Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG (TID), ORAL		Hypoaesthesia		Telmisartan (Telmisartan)	C		
		Pain		Esomeprazole (Esomeprazole)	C		
		Paraesthesia		Atorvastatin (Atorvastatin)	C		
		Post Procedural Complication		Acetylsalicylic Acid (Acetylsalicylic Acid)	C		

Date:01/20/04ISR Number: 4276488-6Report Type:Expedited (15-DaCompany Report #2004001682
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Aneurysm Aortic Aneurysm	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL				Dilantin Suspension (Phenytoin Sodium)	SS		

Date:01/20/04ISR Number: 4276500-4Report Type:Expedited (15-DaCompany Report #2004001906
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Anorgasmia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL		Caesarean Section Dizziness Drug Effect Decreased Feeling Abnormal Medication Error Panic Disorder Pregnancy Sexual Abuse Somnolence					

Date:01/20/04ISR Number: 4276910-5Report Type:Expedited (15-DaCompany Report #2004001690
Age:75 YR Gender:Male I/FU:I

Outcome	PT
Other	Bronchitis Confusional State

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Delusion Dry Mouth Dry Throat	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Dysphagia Hallucination					
		Pneumonia Pyrexia		Heparin-Fraction, Sodium Salt (Heparin-Fraction, Sodium Salt)	C		
				Erythropoietin (Erythropoietin)	C		
				Lorazepam (Lorazepam)	C		
				Amantadine Hydrochloride (Amantadine Hydrochloride)	C		

Date:01/21/04ISR Number: 4277610-8Report Type:Expedited (15-DaCompany Report #2003116124
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (DAILY), ORAL		Abnormal Dreams Angina Pectoris Feeling Abnormal Hallucination Heart Rate Increased Suicidal Ideation Tremor	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:01/21/04ISR Number: 4277611-XReport Type:Expedited (15-DaCompany Report #2004001757
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Electromyogram Abnormal Liver Function Test Abnormal	Health Professional Company	Neurontin (Gabapentin) Panadeine Co	PS		

Respiratory Arrest

Representative

(Codeine Phosphate,
Paracetamol) C
Amitriptyline
Hydrochloride
(Amitriptyline
Hydrochloride) C

Date:01/21/04ISR Number: 4278021-1Report Type:Expedited (15-DaCompany Report #2004001909

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (TID), ORAL		Intentional Misuse Suicidal Ideation Suicide Attempt	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Trazodone Citalopram Hydrobromide (Citalopram Hydrobromide) Methylphenidate Hydrochloride	SS C		

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Freedom Of Information (FOI) Report

(Methylphenidate
Hydrochloride) C

Date:01/22/04ISR Number: 4278865-6Report Type:Expedited (15-DaCompany Report #2012272

Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Consumer	Oxycontin Tablets			
Hospitalization - Initial or Prolonged		Apnoea	Health	(Oxycodone			
Other		Back Pain	Professional	Hydrochloride) Cr			
SEE IMAGE		Drug Abuser	Other	Tablet	PS		
		Haematemesis		Diazepam (Diazepam)	SS		
		Headache		Oxazepam (Oxazepam)	SS		
		Polysubstance Abuse		Temazepam			
				(Temazepam)	SS		
				Lorazepam			
				(Lorazepam)	SS		
				Cannabis (Cannabis)	SS		
				Diphenhydramine			
				Hydrochloride			
				(Diphenhydramine			
				Hydrochloride)	SS		
				Gabapentin			
				(Gabapentin)	SS		
				Claritin			
				(Loratadine)	C		
				Zoloft (Sertraline			
				Hydrochloride)	C		
				Altace (Ramipril)	C		
				Allopurinol			
				(Allopurinol)	C		
				Depakote (Valproate			
				Semisodium)	C		
				Soma (Carisoprodol)	C		

Date:01/22/04ISR Number: 4279044-9Report Type:Expedited (15-DaCompany Report #2003112401

Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alanine Aminotransferase	Health	Neurontin			

200 MG (BID)

Increased	Professional	(Gabapentin)	PS
Aspartate	Company	Lamotrigine	
Aminotransferase	Representative	(Lamotrigine)	SS
Increased		Alprazolam	
Biopsy Liver Abnormal		(Alprazolam)	C
Blood Alkaline		Phenelzine Sulfate	
Phosphatase Increased		(Phenelzine Sulfate)	C
Liver Function Test		Olanzapine	
Abnormal		(Olanzapine)	C
		Estradiol	
		(Estradiol)	C
		Estrogens Conjugated	
		(Estrogens	
		Conjugated)	C
		Medroxyprogesterone	
		Acetate	
		(Medroxyprogesterone	
		Acetate)	C
		Guaifenesin	
		(Guaifenesin)	C

Freedom Of Information (FOI) Report

Seretide Mite
 (Fluticasone
 Propionate,
 Salmeterol
 Xinafoate) C
 Ipratropium Bromide
 (Ipratropium
 Bromide) C
 Macrogol (Macrogol) C
 Docusate Sodium
 (Docusate Sodium) C
 Fludrocortisone
 Acetate
 (Fludrocortisone
 Acetate) C
 Esomeprazole
 (Esomeprazole) C
 Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C

Date:01/22/04ISR Number: 4279279-5Report Type:Expedited (15-DaCompany Report #2002069039
 Age:42 YR Gender:Male I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG		Anxiety Grand Mal Convulsion	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
DAILY, ORAL		Loss Of Consciousness	Professional				
800 MG,		Myoclonic Epilepsy Petit Mal Epilepsy		Clozapine (Clozapine)	SS		ORAL
DAILY, ORAL		Urinary Incontinence		Venlafaxine (Venlafaxine)	C		
				Panadine Co (Codeine Phosphate, Paracetamol)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (TID), ORAL	Stevens-Johnson Syndrome Toxic Skin Eruption	Foreign Health Professional	Neurontin (Gabapentin) Omeprazole (Omeprazole) Metoprolol Tartrate (Metoprolol Tartrate) Alprazolam (Alprazolam) Atenolol (Atenolol) Fluvastatin (Fluvastatin) Endotelon (Vitis Vinifera, Herbal Extracts Nos) Methylprednisolone Sodium Succinate	PS C C C C C C		ORAL

Freedom Of Information (FOI) Report

(Methylprednisolone
Sodium Succinate) C

Date:01/23/04ISR Number: 4278332-XReport Type:Expedited (15-DaCompany Report #PHRM2004FR00564
Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia Chlamydial Infection Diastolic Dysfunction		Hydergine	PS	Novartis Sector: Pharma	ORAL
400 mg, TID	87840MIN	Dyspnoea Dyspnoea At Rest Dyspnoea Exertional Headache Hypertrophic Cardiomyopathy Inflammation Interstitial Lung Disease Pulmonary Embolism Pulmonary Fibrosis		Neurontin	SS		ORAL
				Fosamax	SS		ORAL

Date:01/23/04ISR Number: 4279205-9Report Type:Expedited (15-DaCompany Report #2004001907
Age:86 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Burning Sensation Cough	Consumer	Neurontin (Gabapentin)	PS		
300 MG (100, TID)		Dysphagia					
		Feeling Abnormal Headache Medication Error Pharyngolaryngeal Pain Sensation Of Pressure Throat Tightness Tinnitus		Losartan Potassium (Losartan Potassium)	C		
				Pantoprazole (Pantoprazole)	C		
				Hydrochlorothiazide (Hydrochlorothiazide)	C		
				Metoprolol Succinate (Metoprolol Syccinate)	C		
				Docusate Sodium			

Date:01/23/04ISR Number: 4279219-9Report Type:Expedited (15-DaCompany Report #2004002865
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatic Failure	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:01/23/04ISR Number: 4279655-0Report Type:Expedited (15-DaCompany Report #2004000436
Age:98 YR Gender:Female I/FU:I

Outcome	PT	Report Source
Death	Blood Glucose Increased	Foreign
Other	Glucose Urine Present Glycosylated Haemoglobin Increased	Health Professional Company

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
ORAL		Neurontin (Gabapentin)	PS		ORAL

Date:01/23/04ISR Number: 4279714-2Report Type:Expedited (15-DaCompany Report #2004002110
Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID),		Coordination Abnormal Face Injury	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Fall Haematoma Vertigo	Professional				

Date:01/26/04ISR Number: 4280042-XReport Type:Expedited (15-DaCompany Report #2004002729
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (TID), ORAL		Sensory Loss	Foreign Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Company Representative				

Date:01/26/04ISR Number: 4280107-2Report Type:Expedited (15-DaCompany Report #2003118184
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2.4 GRAM		Blood Glucose Fluctuation Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		ORAL

	Diabetes Mellitus	Professional			
(DAILY); ORAL		Company	Celebrex (Celecoxib)	SS	ORAL
400 MG (BID);		Representative			
ORAL			Sinequan (Doxepin)	SS	ORAL
30 MG (AT					
NIGHT); ORAL					
			Tramadol Hydrochloride (Tramadol Hydrochloride)	SS	ORAL
200 MG (BID);					
ORAL					
			All Other Therapeutic Products	C	
			All Other Therapeutic Products	C	
			All Other Therapeutic Products	C	
			All Other Therapeutic Products	C	
			All Other Therapeutic Products	C	
			All Other Therapeutic Products	C	
			All Other Therapeutic Products	C	
			All Other Therapeutic Products	C	
			All Other Therapeutic Products	C	
			All Other Therapeutic Products	C	
			All Other		

Freedom Of Information (FOI) Report

Therapeutic Products C
 All Other
 Therapeutic Products C
 All Other
 Therapeutic Products C
 All Other
 Therapeutic Products C
 Drug Used In
 Diabetes C

Date:01/26/04ISR Number: 4280147-3Report Type:Expedited (15-DaCompany Report #2003038770
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Depression	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Hospitalization - 1200 MG		Mania					
Initial or Prolonged (TID), ORAL		Psychotic Disorder	Professional				
Other			Company Representative	Ethanol (Ehtanol)	SS		
				Opioids	C		
				Fentanyl (Fentanyl)	C		
				Paracetamol (Paracetamol)	C		
				Morphine (Morphine)	C		
				Diclofenac Sodium (Diclofenac Sodium)	C		

Date:01/26/04ISR Number: 4280175-8Report Type:Expedited (15-DaCompany Report #2004002253
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional	Neurontin (Gabapentin)	PS		
			Company Representative				

Date:01/26/04ISR Number: 4280177-1Report Type:Expedited (15-DaCompany Report #2003127209
 Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Body Height Decreased Loss Of Consciousness	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
3200 MG		Pain	Professional				
(DAILY), ORAL				Carisoprodol (Carisoprodol) Diazepam (Diazepam)	C C		

Date:01/26/04ISR Number: 4280219-3Report Type:Expedited (15-DaCompany Report #2004002731
Age: Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Chest Pain
Initial or Prolonged	Complex Partial Seizures
Other	Fall Fatigue Loss Of Consciousness Overdose Pain Paraesthesia

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Freedom Of Information (FOI) Report

Dose	Duration	Thrombosis Tremor Weight Decreased	Report Source	Product	Role	Manufacturer	Route
2800 MG (7 TIMES A DAY), ORAL			Consumer	Dilantin Suspension (Phenytoin Sodium) Neurontin (Gabapentin)	PS SS		ORAL
				Lamotrigine (Lamotrigine)	C		
				Atorvastatin (Atorvastatin)	C		
				Benazepril Hydrochloride (Benazepril Hydrochloride)	C		
				Escitalopram (Escitalopram)	C		
				Carbamazepine (Carbamazepine)	C		
				Atenolol (Atenolol)	C		
				Warfarin (Warfarin)	C		

Date:01/26/04ISR Number: 4280220-XReport Type:Expedited (15-DaCompany Report #2004002262
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (ONCE), ORAL		Amnesia Anxiety Back Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Bradyphrenia Cerebrovascular Accident Cognitive Deterioration Colitis Ischaemic Depression Dysarthria Eyelid Ptosis		Valproate Semisodium (Valproate Semisodium)	SS		

Facial Palsy
 Headache
 Hypoaesthesia
 Impaired Driving Ability
 Intervertebral Disc
 Degeneration
 Paraesthesia

Date:01/26/04ISR Number: 4280226-0Report Type:Expedited (15-DaCompany Report #2004002059

Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG		Accidental Overdose Coma	Consumer	Doxepin (Caps) (Doxepin)	PS		ORAL
Other (DAILY), ORAL		Dyskinesia					
900 MG		Electroencephalogram Abnormal		Neurontin (Gabapentin)	SS		ORAL
(DAILY), ORAL		Epilepsy					
		Facial Bones Fracture Fall Limb Operation		Levothyroxine Sodium (Levothyroxine Sodium)	C		

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Freedom Of Information (FOI) Report

Omeprazole
 (Omeprazole) C
 Verapamil
 Hydrochloride
 (Verapamil
 Hydrochloride) C
 Temazepam
 (Temazepam) C
 Baclofen (Baclofen) C
 Carisoprodol
 (Carisoprodol) C
 Parafon Forte
 (Chlorzoxazone,
 Paracetamol) C
 Hydroxyzine Embonate
 (Hydroxyzine
 Embonate) C

Date:01/26/04ISR Number: 4280245-4Report Type:Expedited (15-DaCompany Report #2003117742

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (TID) Other	Atrial Fibrillation Hypotension	Health Professional	Neurontin (Gabapentin) Verapamil (Verapamil)	PS C		

Date:01/26/04ISR Number: 4280281-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200657

Age:44 YR Gender:Female I/FU:F

Outcome	PT
Disability Other	Aphasia Aphonia Blood Thyroid Stimulating Hormone Decreased Brain Damage Carpal Tunnel Syndrome Depression Dissociative Disorder Disturbance In Attention Drug Hypersensitivity

Drug Toxicity
Dysarthria
Dyskinesia
Eustachian Tube
Dysfunction
Feeling Jittery
Gingival Recession
Hypersensitivity
Intentional Misuse
Intervertebral Disc
Disorder
Joint Injury
Limb Discomfort
Medication Error
Meniere'S Disease
Mental Impairment
Muscle Spasms
Polyneuropathy

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Adverse Event	Report Source	Product	Role	Manufacturer	Route
1800 MG (DAILY), ORAL		Radiculopathy					
		Road Traffic Accident					
		Sedation	Report Source	Product	Role	Manufacturer	Route
		Speech Disorder	Consumer	Neurontin			
		Swollen Tongue	Health	(Gabapentin)	PS		ORAL
		Syncope	Professional				
		Temperature Intolerance		Lamotrigine			
		Tendonitis		(Lamotrigine)	C		
		Thyroid Disorder		Lithium Carbonate			
		Tremor		(Lithium Carbonate)	C		
		Viral Labyrinthitis		Clonazepam			
				(Clonazepam)	C		
				Methylphenidate			
				Hydrochloride			
			(Methylphenidate				
			Hydrochloride)	C			
			Levothyroxine Sodium				
			(Levothyroxine				
			Sodium)	C			
			Liothyronine Sodium				
			(Liothyronine				
			Sodium)	C			
			Sertraline				
			Hydrochloride				
			(Sertraline				
			Hydrochloride)	C			
			Pilocarpine				
			Hydrochloride				
			(Pilocarpine				
			Hydrochloride)	C			
			Metoprolol Succinate				
			(Metoprolol				
			Succinate)	C			
			Lansoprazole				
			(Lansoprazole)	C			
			Hyoscyamine Sulfate				
			(Hyoscyamine				
			Sulfate)	C			
			Diltiazem				
			Hydrochloride				
			(Diltiazem				
			Hydrochloride)	C			
			Bupropion				
			Hydrochloride				

(Bupropion Hydrochloride)	C
Celebrex (Celecoxib)	C
Quetiapine Fumarate (Quetiapine Fumarate)	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C
Olanzapine	

Freedom Of Information (FOI) Report

(Olanzapine) C
 Omeprazole
 (Omeprazole) C
 Nasal Preparations C
 Nefazodone
 Hydrochloride
 (Nefazodone
 Hydrochloride) C
 Totolin
 (Guaifenesin,
 Phenylpropanolamine
 Hydrochloride) C
 Meclozine
 (Meclozine) C
 Fluticasone
 Propionate
 (Fluticasone
 Propionate) C
 Trimethobenzamide
 Hydrochloride
 (Trimethobenzamide
 Hydrochloride) C
 Respaire-Sr-120
 (Guaifenesin,
 Pseudoephedrine
 Hydrochloride) C

Date:01/26/04ISR Number: 4280294-6Report Type:Expedited (15-DaCompany Report #2003126804

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Contusion	Consumer	Neurontin			
Other		Coordination Abnormal	Health	(Gabapentin)	PS		ORAL
600 MG		Difficulty In Walking	Professional				
(DAILY), ORAL		Dizziness		Fentanyl (Fentanyl)	C		
		Drug Toxicity		Fluoxetine			
		Fall		Hydrochloride			
		Feeling Abnormal		(Fluoxetine			
		Head Injury		Hydrochloride)	C		
		Myoclonus		Bupropion			
		Pain In Extremity		Hydrochloride			
				(Bupropion			
				Hydrochloride)	C		

Fexofenadine Hydrochloride (Fexofenadine Hydrochloride)	C
Pantoprazole (Pantoprazole)	C
Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C
Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride)	C
Rofecoxib (Rofecoxib)	C
Trazodone (Trazodone)	C

Freedom Of Information (FOI) Report

Modafinil
 (Modafinil) C
 Dyazide
 (Hydrochlorothiazide
 , Triamterene) C
 All Other
 Therapeutic Product C

Date:01/26/04ISR Number: 4280322-8Report Type:Expedited (15-DaCompany Report #2002050043
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abdominal Distension	Consumer	Neurontin			
Other		Adrenal Adenoma	Health	(Gabapentin)	PS		ORAL
(TWICE		Blood Arsenic Increased	Professional				
DAILY), ORAL		Blood Mercury		Lamotrigine			
ORAL		Cataract		(Lamotrigine)	SS		ORAL
		Creatinine Urine		Clonazepam			
		Decreased		(Clonazepam)	SS		
		Drug Ineffective		Nifedipine			
		Dry Mouth		(Nifedipine)	C		
		Dyskinesia					
		Dyspnoea					
		Eating Disorder					
		Glaucoma					
		Helicobacter Infection					
		Hunger					
		Micturition Disorder					
		Nausea					
		Pharmaceutical Product					
		Complaint					
		Pollakiuria					
		Sensation Of Pressure					
		Treatment Noncompliance					
		Weight Decreased					
		Weight Increased					

Date:01/27/04ISR Number: 4280560-4Report Type:Expedited (15-DaCompany Report #2004001236
 Age:80 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG		Asthenia Cardiomegaly	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other (TID), ORAL		Chlamydial Infection	Professional				
ORAL		Computerised Tomogram Abnormal		Dihydroergotoxine (Dihydroergotoxine)	SS		ORAL
ORAL		Dyspnoea Exertional Headache		Alendronate Sodium (Alendronate Sodium)	SS		ORAL
		Interstitial Lung Disease Pulmonary Embolism Pulmonary Fibrosis					

Date:01/27/04ISR Number: 4280603-8Report Type:Expedited (15-DaCompany Report #2003116568
Age: Gender:Female I/FU:I

Outcome
Hospitalization -
Initial or Prolonged

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Other

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain	Consumer	Neurontin			
900 MG TID		Balance Disorder	Health	(Gabapentin)	PS		ORAL
ORAL		Convulsion	Professional				
		Dizziness		Citalopram			
		Fall		Hydrobromide	C		
		Feeling Abnormal		Atenolol	C		
		Headache					
		Loss Of Consciousness					

Date:01/28/04ISR Number: 4280477-5Report Type:Direct Company Report #CTU 210953
 Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation		Ms Contin	PS		ORAL
Hospitalization -							
90 MG TID PO		Confusional State		Neurotin	SS		ORAL
Initial or Prolonged		Hostility					
300 MG Q HS							
PO							

Date:01/28/04ISR Number: 4280513-6Report Type:Direct Company Report #CTU 210992
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression		Neurontin 800 Mg	PS		
Hospitalization -							
4 TIMES		Suicide Attempt					
Initial or Prolonged							

Date:01/29/04ISR Number: 4282471-7Report Type:Expedited (15-DaCompany Report #2004004401
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Dialysis	Health	Neurontin	
	Hypoglycaemia	Professional	(Gabapentin)	PS
	Therapy Non-Responder			

Date:01/29/04ISR Number: 4282495-XReport Type:Expedited (15-DaCompany Report #2003120051
Age:74 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG, TID, Other ORAL	Diarrhoea Fatigue Feeling Abnormal Influenza Like Illness Nausea	Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Rosiglitazone Maleate	SS		
			Albyl Med Kodein (Acetylsalicylic Acid, Codeine Phosphate)	SS		
			Oxycocet (Paracetamol, Oxycodone Hydrochloride)	C		
			Desloratadine	C		
			Prochlorperazine	C		
			Cetirizine			
			Hydrochloride	C		
			Drug, Unspecified	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/29/04ISR Number: 4282510-3Report Type:Expedited (15-DaCompany Report #2003119698

Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Body Height Decreased	Consumer	Neurontin			
Other		Cataract	Health	(Gabapentin)	PS		ORAL
600 MG, ORAL		Choroidal Neovascularisation	Professional	Feldene (Piroxicam)	C		
		Somnolence		Lipitor (Atorvastatin)	C		
				Lisinopril (Lisinopril)	C		
				Verapamil (Verapamil)	C		
				Furosemide (Furosemide)	C		
				Potassium (Potassium)	C		

Date:01/29/04ISR Number: 4282512-7Report Type:Expedited (15-DaCompany Report #2004003767

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Respiratory Arrest	Health	Neurontin			
Other		Respiratory Depression	Professional	(Gabapentin)	PS		

Date:01/29/04ISR Number: 4282515-2Report Type:Expedited (15-DaCompany Report #2004003765

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Respiratory Arrest	Health	Neurontin			
Other		Respiratory Depression	Professional	(Gabapentin)	PS		

Date:01/29/04ISR Number: 4282517-6Report Type:Expedited (15-DaCompany Report #2004003763

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Respiratory Arrest	Health	Neurontin	
Other	Respiratory Depression	Professional	(Gabapentin)	PS

Date:01/29/04ISR Number: 4282540-1Report Type:Expedited (15-DaCompany Report #2003008068
Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Aplastic Anaemia
Initial or Prolonged	Arthralgia
Disability	Asthenia
Other	Balance Disorder
	Burns Second Degree
	Coma
	Coordination Abnormal
	Crying
	Disturbance In Attention
	Dizziness
	Dry Mouth
	Epilepsy
	Fall
	Grand Mal Convulsion
	Headache

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 1200 MG (TID), ORAL		Heart Rate Increased	Consumer Health Professional	Neurontin (Gabapentin)	PS		
		Muscle Spasms					
		Nerve Injury					
		Nervousness		Dilantin (Phenytoin Sodium)	SS		ORAL
		Pain					
		Pain In Extremity					
		Panic Attack		Acetylsalicylic Acid (Acetylsalicylic Acid)	SS		
		Paralysis					
		Speech Disorder					
				Vitamins With Minerals	SS		
				Megestrol	C		
				(Megestrol)			

Date:01/29/04ISR Number: 4283190-3Report Type:Expedited (15-DaCompany Report #2004003787
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 1800 MG Initial or Prolonged (TID),		Pneumonia	Foreign Health Professional	Neurontin (Gabapentin)	PS		
		Septic Shock					
		Status Epilepticus					
			Company Representative	Bromazepam	C		
				Antihypertensives	C		
				Hypnotics And Sedatives	C		

Date:01/29/04ISR Number: 4283295-7Report Type:Expedited (15-DaCompany Report #2004003778
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2600 MG (BID)		Blood Pressure Decreased	Foreign Consumer	Neurontin (Gabapentin)	PS		
		Headache					

Somnolence
Weight Increased

Hydrocortisone
(Hydrocortisone) C

Date:01/29/04ISR Number: 4283296-9Report Type:Expedited (15-DaCompany Report #2004003783
Age:72 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Convulsion	Foreign	Neurontin			
Initial or Prolonged	Dystonia	Health	(Gabapentin)	PS		ORAL
900 MG						
Other	Epilepsy	Professional				
(DAILY), ORAL						
	Myoclonic Epilepsy		Teicoplanin			
			(Teicoplanin)	SS		
INTRAVENOUS	8 MG/KG					
(ONCE),						
INTRAVENOUS						

Date:01/29/04ISR Number: 4283627-XReport Type:Direct Company Report #CTU 211127
Age:37 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Azotaemia
Initial or Prolonged	Chromaturia
	Decreased Appetite

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dehydration					
		Mental Status Changes					
		Overdose					
500 BID, CHRONIC 80 QD 600 TID CHRONIC 100 QD		Somnolence		Naprosyn	PS		
		Urinary Tract Infection					
		Urine Output Decreased		Lasix	SS		
				Neurontin	SS		
				Spironolactone	SS		
				Folic Acid	C		
				Darvon	C		
				Darvocet	C		
				Zyrtec	C		
				Paxil	C		
				Trazodone	C		
				Elavil	C		
				Bupropion	C		
				Risperidal	C		
				Flucanionide	C		
				Cotrimazole	C		
				Soma	C		

Date:01/30/04 ISR Number: 4282121-X Report Type:Periodic Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12372553
 Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nervousness		Trazodone Hcl Tabs	PS	Apothecon	ORAL
				Effexor	SS		
				Xanax	SS		
				Lamictal	SS		
				Neurontin	SS		
				Synthroid	SS		

Date:01/30/04 ISR Number: 4282838-7 Report Type:Periodic Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12286415
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Serzone Tabs 100 Mg	PS	Bristol-Myers Squibb Company	ORAL
100 mg daily							
in the							
evening, meds							
not taken for							
1 wk as of	1	YR					
				Effexor	SS		
				Neurontin	SS		
				Requip	SS		
				Flexeril	SS		
"Adderall XR"				Adderall	SS		
				Vitamin B12	C		
PARENTERAL							

Date:01/30/04ISR Number: 4282902-2Report Type:Periodic Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12413118
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Serzone Tabs	PS	Bristol-Myers Squibb Company	ORAL
Other		Hepatitis					
100mg							
tablets; 1 to							
4 tabs at hs.							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Stopped

Dec-2000,

Lipitor

SS

On 10mg/day

as of

06-Feb-2000.

Stopped

Dec-2000,

Gabapentin

SS

ORAL

100mg,300mg,8

00mg caps.1-3

hs-tid

prn.Stop

12/00,restart

Gabapentin

SS

ORAL

100mg,300mg,8

00mg caps.1-3

hs-tid

prn.Stop

12/00,restart

Gabapentin

SS

ORAL

100mg,300mg,8

00mg caps.1-3

hs-tid

prn.Stop

12/00,restart

Date:01/30/04ISR Number: 4284492-7Report Type:Expedited (15-DaCompany Report #2004004409

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysphagia	Health	Neurontin (Tablets)			
		Malignant Neoplasm	Professional	(Gabapentin)	PS		ORAL
ORAL		Progression Neoplasm Malignant					

Date:01/30/04ISR Number: 4284579-9Report Type:Expedited (15-DaCompany Report #2004003994

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Developmental Delay	Foreign	Neurontin			
		Maternal Drugs Affecting	Health	(Gabapentin)	PS		
PERIARTICULAR	2400 MG	Foetus	Professional				

(TID),

PLACENTAL

Date:01/30/04ISR Number: 4284585-4Report Type:Expedited (15-DaCompany Report #2004003772

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dizziness	Foreign	Neurontin			
Initial or Prolonged		Fall	Health	(Gabapentin)	PS		ORAL
300 MG							
Other		Paralysis	Professional				

(DAILY), ORAL

Company
Representative

Date:01/30/04ISR Number: 4284706-3Report Type:Expedited (15-DaCompany Report #2003121316

Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT
Hospitalization -		Biliary Dilatation
Initial or Prolonged		Cytolytic Hepatitis
		Drug Interaction

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hepatitis Cholestatic Psychotic Disorder Treatment Noncompliance	Report Source	Product	Role	Manufacturer	Route
300 MG (TID), ORAL			Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Trimetazidine Hydrochloride (Trimetazidine Hydrochloride)	C		
				Trimebutine (Trimebutine)	C		
				Domperidone (Domperidone)	C		
				Lisinopril (Lisinopril)	C		
				Milnacipran (Milnacipran)	C		
				Lormetazepam (Lormetazepam)	C		
				Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
				Clonazepam (Clonazepam)	C		

Date:02/02/04ISR Number: 4284252-7Report Type:Direct
Age:81 YR Gender:Female I/FU:I

Company Report #CTU 211285

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG BID ORAL			Dyskinesia Tremor		Neurontin 300 Mg Parke-Davis	PS	Parke-Davis	ORAL

Date:02/02/04ISR Number: 4285018-4Report Type:Expedited (15-DaCompany Report #2004003799
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3600 MG ORAL Other		Alanine Aminotransferase Increased	Foreign Literature	Gabapentin (Gabapentin)	PS		ORAL
		Anaesthesia Aspartate Aminotransferase Abnormal Blood Alkaline Phosphatase Increased Blood Lactate Dehydrogenase Abnormal Gamma-Glutamyltransferase Increased Hepatotoxicity Hypoaesthesia Neuropathic Pain	Health Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/04ISR Number: 4285359-0Report Type:Expedited (15-DaCompany Report #2003119508

Age:31 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID), Other ORAL	Abnormal Behaviour Back Injury Contusion	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL	Depressed Mood Diarrhoea	Company Representative	Diazepam (Diazepam) Tramadol (Tramadol)	SS SS		ORAL
ORAL	Euphoric Mood Fall		Temazepam (Temazepam)	SS		ORAL
	Feeling Abnormal Judgement Impaired Mood Altered Muscle Spasms Nausea Overdose Pain Self-Medication		Diamorphine (Diamorphine) Alimemazine Tartrate (Alimemazine Tartrate) Indometacin (Indometacin) Panadeine Co (Codeine Phosphate, Paracetamol) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C C C C C		

Date:02/02/04ISR Number: 4285407-8Report Type:Expedited (15-DaCompany Report #PHRM2004FR00564

Age:80 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Asthenia Chlamydial Infection Diastolic Dysfunction	Foreign Health Professional	Hydergine (Co-Dergocrine Mesylate) Unknown	PS		ORAL
400 MG TID ORAL	Dyspnoea Dyspnoea Exertional Headache	Other	Neurontin (Gabapentin) Table	SS		ORAL

Hypertrophic
Cardiomyopathy

Fosamax (Alendronate
Sodium) Tablet SS

ORAL

ORAL

Iatrogenic Injury
Inflammation
Interstitial Lung Disease
Pulmonary Embolism
Pulmonary Fibrosis

Date:02/02/04ISR Number: 4285621-1Report Type:Expedited (15-DaCompany Report #2004002698

Age: Gender:Female I/FU:I

Outcome

PT

Other

Asthenia
Blood Glucose Fluctuation
Convulsion
Dizziness
Drug Interaction
Dysgeusia
Ear Pain
Fatigue
Fluid Retention
Glucose Tolerance
Decreased
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
800 MG	203 MON	Hypoglycaemia Impaired Work Ability Lethargy	Health Professional	Neurontin (Gabapentin)	PS		
		Oedema Peripheral Rhinitis		Zithromax (Azithromycin)	SS		
		Somnolence Thinking Abnormal Treatment Noncompliance		Elavil (Amitriptyline Hydrochloride)	SS		ORAL
				Ibuprofen (Ibuprofen)	SS		
				Mometasone Furoate (Mometasone Furoate)	SS		
				Caffeine (Caffeine)	SS		ORAL

(DAILY), ORAL

Date:02/03/04ISR Number: 4285956-2Report Type:Expedited (15-DaCompany Report #2004005100
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN		Diplopia Paralysis	Health Professional	Neurontin (Gabapentin)	PS		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C		
				Folic Acid (Folic Acid)	C		
				Atorvastatin (Atorvastatin)	C		
				Budesonide (Budesonide)	C		
				Salbutamol (Salbutamol)	C		
				Cetirizine Hydrochloride			

(Cetirizine Hydrochloride)	C
Mometasone Furoate (Mometasone Furoate)	C
Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride)	C
Vicodin (Paracetamol, Hydrochloride Bitartrate)	C
Nortriptyline Hydrochloride (Nortriptyline Hydrochloride)	C
Metaxalone (Metaxalone)	C
Ranitidine Hydrochloride (Ranitidine	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C

Date:02/03/04ISR Number: 4285994-XReport Type:Expedited (15-DaCompany Report #2003041722

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Coordination Abnormal	Consumer	Neurontin			
Other		Dystonia		(Gabapentin)	PS		ORAL
600 MG (300,		Memory Impairment					
BID), ORAL		Nervous System Disorder		Lipitor			
		Tremor		(Atorvastatin)	C		
				Clopidogrel Sulfate			
				(Clopidogrel			
				Sulfate)	C		
				Toprol (Metoprolol)	C		
				Ace Inhibitor Nos	C		

Date:02/03/04ISR Number: 4286040-4Report Type:Expedited (15-DaCompany Report #2004004406

Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Arteriospasm Coronary	Consumer	Neurontin			
Initial or Prolonged		Chest Discomfort		(Gabapentin)	PS		ORAL
ORAL		Laboratory Test Abnormal		Lipitor			
Other		Pallor		(Atrovastatin)	C		
				Multivitamins			
				(Ergocalciferol,			
				Ascorbic Acide,			
				Folic Acid, Thiamine			
				Hydrochloride,	C		

Date:02/03/04ISR Number: 4286064-7Report Type:Expedited (15-DaCompany Report #2004004686

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder	Consumer	Neurontin			

DAILY , ORAL	Cognitive Disorder	(Gabapentin)	PS	ORAL
	Coordination Abnormal	Prednisone		
	Dizziness	(Prednisone)	SS	
	Neuropathy Peripheral	Levothyroxine Sodium	C	
	Somnolence	Estrogens Conjugated	C	
	Vasculitis	Celecoxib	C	
	Weight Increased	Acetylsalicylic Acid	C	

Date:02/03/04ISR Number: 4286180-XReport Type:Expedited (15-DaCompany Report #2004004679
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dependence	Foreign	Neurontin			
Other		Drug Withdrawal Syndrome	Health	(Gabapentin)	PS		ORAL
1200 MG		Feeling Cold	Professional				
(TID), ORAL		Pain		Valoron N (Naloxone			
		Tremor		Hydrochloride,			
				Tilidine			
				Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/04/04ISR Number: 4286769-8Report Type:Expedited (15-DaCompany Report #2004005558

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Dilantin Kapseals			
		Convulsion		(Phenytoin Sodium)	PS		ORAL
400 MG (QID),							
ORAL		Pharmaceutical Product					
		Complaint		Neurontin			
900 MG (TID),				(Gabapentin)	SS		ORAL
ORAL							
				Primidone			
				(Primidone)	C		
				Carbamazepine			
				(Carbamazepine)	C		
				Ranitidine			
				(Ranitidine)	C		
				Alprazolam			
				(Alprazolam)	C		
				Levothyroxine Sodium			
				(Levothyroxine			
				Sodium)	C		
				Fluoxetine			
				Hydrochloride			
				(Fluoxetine			
				Hydrochloride)	C		

Date:02/04/04ISR Number: 4286786-8Report Type:Expedited (15-DaCompany Report #2004005466

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Consumer	Dilantin Kapseals			
		Asthenia		(Phenytoin Sodium)	PS		ORAL
ORAL							
		Chest Pain		Neurontin			
600 MG (BID),		Fatigue		(Gabapentin)	SS		ORAL
ORAL		Insomnia					
		Vitamin B Complex		Phenytoin			

Deficiency

(Phenytoin) SS
Phenobarbital
(Phenobarbital) C

Date:02/04/04ISR Number: 4286845-XReport Type:Expedited (15-DaCompany Report #2004005992
Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Hepatitis Fulminant	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Professional	Paracetamol (Paracetamol)	SS		ORAL
4 GRAM, ORAL			Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		ORAL
ORAL			Morphine Sulfate (Morphine Sulfate)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/04/04ISR Number: 4287044-8Report Type:Expedited (15-DaCompany Report #2003124917

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ageusia	Foreign	Neurontin			
ORAL		General Physical Health	Health	(Gabapentin)	PS		ORAL
		Deterioration	Professional	Tamoxifen			
		Neoplasm Progression		(Tamoxifen)	C		
		Tinnitus		Morphine (Morphine)	C		

Date:02/04/04ISR Number: 4287045-XReport Type:Expedited (15-DaCompany Report #2003124090

Age:81 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Aphasia	Foreign	Neurontin			
Initial or Prolonged		Transient Ischaemic	Health	(Gabapentin)	PS		ORAL
400 MG (BID),		Attack	Professional				
ORAL			Company	Aporex (Paracetamol			
			Representative	Dextropropoxyphene			
				Hydrochloride)	C		
				Diclofenac Sodium			
				(Diclofenac Sodium)	C		
				Calcifediol			
				(Calcifediol)	C		
				Pantoprzole			
				(Pantoprazole)	C		
				Dimeticone			
				(Dimeticone)	C		
				Mebeverine			
				(Mebeverine)	C		

Date:02/04/04ISR Number: 4287046-1Report Type:Expedited (15-DaCompany Report #2004003783

Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion	Foreign	Neurontin			

Initial or Prolonged Epilepsy Health (Gabapentin) PS ORAL
900 MG
Other Professional
(DAILY), ORAL
INTRAVENTOUS 8 MG/KG Teicoplanin (Teicoplanin) SS
(ONCE),
INTRAVENTOUS

Date:02/04/04ISR Number: 4287180-6Report Type:Expedited (15-DaCompany Report #2004004691
Age:49 YR Gender:Male I/FU:I

Outcome PT
Life-Threatening Blood Calcium Increased
Hospitalization - Blood Creatinine
Initial or Prolonged Increased
Other Blood Potassium Increased
Blood Sodium Increased
Cardiac Arrest
Life Support
Neurological Examination
Abnormal
Pupil Fixed
Respiratory Failure
Sedation

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Unresponsive To Pain Stimuli	Report Source	Product	Role	Manufacturer	Route
1800 (TID),			Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
ORAL				Lorazepam (Lorazepam)	SS		
				Haloperidol (Haloperidol)	SS		
				Valproic Acid (Valproic Acid)	SS		
				Salbutamol (Salbutamol)	C		
				Amlodipine (Amlodipine)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
				Enalapril (Enalapril)	C		
				Ezetimibe (Ezetimibe)	C		
				Glibenclamide (Glibenclamide)	C		
				Isosorbide Mononitrate (Isosorbide Mononitrate)	C		
				Levofloxacin (Levofloxacin)	C		
				Olanzapine (Olanzapine)	C		
				Prenatal Vitamins (Ascorbic Acid, Biotin , Tocopherol, Nicotinic Acid, Retinol, Vitamin D	C		
				Rofecoxib (Rofecoxib)	C		
				Simvastatin			

(Simvastatin) C
Rosiglitazone
(Rosiglitazone) C

Date:02/04/04ISR Number: 4287184-3Report Type:Expedited (15-DaCompany Report #2004001906
Age:44 YR Gender:Female I/FU:F

Outcome PT
Other Amnesia
Anorgasmia
Caesarean Section
Dizziness
Feeling Abnormal
Neuropathic Pain
Panic Reaction
Paraesthesia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pregnancy Self-Medication Sexual Abuse	Report Source	Product	Role	Manufacturer	Route
900 MG (TID), ORAL		Somnolence	Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:02/04/04ISR Number: 4287187-9Report Type:Expedited (15-DaCompany Report #2004005958
Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2400 MG (QID), ORAL		Abnormal Behaviour Bronchitis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1000 MG (BID), ORAL		Drug Effect Decreased Drug Interaction		Naproxen (Naproxen)	SS		ORAL
		Pharmaceutical Product		Clonazepam (Clonazepam)	C		
		Complaint Sinusitis Tonic Clonic Movements		Lamotrigine (Lamotrigine)	C		
				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Calcium (Calcium) Fish Oil (Fish Oil)	C C C		

Date:02/04/04ISR Number: 4287189-2Report Type:Expedited (15-DaCompany Report #2003114753
Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization -		Bone Neoplasm Malignant Lung Cancer Metastatic	Consumer Health	Neurontin (Gabapentin)	PS		

Initial or Prolonged	Lung Carcinoma Cell Type	Professional	Omeprazole	
Other	Unspecified Stage Iv		(Omeprazole)	C
	Pulmonary Oedema		Clonazepam	
			(Clonazepam)	C

Date:02/05/04ISR Number: 4286257-9Report Type:Direct Company Report #CTU 211566
 Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Pruritus		Neurontin 300 Mg	PS		ORAL
300 MG PO TID		Rash		Lisinopril	C		

Date:02/05/04ISR Number: 4286939-9Report Type:Expedited (15-DaCompany Report #2012272
 Age:42 YR Gender:Male I/FU:F

Outcome	PT
Death	Accident
Hospitalization -	Accidental Overdose
Initial or Prolonged	Back Pain
Other	Drug Abuser
	Drug Toxicity
	Haematemesis
	Headache

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Neck Pain

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
SEE IMAGE		Consumer Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		
			Diazepam (Diazepam)	SS		
			Oxazepam (Oxazepam)	SS		
			Temazepam (Temazepam)	SS		
			Lorazepam (Lorazepam)	SS		
			Cannabnoids (Cannabis)	SS		
			Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride)	SS		
			Gabapentin (Gabapentin)	SS		
			Claritin (Loratadine)	C		
			Zoloft (Sertraline Hydrochloride)	C		
			Altace (Ramipril)	C		
			Allopurinol (Allopurinol)	C		
			Depakote (Valproate Semisodium)	C		
			Soma (Carisoprodol)	C		
			Relafen (Nabumetone)	C		

Date:02/05/04ISR Number: 4287009-6Report Type:Expedited (15-DaCompany Report #USA-2004-0012636
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Bicarbonate Increased Blood Chloride Decreased Blood Creatinine	Study Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		

60 MG, BID

62.5 MG, BID,	Increased	Tracleer (Bosentan)	SS	ORAL
ORAL	Blood Sodium Decreased			
	Blood Urea Increased	Percocet		
	Encephalopathy	(Paracetamol,		
	Mental Status Changes	Oxycodone		
2 TABLET, Q4H	Myoclonus	Hydrochloride)	SS	
	Oedema Peripheral	Neurontin		
200 MG, TID,	Pain	(Gabapentin)	SS	ORAL
ORAL	Somnolence			
		Coumadin "Endo"		
		(Warfarin Sodium)	C	
		Lasix (Furosemide)	C	
		Zaroxolyn		
		(Metolazone)	C	
		Prednisone		
		(Prednisone)	C	
		Epoprostenol		
		(Epoprostenol)	C	
		Protonix		

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Freedom Of Information (FOI) Report

(Pantoprazole) C
 Glucophage
 (Metformin
 Hydrochloride) C
 Bactrim
 (Sulfamethoxazole,
 Trimethoprim) C
 Oxygen (Oxygen) C
 Lisinopril
 (Lisinopril) C

Date:02/05/04ISR Number: 4288447-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Biopsy Tongue Abnormal	Consumer	Neurontin			
Other		Blood Thyroid Stimulating	Health	(Gabapentin)	PS		ORAL
1800 MG		Hormone Decreased	Professional				
(DAILY), ORAL		Carpal Tunnel Syndrome		Hyoscyamine Sulfate			
		Depression		(Hyoscyamine			
		Dissociative Disorder		Sulfate)	SS		
		Dyskinesia		Nasal Preparations	SS		
		Feeling Jittery		Lamotrigine			
		Hypersensitivity		(Lamotrigine)	C		
		Injury		Lithium Carbonate			
		Intervertebral Disc		(Lithium Carbonate)	C		
		Disorder		Clonazepam			
		Meniere'S Disease		(Clonazepam)	C		
		Nervous System Disorder		Methylphenidate			
		Neuropathy		Hydrochloride (
		Overdose		Methylphenidate			
		Radiculopathy		Hydrochloride)	C		
		Road Traffic Accident		Levothyroxine Sodium			
		Sedation		(Levothyroxine			
		Speech Disorder		Sodium)	C		
		Swollen Tongue		Sertraline			
		Syncope		Hydrochloride			
		Tendonitis		(Sertraline			
		Viral Labyrinthitis		Hydrochloride)	C		
				Pilocarpine			
				Hydrochloride			
				(Pilocarpine			
				Hydrochloride)	C		

Metoprolol Succinate	
(Metoprolol	
Succinate)	C
Lansoprazole	
(Lansoprazole)	C
Hyoscyamine Sulfate	
(Hyoscyamine	
Sulfate)	C
Diltiazem	
Hydrochloride	
(Diltiazem	
Hdrochloride)	C
Bupropion	
Hydrochloride	
(Bupropion	
Hydrochloride)	C
Celebrex (Celecoxib)	C
Quetiapine Fumarate	

Freedom Of Information (FOI) Report

(Quetiapine Fumarate)	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
Risperidone (Risperidone)	C
Respaire-Sr-120 (Guaifenesin, Pseudoepedrine Hydrochloride)	C
Trimethobenazamide Hydrochloride (Trimethobenzamide Hydrochloride)	C
Fluticasone Propionate (Fluticasone Propionate)	C
Meclozine (Meclozine)	C
Totolin (Guaifenesin, Phenylpropanolamine Hydrochloride)	C
Nefazodone Hydrochloride (Nefazodone Hydrochloride)	C
Olanzapine (Olanzapine)	C
Omeprazole (Omeprazole)	C

Date:02/05/04ISR Number: 4288452-1Report Type:Expedited (15-DaCompany Report #2004005040
Age:84 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Condition Aggravated Fatigue	Consumer	Neurontin (Gabapentin)	PS		ORAL

Neuropathy
Rash

Metformin (Metformin)	C
Tamsulosin Hydrochloride (Tamsulosin Hydrochloride)	C
Lansoprazole (Lansoprazole)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C
Atorvastatin (Atorvastatin)	C
Glimepiride (Glimepiride)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/05/04ISR Number: 4288747-1Report Type:Expedited (15-DaCompany Report #2004003351
Age:73 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 600 MG ORAL Initial or Prolonged 600 MG (BID), ORAL	Cross Sensitivity Reaction Drug Eruption Eyelid Oedema	Foreign Health Professional	Neurontin (Gabapentin) Carbamazepine (Carbamazepine)	PS SS		ORAL ORAL
			Phenprocoumon (Phenprocoumon) Salutec (Hydrochlorothiazide , Ramipril) Torasemide (Torasemide)	C C C		

Date:02/06/04ISR Number: 4289167-6Report Type:Expedited (15-DaCompany Report #2004006906
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Renal Impairment	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/06/04ISR Number: 4289194-9Report Type:Expedited (15-DaCompany Report #2003037220
Age:74 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 900 MG TID ORAL	Acute Sinusitis C-Reactive Protein Increased Diplopia Red Blood Cell	Consumer Health Professional	Neurontin (Gabapentin) Levothyroxine Ezetimibe	PS C C		ORAL

Sedimentation Rate
Increased
Strabismus

Cefalexin C
Etodolac C
Pethidine
Hydrochloride C

Date:02/06/04ISR Number: 4289217-7Report Type:Expedited (15-DaCompany Report #2004001909
Age:48 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (TID), ORAL	Anorexia Intentional Misuse Marital Problem Nausea Suicidal Ideation Suicide Attempt Weight Decreased	Health Professional	Neurontin (Gabapentin) Trazodone (Trazodone) Citalopram Hydrobromide (Citalopram Hydrobromide) Methylphenidate Hydrochloride (Methylphenidate Hydrochloride)	PS SS C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/06/04ISR Number: 4289218-9Report Type:Expedited (15-DaCompany Report #2004005466

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Anxiety Asthenia	Consumer	Dilantin Kapseals (Phenytoin Sodium)	PS		ORAL
Other 1200 MG (BID), ORAL		Chest Discomfort Chest Pain		Neurontin (Gabapentin)	SS		ORAL
		Dementia					
300 MG (DAILY)		Drug Interaction Fatigue		Phenytoin (Phenytoin)	SS		
		Insomnia					
30 MG (DAILY)		Paraesthesia Vitamin B Complex Deficiency		Phenobarbital (Phenobarbital)	SS		

Date:02/06/04ISR Number: 4289244-XReport Type:Expedited (15-DaCompany Report #2004005672

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dyspnoea	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/06/04ISR Number: 4289246-3Report Type:Expedited (15-DaCompany Report #2004005992

Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Hepatitis Fulminant	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
			Professional	Paracetamol (Paracetamol)	SS		ORAL
4 GRAM , ORAL				Amitriptyline			

ORAL

Hydrochloride (Amitriptyline Hydrochloride)	SS	ORAL
Morphine Sulfate (Morphine Sulfate)	SS	

Date:02/06/04ISR Number: 4289407-3Report Type:Expedited (15-DaCompany Report #2004005673
 Age:59 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Condition Aggravated Dyspnoea	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/06/04ISR Number: 4289417-6Report Type:Expedited (15-DaCompany Report #2004006265
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Renal Disorder Renal Pain	Consumer	Neurontin (Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:02/06/04ISR Number: 4289910-6Report Type:Expedited (15-DaCompany Report #2004006431
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG (BID)		Cardiomegaly Myalgia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/06/04ISR Number: 4291338-XReport Type:Direct Company Report #CTU 211706
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 100 MG BID PO		Oedema Peripheral Pruritus Swelling Face		Neurontin 100 Mg-Parkdavis Lactulose Docusate Sodium Senna Ranitidine Levothyroid Celexa Kcl Lasix Zyrtec Multivit Norvasc Ferrous Sulfate Sulfatcim Ds Oxycontin	PS C C C C C C C C C C C C C	Parkdavis	ORAL

Date:02/09/04ISR Number: 4290234-1Report Type:Expedited (15-DaCompany Report #2004005157
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG (200,		Blood Creatinine Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL

Other TID), ORAL	Encephalopathy			
125 MG (62.5, BID), ORAL	Myoclonus Oedema Peripheral Somnolence		Bosentan (Bosentan) SS	ORAL
120 MG (BID), (Q4H, ORAL)			Oxycodone Hydrochloride (Oxycodone Hydrochloride) SS	
			Oxycocet (Paracetamol, Oxycodone Hydrochloride) SS	ORAL
			Warfarin Sodium (Warfarin Sodium) C	
			Furosemide (Furosemide) C	
			Metolazone (Metolazone) C	
			Prednisone (Prednisone) C	
			Epoprostenol Sodium (Epoprostenol Sodium) C	
			Pantoprazole	

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(Pantoprazole) C
 Metformin
 Hydrochloride
 (Metformin
 Hydrochloride) C
 Bactrim
 (Sulfamethoxazole,
 Trimethoprim) C
 Oxygen (Oxygen) C
 Lisinopril
 (Lisinopril) C

Date:02/09/04ISR Number: 4290245-6Report Type:Expedited (15-DaCompany Report #2004005888
 Age:69 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Abdominal Tenderness Abnormal Faeces	Health Professional	Neurontin (Gabapentin)	PS		ORAL
	Alopecia		Ketorolac			
	Anxiety		Tromethamine			
	Blood Pressure Systolic Increased		(Ketorolac Tromethamine)	C		
	Constipation		Esomeprazole			
	Depression		(Esomeprazole)	C		
	Diverticulitis		Atorvastatin			
	Drug Tolerance Decreased		(Atorvastatin)	C		
	Dysuria		Metronidazole			
	Ear Pain		(Metronidazole)	C		
	Haematuria		Levothyroxine Sodium			
	Insomnia		(Levothyroxine Sodium)	C		
	Middle Ear Effusion		Indapamide			
	Nausea		(Indapamide)	C		
	Oral Intake Reduced		Panadeine Co			
	Osteoarthritis		(Codeine Phosphate, Paracetamol)	C		
	Protein Urine		Zolpidem Tartrate			
	Rectal Haemorrhage		(Zolpidem Tartrate)	C		
	Sinusitis		Ultracet			
	Tenderness		(Paracetamol., Tramadol			
	Weight Decreased		Hydrochloride)	C		
	White Blood Cells Urine Positive		Alprazolam (Alprazolam)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Health	Neurontin			
		Depression	Professional	(Gabapentin)	PS		ORAL
2400 MG		Disturbance In Attention					
(QID), ORAL		Suicidal Ideation		Prednisone			
		Weight Increased		(Prednisone)	C		
				Clopidogrel Sulfate			
				(Clopidogrel			
				Sulfate)	C		
				Dipyridamole			
				(Dipyridamole)	C		
				Diltiazem			

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(Diltiazem) C
 Omeprazole
 (Omeprazole) C
 Hydrochlorothiazide
 (Hydrochlorothiazide
) C
 Hydroxyzine
 Hydrochloride
 (Hydroxyzine
 Hydrochloride) C
 Progesterone
 (Progesterone) C
 Metoprolol
 (Metoprolol) C
 Fludrocortisone
 Acetate
 (Fludrocortisone
 Acetate) C
 Zolpidem Tartrate
 (Zolpidem Tartrate) C
 Fentanyl (Fentanyl) C
 Alprazolam
 (Alprazolam) C

Date:02/09/04ISR Number: 4290258-4Report Type:Expedited (15-DaCompany Report #2003126804
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (DAILY), ORAL		Contusion Coordination Abnormal Difficulty In Walking	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Dizziness Drug Toxicity Dysstasia Fall Feeling Abnormal Head Injury Memory Impairment Myoclonus		Fentanyl (Fentanyl) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Bupropion Hydrochloride (Bupropion Hydrochloride) Fexofenadine Hydrochloride (Fexofenadine	C C C		

Hydrochloride)	C
Pantoprazole	
(Pantoprazole)	C
Oxycodone	
Hydrochloride	
(Oxycodone	
Hydrochloride)	C
Cyclobenzaprine	
Hydrochloride	
(Cyclobenzaprine	
Hydrochloride)	C
Rofecoxib	
(Rofecoxib)	C
Trazodone	
(Trazodone)	C
Modafinil	
(Modafinil)	C
Dyazide	

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(Hydrochlorothiazide
 , Triamterene) C
 All Other
 Therapeutic Products C
 Oxycodone
 Hydrochloride
 (Oxycodone
 Hydrochloride) C
 Morphine (Morphine) C

Date:02/09/04ISR Number: 4290261-4Report Type:Expedited (15-DaCompany Report #2003123581
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Cancer Stage I	Health	Neurontin			
Other		Cognitive Disorder	Professional	(Gabapentin)	PS		ORAL
100 MG (DAILY		Drug Dependence					
AT BEDTIME),		Drug Tolerance					
ORAL		Dysarthria		Propacet			
		Feeling Abnormal		(Paracetamol,			
		Food Craving		Dextropropoxyphene			
		Somnolence		Napsilate)	C		
		Weight Increased		Alendronate Sodium			
				(Alendronate Sodium)	C		
				Methocarbamol			
				(Methocarbamol)	C		
				Methylprednisolone			
				(Methylprednisolone)	C		
				Hydrochlorothiazide			
				(Hydrochlorothiazide			
)	C		
				Atenolol (Atenolol)	C		
				All Other			
				Therapeutic Products	C		
				Fentanyl. (Fentanyl)	C		
				Clonazepam			
				(Clonazepam)	C		
				Celecoxib			
				(Celecoxib)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hepatic Failure Hepatitis Fulminant	Foreign Health Professional Other	Morphine Sulfate(Morphine Sulfate) Unknown Efferalgen(Paracetam ol)	PS SS		ORAL
4 GRAM, DAILY, ORAL							
ORAL				Laroxyl (Amitriptyline Hydrochloride)	SS		ORAL
ORAL				Neurontin(Gabapentin)	SS		ORAL

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Date:02/09/04ISR Number: 4290734-4Report Type:Expedited (15-DaCompany Report #2003035798
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthropathy	Foreign	Neurontin			
1800 MG, ORAL		Condition Aggravated	Health	(Gabapentin)	PS		ORAL
		Psoriasis	Professional	Oxcarbazepine			
				(Oxcarbazepine)	C		

Date:02/10/04ISR Number: 4293215-7Report Type:Expedited (15-DaCompany Report #2004006267
Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Bone Neoplasm Malignant	Consumer	Neurontin			
ORAL		Hallucination, Visual		(Gabapentin)	PS		ORAL
		Somnolence		Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	SS		

Date:02/10/04ISR Number: 4293632-5Report Type:Expedited (15-DaCompany Report #04P-062-0249263-00
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Arthralgia	Foreign	Ergenyl (Sodium			
Intervention to		Bone Marrow Oedema	Health	Valproate) (Sodium			
Prevent Permanent		Osteochondrosis	Professional	Valproate) (Sodium			
Impairment/Damage				Valproate)	PS		ORAL
3 DOSAGE							
FORMS, 2 IN 1							
D, PER ORAL							
				Gabapentin	SS		ORAL
2 DOSAGES							
FORMS, 1 IN 1							

D, PER ORAL

Date:02/11/04ISR Number: 4291187-2Report Type:Periodic
Age: Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0438903A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Neck Pain		Paxil Neurontin	PS SS	Glaxosmithkline	ORAL

Date:02/11/04ISR Number: 4294186-XReport Type:Expedited (15-DaCompany Report #2004002731
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Abdominal Pain Upper Arthralgia Asthenia Chest Pain	Consumer	Dilantin Suspension (Phenytoin Sodium) Neurontin (Gabapentin)	PS SS		ORAL
2800 MG (7 TIMES A DAY), ORAL		Complex Partial Seizures Fall Loss Of Consciousness Pain In Extremity Paraesthesia Skin Discolouration Tremor Vascular Occlusion Weight Decreased		Lamotrigine (Lamotrigine) Atorvastatin (Atorvastatin) Benazepril Hydrochloride (Benzepril	C C		

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Hydrochloride) C
 Escitalopram
 (Escitalopram) C
 All Other
 Therapeutic Products C
 Carbamazepine
 (Carbamazepine) C
 Atenolol (Atenolol) C
 Warfarin (Warfarin) C

Date:02/11/04ISR Number: 4294200-1Report Type:Expedited (15-DaCompany Report #001-0945-M0000544
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract	Consumer	Neurontin			
500 MG		Muscle Spasms		(Gabapentin)	PS		ORAL
(DAILY), ORAL		Vision Blurred					
				Acetylsalicylic Acid			
				(Acetylsalicylic			
				Acid)	C		
				Indometacin			
				(Indometacin)	C		
				Amlodipine Besilate			
				(Amlodipine			
				Besilate)	C		
				Atenolol (Atenolol)	C		
				Nicotinic Acid			
				(Nicotinic Acid)	C		
				Estrogens Conjugated			
				(Estrogens			
				Conjugated)	C		
				Magnesium			
				(Magnesium)	C		
				Levothyroxine Sodium			
				(Levothyroxine			
				Sodium)	C		

Date:02/12/04ISR Number: 4293808-7Report Type:Expedited (15-DaCompany Report #PHNU2004DE00766
 Age: Gender:Male I/FU:I

Outcome PT

Other

Abnormal Behaviour
Alopecia
Bradyphrenia
Condition Aggravated
Coordination Abnormal
Depression
Difficulty In Walking
Disturbance In Attention
Dyskinesia
Erectile Dysfunction
Fall
Fatigue
Gait Disturbance
Memory Impairment
Mental Impairment
Motor Dysfunction
Movement Disorder
Muscle Spasms
Muscle Twitching

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Dose	Duration	Myalgia Neuroleptic Malignant Syndrome	Report Source	Product	Role	Manufacturer	Route
1600mg/day		Paralysis Parkinson'S Disease	Consumer	Tegretal	PS	Novartis Sector: Pharma	ORAL
UNKNOWN		Thinking Abnormal		Zentropil	SS		
UNKNOWN		Tremor		Ergenyl "Sanofi-Synthelabo"	SS		
UNKNOWN				Neurontin	SS		
UNKNOWN				Lamictal "Glaxosmithkline"	SS		
UNKNOWN				Gabitril	SS		
UNKNOWN				Topamax	SS		
UNKNOWN				Keppra	SS		
UNKNOWN				Carbabetta	SS		
UNKNOWN				Nacom "Dupont Pharma"	SS		
UNKNOWN				Movergan	SS		
UNKNOWN				Madopar	SS		
UNKNOWN				Levodopa	SS		
UNKNOWN				Amantadin	SS		
UNKNOWN				Comtess	SS		
UNKNOWN				Parkotil	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hepatic Steatosis Hepatocellular Damage	Foreign Literature	Neurontin (Gabapentin)	PS		ORAL
Other 900 MG (300 MG, TID),		Jaundice Cholestatic	Health Professional				

Human Mixtard (Insulin Human, Isophane)	C
Metformin (Metformin)	C
Amitriptyline (Amitriptyline)	C
Dihydrocodeine (Dihydrocodeine)	C
Ramipril (Ramipril)	C

Date:02/13/04ISR Number: 4296968-7Report Type:Expedited (15-DaCompany Report #2004007389
Age:82 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2400 MG (TID), ORAL		Condition Aggravated Dry Mouth Epilepsy Glossitis Somnolence	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/13/04ISR Number: 4297118-3Report Type:Expedited (15-DaCompany Report #2004007885
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG		Convulsion					
(TID), ORAL		Vaginal Haemorrhage					
				Levetiracetam (Levetiracetam)	C		
				Fluoxetine (Fluoxetine)	C		

Date:02/16/04ISR Number: 4295011-3Report Type:Expedited (15-DaCompany Report #PHNU2004DE00766
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Tegretal	PS	Novartis Sector: Pharma	ORAL
400 mg, QOD		Alopecia					
		Anosmia		Ergenyl			
UNKNOWN		Bradyphrenia		"Sanofi-Synthelabo"	SS		
		Chills		Neurontin	SS		
		Condition Aggravated		Gabitril	SS		ORAL
60mg/day		Coordination Abnormal		Topamax	SS		ORAL
100mg/day	549 DAY	Depression		Carbabeta	SS		ORAL
400 mg, QID		Difficulty In Walking		Madopar	SS		ORAL
62.5 mg, TID		Disturbance In Attention		Amantadin	SS		ORAL
		Drug Interaction		Parkotil	SS		ORAL
0.20mg/day		Dyskinesia		Parkotil	SS		ORAL
1mg/day		Erectile Dysfunction		Comtess	SS		ORAL
200 mg, QID		Fall		Levodopa	SS		ORAL

1DF/day		Fatigue	Movergan	SS	ORAL
		Gait Disturbance	Zentropil	C	
UNKNOWN	3DF/Day	Grand Mal Convulsion	Selegiline	C	
5mg/day		Memory Impairment	Remergil	C	ORAL
15mg/day	366 DAY	Mental Impairment	Tasmar	C	
		Motor Dysfunction	Keppra	C	
UNKNOWN	1000 -	Movement Disorder			
1500mg/day		Muscle Spasms	Nacom "Dupont Pharma"	C	
UNKNOWN		Muscle Twitching			
		Myalgia	Phenhydan	C	
		Nerve Conduction Studies Abnormal	Lamictal "Glaxosmithkline"	C	
UNKNOWN		Neuroleptic Malignant Syndrome			
		Paralysis			
		Parkinson'S Disease			
		Salivary Hypersecretion			
		Simple Partial Seizures			
		Tremor			

Date:02/16/04ISR Number: 4295766-8Report Type:Periodic Company Report #US-GLAXOSMITHKLINE-A0432317A
Age:44 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
25MG Twice		Pregnancy		Lamictal	PS	Glaxosmithkline	ORAL
per day				Depakote Er	SS		ORAL
150MG Unknown				Effexor	SS		ORAL
150MG Per day				Neurontin	SS		ORAL
100MG Per day				Lorazepam	SS		ORAL
.5MG As							

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required

Date:02/17/04ISR Number: 4299115-0Report Type:Expedited (15-DaCompany Report #2004UW02477
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coronary Artery Atherosclerosis Therapeutic Agent Toxicity	Health Professional	Seroquel Neurontin Lexapro Clonazepam	PS SS SS SS		

Date:02/17/04ISR Number: 4299176-9Report Type:Expedited (15-DaCompany Report #2004007120
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other (DAILY), ORAL		Agitation Depression Intervertebral Disc Degeneration	Consumer	Neurontin (Gabapentin) Tramadol Hydrochloride (Tramadol Hydrochloride) Propacet (Paracetamol, Dextropropoxyphene Napsilate) Provella-14 (Medroxyprogesterone Acetate, Estrogens Conjugated) Antidepressants	PS C C C		ORAL

Date:02/17/04ISR Number: 4299218-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
 Age:44 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability	Abdominal Pain Upper Animal Bite Anorexia

Other

Asthenia
Bronchitis
Cellulitis
Cough
Cyst Rupture
Depression
Drug Hypersensitivity
Dry Mouth
Dysphagia
Dysuria
Ear Pain
Facial Pain
Fatigue
Granuloma
Headache
Hypothyroidism
Insomnia
Laceration
Localised Infection
Loss Of Consciousness
Mania

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Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1800 MG	(DAILY), ORAL	Nausea Ovarian Mass Pelvic Inflammatory Disease	Neurontin (Gabapentin)	PS		ORAL
		Pharyngolaryngeal Pain				
		Professional				
			Lamotrigine (Lamotrigine)	C		
			Lithium Carbonate (Lithium Carbonate)	C		
			Clonazepam (Clonazepam)	C		
			Methylphenidate Hydrochloride (Methylphenidate Hydrochloride)	C		
			Levothyroxine Sodium (Levothyroxine Sodium)	C		
			Liothyronine Sodium (Liothyronine Sodium)	C		
			Sertraline Hydrochloride (Sertraline Hydrochloride)	C		
			Pilocarpine Hydrochloride (Pilocarpine Hydrochloride)	C		
			Metoprolol Succinate (Metoprolol Succinate)	C		
			Lansoprazole (Lansoprazole)	C		
			Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C		
			Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C		
			Bupropion Hydrochloride			

(Bupropion Hydrochloride)	C
Celebrex (Celecoxib)	C
Quetiapine Fumarate (Quetiapine Fumarate)	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
Olanzapine (Olanzapine)	C
Omeprazole (Omeprazole)	C

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Nasal Preparations	C
Nefazodone	
Hydrochloride	
(Nefazodone	
Hydrochloride)	C
Totolin	
(Guaifenesin,	
Phenylpropanolamine	
Hydrochloride)	C
Meclozine	
(Meclozine)	C
Fluticasone	
Propionate	
(Fluticasone	
Propionate)	C
Trimethobenzamide	
Hydrochloride	
(Trimethobenzamide	
Hydrochloride)	C
Respaire-Sr-120	
(Guaifenesin,	
Pseudoephedrine	
Hydrochloride)	C
Risperidone	
(Risperidone)	C

Date:02/17/04ISR Number: 4299223-4Report Type:Expedited (15-DaCompany Report #2003124521
 Age:47 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Aggression	Health	Neurontin			
Initial or Prolonged	Condition Aggravated	Professional	(Gabapentin)	PS		ORAL
900 MG (TID),						
Disability	Deafness					
ORAL						
Other	Dysgraphia		Fluoxetine			
	Dysphagia		Hydrochloride			
	Fatigue		(Fluoxetine			
	Fear		Hydrochloride)	C		
	Hearing Impaired		Fentanyl (Fentanyl)	C		
	Injury		Clonazepam			
	Neuralgia		(Clonazepam)	C		
	Pain		Hydromorphone			
	Panic Disorder		Hydrochloride			

Sluggishness
Suicide Attempt
Vision Blurred
Weight Increased

(Hydromorphone
Hydrochloride) C
Morphine Sulfate
(Morphine Sulfate) C

Date:02/17/04ISR Number: 4299248-9Report Type:Expedited (15-DaCompany Report #2004008023

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3000 MG Other (DAILY)		Leukaemia Pancreatitis	Health Professional	Neurontin (Gabapentin)	PS		

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Date:02/17/04ISR Number: 4299258-1Report Type:Expedited (15-DaCompany Report #2004007906
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Mouth	Consumer	Neurontin			
		Tooth Extraction		(Gabapentin)	PS		
		Toothache		Methadone			
				(Methadone)	C		
				Venlafaxine			
				Hydrochloride			
				(Venlafaxine			
				Hydrochloride)	C		
				Nabumetone			
				(Nabumetone)	C		
				Senna Fruit (Senna			
				Fruit)	C		
				Lidocaine Infusion	C		
				Cilest			
				(Ethinylestradiol,			
				Norgestimate)	C		

Date:02/17/04ISR Number: 4299290-8Report Type:Expedited (15-DaCompany Report #2004002059
Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Abasia	Health	Doxepin (Caps)			
Initial or Prolonged		Accidental Overdose	Professional	(Doxepin)	PS		ORAL
300 MG							
Other		Coma					
(DAILY), ORAL							
		Concussion		Neurontin			
		Dyskinesia		(Gabapentin)	SS		ORAL
900 MG							
(DAILY), ORAL		Electroencephalogram					
		Abnormal		Phenobarbital			
		Epilepsy		(Phenobarbital)	SS		ORAL
30 MG (BID),							
ORAL		Facial Bones Fracture					
		Fall		Levothyroxine Sodium			
		Fatigue		(Levothyroxine			
		Head Injury		Sodium)	C		

Limb Operation	Omeprazole	
Memory Impairment	(Omeprazole)	C
Mobility Decreased	Verapamil	
Myoclonus	Hydrochloride	
Neck Pain	(Verapamil	
Overdose	Hydrochloride)	C
Pain	Temazepam	
Scar	(Temazepam)	C
Shoulder Operation	All Other	
Sinus Disorder	Therapeutic Products	C
Sinusitis	All Other	
Somnolence	Therapeutic Products	C
Thoracic Vertebral	Baclofen (Baclofen)	C
Fracture	Carisoprodol	
	(Carisoprodol)	C
	Parafon Forte	
	(Chlorzoxazone,	
	Paracetamol)	C
	Hydroxyzine Embonate	
	(Hydroxyzine	
	Embonate)	C
	Prochlorperazine	
	Edisylate	
	(Prochlorperazine	

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Edisylate)	C
Sumatriptan Succinate (Sumatriptan Succinate)	C
Dyazide (Hydrochlorothiazide , Triamterene)	C
Pravastatin Sodium (Pravastatin Sodium)	C
Tegaserod (Tegaserod)	C
Percodan (Acetylsalicylic Acid, Caffeine, Phenacetin, Oxycodone Vicodin (Paracetamol, Hydrocodone Bitartrate)	C

Date:02/17/04ISR Number: 4299457-9Report Type:Expedited (15-DaCompany Report #2003039498
 Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Phenobarbital Sodium (Phenobarbital Sodium)	PS		ORAL
Other		Disease Recurrence Drug Interaction Motor Dysfunction			SS		
150 MG							
(DAILY), ORAL				Baclofen Paracetamol	C		

Date:02/17/04ISR Number: 4300541-1Report Type:Expedited (15-DaCompany Report #2004003778
 Age:21 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Asthenia	Foreign	Neurontin			

Initial or Prolonged	Epilepsy	Consumer	(Gabapentin)	PS
1600 MG (BID)				
Other	Headache	Health	Hyrdocortisone	
	Meningitis	Professional	(Hydrocortisone)	C
	Platelet Count Decreased		Paracetamol	
	Post Procedural		(Paracetamol)	C
	Complication		Homeopathic	
	Somnolence		Preparation	C
	Staphylococcal Infection			
	Tinnitus			
	Urinary Incontinence			
	Vertigo			
	Weight Increased			

Date:02/18/04ISR Number: 4300194-2Report Type:Expedited (15-DaCompany Report #2003036100
Age:64 YR Gender:Male I/FU:I

Outcome	PT
Other	Allodynia
	Burning Sensation

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Dose	Duration	Condition	Report Source	Product	Role	Manufacturer	Route
900 MG	(DAILY), ORAL	Chronic Lymphocytic Leukaemia Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
400 MG	(DAILY)	Paraesthesia Post Herpetic Neuralgia Urticaria	Professional Company Representative	Chlorambucil (Chlorambucil)	SS		

Date:02/18/04ISR Number: 4300330-8Report Type:Expedited (15-DaCompany Report #2003038198
Age:78 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID)		Dysphagia Face Oedema	Foreign Health	Neurontin (Gabapentin)	PS		
		Gastrointestinal Disorder	Professional Company Representative	Glyceryl Trinitrate Gliclazide Tiapride Alprazolam Panadeine Co (Codeine Phosphate, Paracetamol) Zolpidem Citalobram Hydrobromide	C C C C C C C C		

Date:02/18/04ISR Number: 4300338-2Report Type:Expedited (15-DaCompany Report #2003126147
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aggression	Consumer	Neurontin			

900 MG (TID),	Anger	Health	(Gabapentin)	PS	ORAL
ORAL	Anxiety	Professional			
	Condition Aggravated		Diazepam (Diazepam)	C	
	Diarrhoea				
	Drug Dependence				
	Drug Withdrawal Syndrome				
	Faeces Discoloured				
	Heart Rate Abnormal				
	Homicidal Ideation				
	Hyperhidrosis				
	Limb Injury				
	Muscle Spasms				
	Nausea				
	Oesophageal Disorder				
	Pain				
	Paraesthesia				
	Respiratory Tract				
	Infection				
	Sensory Disturbance				
	Suicidal Ideation				
	Vomiting				

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Date:02/18/04ISR Number: 4300345-XReport Type:Expedited (15-DaCompany Report #2003114753

Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bone Neoplasm Malignant	Consumer	Neurontin			
Hospitalization -		Lung Cancer Metastatic	Health	(Gabapentin)	PS		
Initial or Prolonged		Lung Carcinoma Cell Type	Professional	Omeprazole			
Other		Unspecified Stage Iv		(Omeprazole)	C		
		Pulmonary Oedema		Clonazepam			
		Treatment Noncompliance		(Clonazepam)	C		

Date:02/18/04ISR Number: 4300499-5Report Type:Expedited (15-DaCompany Report #001-0945-M0200657

Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Acute Sinusitis	Consumer	Neurontin			
Initial or Prolonged		Astigmatism	Health	(Gabapentin)	PS		ORAL
1800 MG							
Disability		Blepharitis	Professional				
(DAILY), ORAL							
Other		Blepharospasm		Lamotrigine			
		Brain Damage		(Lamotrigine)	C		
		Bronchospasm		Lithium Carbonate			
		Burning Sensation		(Lithium Carbonate)	C		
		Depression		Clonazepam			
		Diplopia		(Clonazepam)	C		
		Dizziness		Methylphenidate			
		Drug Hypersensitivity		Hydrochloride			
		Drug Level Increased		(Methylphenidate			
		Ear Pain		Hydrochloride)	C		
		Eyelid Ptosis		Levothyroxine Sodium			
		Facial Bones Fracture		(Levothyroxine			
		Fall		Sodium)	C		
		Hordeolum		Liothyronine Sodium			
		Hypersensitivity		(Liothyronine			
		Keratoconjunctivitis		Sodium)	C		
		Sicca		Sertraline			
		Lid Lag		Hydrochloride			
		Myopia		(Sertraline			
		Photophobia		Hydrochloride)	C		
		Photopsia		Pilocarpine			
		Prescribed Overdose		Hydrochloride			
		Road Traffic Accident		(Pilocarpine			

Sensation Of Pressure
Speech Disorder
Swollen Tongue
Vestibular Neuronitis

Hydrochloride) C
Metoprolol Succinate
(Metoprolol
Succinate) C
Lansoprazole
(Lansoprazole) C
Hyoscyamine Sulfate
(Hyoscyamine
Sulfate) C
Diltiazem
Hydrochloride
(Diltiazem
Hydrochloride) C
Bupropion
Hydrochloride
(Bupropion
Hydrochloride) C
Celebrex (Celecoxib) C
Quetiapine Fumarate
(Quetiapine

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Fumarate)	C
Metoprolol Tartrate	
(Metoprolol	
Tartrate)	C
Citalopram	
Hydrobromide	
(Citalopram	
Hydrobromide)	C
Hyoscyamine Sulfate	
(Hyoscyamine	
Sulfate)	C
Olanzapine	
(Olanzapine)	C
Omeprazole	
(Omeprazole)	C
Nasal Preparations	C
Nefazodone	
Hydrochloride	
(Nefazodone	
Hydrochloride)	C
Totolin	
(Guaifenesin,	
Phenylpropanolamine	
Hydrochloride)	C
Meclozine	
(Meclozine)	C
Fluticasone	
Propionate	
(Fluticasone	
Propionate)	C
Trimethobenzamide	
Hydrochloride	
(Trimethobenzamide	
Hydrochloride)	C
Respaire-Sr-120	
(Guaifenesin,	
Pseudoephedrine	
Hydrochloride)	C
Risperidone	
(Risperidone)	C

Date:02/20/04ISR Number: 4302034-4Report Type:Expedited (15-DaCompany Report #2004008693

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					

Other	Dizziness	Foreign	Neurontin	
900 MG TIC	Fall	Consumer	(Gabapentin)	PS
	Hypoaesthesia		Oral Contraceptives	
	Incontinence		Nos	C
	Loss Of Consciousness		Omeprazole	
			(Omeprazole)	C

Date:02/20/04ISR Number: 4302051-4Report Type:Expedited (15-DaCompany Report #2004003992
Age:29 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Asthenia
Initial or Prolonged	Muscle Spasms
Other	Muscle Twitching
	Pollakiuria

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Dose	Duration	Sensory Disturbance Urinary Incontinence	Report Source	Product	Role	Manufacturer	Route
ORAL			Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:02/20/04ISR Number: 4302313-0Report Type:Expedited (15-DaCompany Report #2004008953
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Cardiac Arrest	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
Other			Diarrhoea					
ORAL			Face Oedema Medication Error		All Other Therapeutic Products	SS		OTHER
OTHER			Oedema		All Other Therapeutic Products	C		

Date:02/20/04ISR Number: 4302522-0Report Type:Expedited (15-DaCompany Report #2004001234
Age: Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Pruritus Rash	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL			Skin Disorder		Vitamins Estrogens Conjugated (Estrogens Conjugated)	C		

Date:02/20/04ISR Number: 4302646-8Report Type:Expedited (15-DaCompany Report #2004008949
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1200 MG	Abnormal Dreams Aggression	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other (QID), ORAL		Anorexia					
		Chapped Lips		Bupropion			
		Cough		Hydrochloride			
		Disease Recurrence		(Bupropion			
		Dizziness		Hydrochloride)	C		
		Drug Effect Decreased		Diphenhydramine			
		Dry Mouth		Hydrochloride			
		Eating Disorder		(Diphehydramine			
		Homicidal Ideation		Hydrochloride)	C		
		Insomnia		Valproate Sodium			
		Myocardial Infarction		(Valproate Sodium)	C		
		Nightmare		Valproate Semisodium			
		Tremor		(Valproate			
		Weight Increased		Semisodium)	C		

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Date:02/20/04ISR Number: 4302961-8Report Type:Expedited (15-DaCompany Report #2003008068

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Activities Of Daily	Consumer	Neurontin			
Initial or Prolonged	Living Impaired	Health	(Gabapentin)	PS		
Disability	Aplastic Anaemia	Professional	Dilantin (Phenytoin			
Other	Arthralgia		Soidum)	SS		ORAL
1200 MG						
(TID), ORAL	Balance Disorder					
	Difficulty In Walking		Acetylsalicylic Acid			
	Dry Mouth		(Acetylsalicylic			
	Fall		Acid)	SS		
	Hypotonia		Vitamins With			
	Muscle Spasms		Minerals	SS		
	Pain		Megestrol			
	Pain In Extremity		(Megestrol)	C		
	Paralysis					
	Red Blood Cell Count					
	Increased					

Date:02/23/04ISR Number: 4303086-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200657

Age:44 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abnormal Sensation In Eye
Initial or Prolonged	Activities Of Daily
Disability	Living Impaired
Other	Acute Sinusitis
	Anger
	Antinuclear Antibody
	Positive
	Anxiety
	Aphasia
	Arthritis
	Back Pain
	Biopsy Tongue Abnormal
	Blepharitis
	Blepharospasm
	Blood Glucose Increased
	Brain Damage
	Bronchospasm
	Bruxism

Burning Sensation
Chest Pain
Coordination Abnormal
Depression
Diplopia
Dizziness
Drug Withdrawal Syndrome
Ear Pain
Eye Irritation
Eyelid Ptosis
Facial Bones Fracture
Fall
Feeling Jittery
Haemorrhage
Halo Vision
Head Injury
Hordeolum
Hypersensitivity
Insomnia

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Dose	Duration	Intentional Self-Injury Intervertebral Disc Degeneration	Report Source	Product	Role	Manufacturer	Route
1800 MG		Irritability Joint Sprain	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL		Keratoconjunctivitis	Professional				
		Sicca		Lamotrigine (Lamotrigine)	C		
		Loss Of Consciousness		Lithium Carbonate (Lithium Carbonate)	C		
		Neck Injury		Clonazepam (Clonazepam)	C		
		Nystagmus		Methylphenidate Hydrochloride			
		Pain		(Methylphenidate Hydrochloride)	C		
		Pco2 Decreased		Levothyroxine Sodium (Levothyroxine Sodium)	C		
		Photophobia		Liothyronine Sodium (Liothyronine Sodium)	C		
		Restless Legs Syndrome		Sertraline Hydrochloride			
		Road Traffic Accident		(Sertraline Hydrochloride)	C		
		Sensation Of Pressure		Pilocarpine Hydrochloride			
		Sensory Disturbance		(Pilocarpine Hydrochloride)	C		
		Speech Disorder		Metoprolol Succinate (Metoprolol Succinate)	C		
		Suicidal Ideation		Lansoprazole (Lansoprazole)	C		
		Swollen Tongue		Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C		
		Vestibular Neuritis		Diltiazem Hydrochloride			
		Viral Infection		(Diltiazem Hydrochloride)	C		
		White Blood Cell Count Abnormal		Bupropion Hydrochloride			

(Bupropion Hydrochloride)	C
Celebrex (Celecoxib)	C
Quetiapine Fumarate (Quetiapine Fumarate)	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C
Olanzapine	

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(Olanzapine)	C
Omeprazole	
(Omeprazole)	C
Nasal Preparations	C
Nefazodone	
Hydrochloride	
(Nefazodone	
Hydrochloride)	C
Totolin	
(Guaifenesin,	
Phenylpropanolamine	
Hydrochloride)	C
Meclozine	
(Meclozine)	C
Fluticasone	
Propionate	
(Fluticasone	
Propionate)	C
Trimethobenzamide	
Hydrochloride	
(Trimethobenzamide	
Hydrochloride)	C
Respaire-Sr-120	
(Guaifenesin,	
Pseudoephedrine	
Hydrochloride)	C
Risperidone	
(Risperidone)	C

Date:02/23/04ISR Number: 4303224-7Report Type:Expedited (15-DaCompany Report #2004009355

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Abdominal Pain	Consumer	Neurontin			
Initial or Prolonged	Back Disorder		(Gabapentin)	PS		ORAL
1800 MG						
Other	Back Pain					
(TID), ORAL						
	Blood Cholesterol		Lipitor			
	Increased		(Atorvastatin)	SS		ORAL
20 MG						
	Chest Pain					
(DAILY), ORAL						
	Drug Effect Decreased		Lithium (Lithium)			
	Drug Intolerance		(Lithium)	SS		

	Erectile Dysfunction	Naproxen (Naproxen)	SS	
	Facial Pain	Rofecoxib		
	Feeling Cold	(Rofecoxib)	SS	
	Gastrooesophageal Reflux Disease	Amoxicillin		
		(Amoxicillin)	SS	ORAL
1000 MG	Hypersomnia			
(BID), ORAL	Medication Error	All Other		
	Neck Pain	Therapeutic Products	C	
	Pain In Jaw	Levothyroxine Sodium		
	Sleep Disorder	(Levothyroxine		
	Somnolence	Sodium)	C	
	Tremor	Vitamins	C	
		Diltiazem		
		Hydrochloride		
		(Diltiazem		
		Hydrochloride)	C	

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Freedom Of Information (FOI) Report

Date:02/23/04ISR Number: 4303334-4Report Type:Expedited (15-DaCompany Report #KII-2003-0007131
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Blood Potassium Decreased Blood Urea Increased Chromaturia Confusional State Depressed Level Of Consciousness Dysarthria	Study Health Professional Other	Morphine Sulfate (Similar To Nda 19-516) (Morphine Sulfate)	PS		ORAL
30 MG, QID, ORAL				Neurontin (Gabapentin)	SS		ORAL
300 MG, SEE TEXT, ORAL		Hypotension					
0.5 MG, QID, ORAL		Hypoxia Lethargy Medication Error		Clonazepam (Clonazepam)	SS		ORAL
QID, ORAL	2 DAY	Mydriasis Non-Cardiogenic Pulmonary Oedema Pneumonia Pruritus Tachypnoea		Benadryl (Diphenhydramine Hydrochloride) Acetylsalicylic Acid (Acetylsalicylic Acid)	SS SS		ORAL ORAL

Date:02/23/04ISR Number: 4303431-3Report Type:Expedited (15-DaCompany Report #KII-2003-0007053
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other ORAL		Abnormal Behaviour Blood Alcohol Increased Blood Calcium Decreased Body Temperature Decreased	Study Health Professional Other	Ms Contin Tablets (Morphine Sulfate) Cr Tablet Neurontin (Gabapentin)	PS SS		ORAL ORAL
ORAL		Bowel Sounds Abnormal Coma		Amitriptyline (Amitriptyline)	SS		ORAL

Confusional State	Phencyclidine	
Depressed Mood	(Phencyclidine)	SS
Hypotension	Alcohol (Ethanol)	SS
Multiple Drug Overdose		
Mydriasis		
Pupillary Reflex Impaired		
Respiratory Depression		
Suicidal Ideation		
Tachycardia		
Urine Output Increased		

Date:02/23/04ISR Number: 4303459-3Report Type:Expedited (15-DaCompany Report #2004009410

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Arrhythmia	Consumer	Neurontin			
Initial or Prolonged	Benign Prostatic		(Gabapentin)	PS		ORAL
ORAL						
Other	Hyperplasia		Venlafaxine			
	Micturition Urgency		Hydrochloride			
	Pollakiuria		(Venlafaxine			
	Residual Urine Volume		Hydrochloride)	C		
	Urinary Incontinence		Buspirone			
			Hydrochloride			
			(Buspirone			
			Hydrochloride)	C		
			Testosterone			
			(Testosterone)	C		

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Date:02/23/04ISR Number: 4303474-XReport Type:Expedited (15-DaCompany Report #2004006906

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Renal Impairment	Consumer Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/23/04ISR Number: 4303560-4Report Type:Expedited (15-DaCompany Report #2004009451

Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Epilepsy	Foreign Health Professional	Neurontin (Gabapentin)	PS		
3200 MG		Overdose					

Date:02/23/04ISR Number: 4303645-2Report Type:Expedited (15-DaCompany Report #2004009409

Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Thyroid Operation Thyroidectomy	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
600 MG				Alendronate Sodium (Alendronate Sodium)	C		
Other (DAILY), ORAL				Fibrinolysin (Fibrinolysin)	C		

Date:02/23/04ISR Number: 4304175-4Report Type:Expedited (15-DaCompany Report #FRWYE593019FEB04

Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged Other	Bradyarrhythmia Electroencephalogram Abnormal Fall Headache	Health Professional Other	Effexor Lp (Venlafaxine Hydrochloride, Capsule, Extended Release, 0)	PS	ORAL
37.5 MG 2X					
PER 1 DAY	Metabolic Disorder				
ORAL	Nervous System Disorder				
1 TABLET 1X	Vertigo		Aprovel (Irbesartan, ,0)	SS	ORAL
PER 1 DAY					
ORAL					
2 TABLET 1X			Contramal (Tramadol Hydrochloride, ,0)	SS	ORAL
PER 1 DAY					
ORAL					
1 DOSE 1X PER			Furosemide (Furosemide, ,0)	SS	ORAL
1 DAY ORAL					
2 DOSE 1X PER			Neurontin (Gabapentin, ,0)	SS	ORAL
1 DAY ORAL					
			Amlor (Amlodipine Besilate)	C	
			Glucor (Acarbose)	C	
			Ogast (Lansoprazole)	C	

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Diffrarel
 (Betacarotene/Myrtil
 lus) C
 Dafalgan
 (Paracetamol) C
 Vastarel
 (Trimetazidine
 Hydrochloride) C

Date:02/23/04ISR Number: 4304193-6Report Type:Expedited (15-DaCompany Report #FRWYE593019FEB04
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other	37.5 MG 2X PER 1 DAY ORAL	Bradyarrhythmia Electroencephalogram Abnormal Fall Headache	Foreign Health Professional Other	Effexor Lp (Venlafaxine Hydrochloride, Capsule, Extended Release, 0)	PS		ORAL
		Metabolic Disorder Nervous System Disorder Vertigo		Aprovel (Irbesartan, ,0)	SS		ORAL
	1 TABLET 1X PER 1 DAY ORAL			Contramal (Tramadol Hydrochloride, ,0)	SS		ORAL
	2 TABLET 1X PER 1 DAY ORAL			Furosemide (Furosemide, ,0)	SS		ORAL
	1 DOSE 1X PER 1 DAY ORAL			Neurontin			

2 DOSE 1X PER

(Gabapentin, ,0) SS

ORAL

1 DAY ORAL

Amlor (Amlodipine Besilate)	C
Glucor (Acarbose)	C
Ogast (Lansoprazole)	C
Difrarel (Betacarotene/Myrtilus)	C
Dafalgan (Paracetamol)	C
Vastarel (Trimetazidine Hydrochloride)	C

Date:02/24/04ISR Number: 4304067-0Report Type:Expedited (15-DaCompany Report #2002051362

Age:55 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Eosinophilia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG	Generalised Erythema					
Other (TID), ORAL	Hypersensitivity	Professional				
	Rash Maculo-Papular					
	Renal Failure Acute					
	Thrombocytopenia					

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Date:02/24/04ISR Number: 4304335-2Report Type:Expedited (15-DaCompany Report #2004005558

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Consumer Health	Dilantin Kapseals (Phenytoin Sodium)	PS		ORAL
400 MG QID, ORAL			Professional				
				Neurontin (Gabapentin)	SS		ORAL
900 MG TID, ORAL							
				Primidone (Primidone)	C		
				Carbamazepine (Carbamazepine)	C		
				Ranitidine (Ranitidine)	C		
				Alprazolam (Alprazolam)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		

Date:02/24/04ISR Number: 4304343-1Report Type:Expedited (15-DaCompany Report #2004005000

Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Loss Of Consciousness Syncope	Health Professional	Neurontin (Gabapentin)	PS		
1800 MG (600 MG TID)			Company				
			Representative	Sertraline Hydrochloride (Sertraline Hydrochloride)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Alkaline Phosphatase Increased	Foreign Literature	Neurontin (Gabapentin)	PS		ORAL
300 MG (QID),		Blood Bilirubin Increased	Health				
ORAL		Cholestasis	Professional	Amitriptyline Hydrochloride (Amitriptyline Hydrchloride)	SS		ORAL
300 MG (QID),		Gamma-Glutamyltransferase Increased International Normalised Ratio Decreased					
ORAL				Atenolol (Atenolol)	SS		ORAL
100 MG							
(DAILY), ORAL				Nifedipine (Nifedipine)	SS		ORAL
30 MG							
(DAILY), ORAL				Lisinopril (Lisinopril)	SS		ORAL
5 MG (DAILY),							
ORAL				Pyridoxine (Pyridoxine)	SS		

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Sodium Polystyrene
Sulfonate (Sodium
Polystyrene
Sulfonate) C
Cyanocobalamin
(Cyanocobalamin) C
Thiamine
Hydrochloride
(Thiamine
Hydrochloride) C

Date:02/24/04ISR Number: 4304522-3Report Type:Expedited (15-DaCompany Report #2004010178

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Hepatic Failure	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/24/04ISR Number: 4304527-2Report Type:Expedited (15-DaCompany Report #2004001504

Age:92 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (TID)	Aphasia Confusional State	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
, ORAL	Drug Level Increased	Professional Company Representative	Enalapril (Enalapril)	C		
	Dysarthria Psychotic Disorder Restlessness		Thiamazole (Thiamazole)	C		
			Doxepin (Doxepin)	C		
			Torasemide	C		
			Taurizine (Taurine, Aspartate Zinc)	C		
			Metamizole Sodium (Metamizole Sodium)	C		
			Tramadol			

Hydrochloride
(Tramadol
Hydrochloride) C

Date:02/24/04ISR Number: 4305110-5Report Type:Direct
Age:83 YR Gender:Male I/FU:I

Company Report #CTU 213075

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pain Swelling		Gabapentin	PS		

Date:02/25/04ISR Number: 4303610-5Report Type:Expedited (15-DaCompany Report #FR-BRISTOL-MYERS SQUIBB COMPANY-12511176
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged tablets		Fall Headache Vertigo		Aprovel Tabs Furosemide	PS SS	Bristol-Myers Squibb Company	ORAL ORAL

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Neurontin	SS	ORAL
Contramal	SS	ORAL
Effexor	SS	ORAL
Amlor	C	
Glucor	C	
Ogast	C	
Difrarel	C	
Dafalgan	C	
Vastarel	C	

Date:02/25/04ISR Number: 4303743-3Report Type:Expedited (15-DaCompany Report #PHNU2004DE00766

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 400 mg, QOD		Abnormal Behaviour Alopecia		Tegretal	PS	Novartis Sector: Pharma	ORAL
UNKNOWN		Anosmia Bradyphrenia		Ergenyl "Sanofi-Synthelabo"	SS		
UNKNOWN		Chills		Neurontin	SS		
60mg/day		Condition Aggravated		Gabitril	SS		ORAL
100mg/day	549 DAY	Depression		Topamax	SS		ORAL
400 mg, QID		Difficulty In Walking		Carbabetta	SS		ORAL
62.5 mg, TID		Disturbance In Attention		Madopar	SS		ORAL
0.20mg/day		Dyskinesia Erectile Dysfunction		Amantadin Parkotil	SS SS		ORAL ORAL
1mg/day		Fall		Parkotil	SS		ORAL
200 mg, QID		Fatigue		Comtess	SS		ORAL
1DF/day		Gait Disturbance Grand Mal Convulsion		Levodopa Movergan	SS SS		ORAL ORAL
UNKNOWN	3DF/Day	Memory Impairment		Zentropil	C		
5mg/day		Mental Impairment		Selegiline	C		

15mg/day	366 DAY	Motor Dysfunction	Remergil	C	ORAL
		Movement Disorder	Tasmar	C	
		Muscle Twitching	Keppra	C	
UNKNOWN	1000 -	Myalgia			
1500mg/day		Nerve Conduction Studies	Nacom "Dupont		
		Abnormal	Pharma "	C	
UNKNOWN		Neuroleptic Malignant	Phenhydan	C	
		Syndrome	Lamictal		
		Paralysis	"Glaxosmithkline"	C	
UNKNOWN		Parkinson'S Disease			
		Salivary Hypersecretion			
		Simple Partial Seizures			
		Tremor			

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
		Arthralgia		Wellbutrin	PS	Glaxosmithkline	ORAL
300MG Per day		Joint Swelling		Neurontin	SS		

Date:02/26/04ISR Number: 4305636-4Report Type:Periodic Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12461349
Age:56 YR Gender:Female I/FU:I

Outcome
Hospitalization -
Initial or Prolonged

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Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1/2 of a 10 mg tablet		Acidosis Convulsion Depressed Level Of Consciousness	Health Professional	Abilify Tabs 10 Mg	PS	Otsuka Pharmaceutical Company, Ltd.	ORAL
1/2 of a 10 mg tablet		Hypotension Overdose Respiratory Depression Suicide Attempt		Abilify Tabs 10 Mg	SS	Otsuka Pharmaceutical Company, Ltd.	ORAL
				Tramadol Neurontin	SS SS		ORAL ORAL

Date:02/26/04ISR Number: 4307311-9Report Type:Expedited (15-DaCompany Report #2003120047
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300, Other TID), ORAL		Blood Albumin Decreased Blood Albumin Increased Blood Bicarbonate Increased Blood Creatinine	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
TID		Increased Blood Glucose Increased Blood Phosphorus Increased Blood Pressure Systolic Increased Blood Urea Increased Bundle Branch Block Right Cardiac Murmur Carotid Bruit Disorientation Dyspnoea Electrocardiogram St		Minerals Nos (Minerals Nos) Amlodipine Besilate (Amlodipine Besilate) Furosemide (Furosemide) Doxazosin (Doxazosin) Metoprolol Succinate (Metoprolol Succinate) Hydrochlorothiazide (Hydrochlorothiazide)	SS C C C C C		

Segment Abnormal
Feeling Abnormal
Haematocrit Decreased
Haemoglobin Decreased
Headache
Hypercalcaemia
Lymphocyte Count
Decreased
Myocardial Infarction
Nausea
Pain In Extremity
Protein Total Decreased
Rales
Red Blood Cell
Sedimentation Rate
Increased
Renal Failure Acute
Thinking Abnormal
Thrombocytopenia

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Date:02/26/04ISR Number: 4307315-6Report Type:Expedited (15-DaCompany Report #2004010452

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Respiratory Arrest	Consumer	Neurontin (Gabapentin)	PS		

Date:02/26/04ISR Number: 4307319-3Report Type:Expedited (15-DaCompany Report #2004009355

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Abdominal Pain Back Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
1800 MG Other (TID), ORAL		Back Pain					
20 MG		Blood Cholesterol Increased		Lipitor (Atorvastatin)	SS		ORAL
(DAILY), ORAL		Chest Pain					
		Drug Ineffective		Lithium (Lithium)			
		Drug Intolerance		(Lithium)	SS		
		Erectile Dysfunction		Naproxen (Naproxen)	SS		
		Facial Pain		Rofecoxib			
		Feeling Cold		(Rofecoxib)	SS		
		Gastroesophageal Reflux Disease		Amoxicillin (Amoxicillin)	SS		ORAL
1000 MG		Glossodynia					
(BID), ORAL		Hypersomnia		All Other			
		Medication Error		Therapeutic Products	C		
		Neck Pain		Levothyroxine Sodium			
		Oesophageal Spasm		(Levothyroxine			
		Pain In Jaw		Sodium)	C		
		Sleep Disorder		Vitamins	C		
		Tooth Abscess		Diltiazem			
		Tremor		Hydrochloride (Diltiazem Hydrochloride)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Body Temperature Decreased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID),		Breath Sounds Decreased					
ORAL		Cold Sweat		Paclitaxel (Paclitaxel)	SS		
INTRA	VENOUS	Drug Ineffective					
		Erythema		Pyridoxine Hydrochloride (Pyridoxine Hydrochloride)	C		
		Feeling Abnormal		Simvastatin (Simvastatin)	C		
		Feeling Cold		Verapamil Hydrochloride (Verapamil Hydrochloride)	C		
		Haematocrit Decreased		Sertraline Hydrochloride (Sertraline Hydrochloride)	C		
		Haemoglobin Decreased		Zolpidem Tartrate			
		Hyperhidrosis					
		Peripheral Sensory Neuropathy					
		Rash					
		White Blood Cell Count Increased					

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(Zolpidem Tartrate) C

Date:02/26/04ISR Number: 4307361-2Report Type:Expedited (15-DaCompany Report #2004010456

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abasia Pain In Extremity	Health Professional	Neurontin (Gabapentin)	PS		

Date:02/26/04ISR Number: 4307362-4Report Type:Expedited (15-DaCompany Report #2004009823

Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis Exostosis Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL				Nateglinide (Nateglinide) Insulin (Insulin) Doxepin Hydrochloride Atenolol (Atenolol) Simvastatin (Simvastatin)	C C C C C		

Date:02/26/04ISR Number: 4307385-5Report Type:Expedited (15-DaCompany Report #2004010848

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 600 MG (200 Other MG TID), ORAL		Blood Creatinine Increased Mental Status Changes Myoclonus Oedema Peripheral Somnolence	Health Professional	Gabapentin (Gabapentin) Oxycocet (Paracetamol, Oxycodone	PS		ORAL

TWO TABLETS	Hydrochloride)	SS	ORAL
Q4H, ORAL			
	Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS	ORAL
60 MG (BID),			
ORAL			
62.5 MG BID	Bosentan (Bosentan)	SS	
	Warfarin Sodium (Warfarin Sodium)	C	
	Furosemide (Furosemide)	C	
	Prednisone (Prednisone)	C	
	Epoprostenol Sodium (Epoprostenol Sodium)	C	
	Pantoprazole (Pantoprazole)	C	
	Metformin Hydrochloride (Metformin Hydrochloride)	C	
	Bactrim		

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(Sulfamethoxazole,
Trimethoprim) C
Oxygen (Oxygen) C
Lisinopril
(Lisinopril) C
Metolazone
(Metolazone) C

Date:02/26/04ISR Number: 4307386-7Report Type:Expedited (15-DaCompany Report #2004009821
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG Other (TID), ORAL		Stevens-Johnson Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Lorazepam (Lorazepam)	C		
				Spiroinolactone (Spiroinolactone)	C		
				Prednisone (Prednisone)	C		
				Oseltamivir (Oseltamivir)	C		
				Folic Acid (Folic Acid)	C		
				Lansoprazole (Lansoprazole)	C		
				Lactulose (Lactulose)	C		
				Levofloxacin (Levofloxacin)	C		
				Propranolol Hydrochloride (Propranolol Hydrochloride)	C		
				Magnesium (Magnesium)	C		
				Neutra-Phos (Sodium Phosphate Dibasic, Sodium Phosphate Monobasic (Anhydrate), Potassium Chloride	C		

(Potassium Chloride) C
 Bumetanide
 (Bumetanide) C
 Octreotide Acetate
 (Octreotide Acetate) C
 Phytomenadione
 (Phytomenadione) C

Date:02/26/04ISR Number: 4307387-9Report Type:Expedited (15-DaCompany Report #2004011640
 Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coma Hypothermia	Literature Health Professional	Gabapentin (Gabapentin) Phenobarbital (Phenobarbital) Carbamazepine	PS C		

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(Carbamazepine) C

Date:02/26/04ISR Number: 4307390-9Report Type:Expedited (15-DaCompany Report #2004005558

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Dilantin Kapseals			
		Pharmaceutical Product	Health	(Phenytoin Sodium)	PS		ORAL
400 MG (QID),		Complaint	Professional				
ORAL		Treatment Noncompliance		Neurontin			
				(Gabapentin)	SS		ORAL
900 MG (TID),							
ORAL				Primidone			
				(Primidone)	C		
				Carbamazepine			
				(Carbamazepine)	C		
				Ranitidine			
				(Ranitidine)	C		
				Alprazolam			
				(Alprazolam)	C		
				Levothyroxine Sodium			
				(Levothyroxine			
				Sodium)	C		
				Fluoxetine			
				Hydrochloride			
				(Fluoxetine			
				Hydrochloride)	C		
				Lovastatin			
				(Lovastatin)	C		

Date:02/26/04ISR Number: 4307427-7Report Type:Expedited (15-DaCompany Report #2004010453

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Foreign	Gabapentin			
		International Normalised	Health	(Gabapentin)	PS		ORAL
ORAL							

10 MG
(DAILY), ORAL
Ratio Increased
Professional
Warfarin (Warfarin) SS
ORAL
Other

Date:02/26/04ISR Number: 4307428-9Report Type:Expedited (15-DaCompany Report #2004002702
Age:17 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Amnesia	Foreign	Neurontin			
ORAL		Viral Infection	Health	(Gabapentin)	PS		ORAL
			Professional	Amitriptyline	C		
				(Amitriptyline)			

Date:02/26/04ISR Number: 4307460-5Report Type:Expedited (15-DaCompany Report #2004009875
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Withdrawal	Foreign	Neurontin			
1800 MG TID,		Convulsions	Health	(Gabapentin)	PS		ORAL
ORAL			Professional				

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Date:02/26/04ISR Number: 4307462-9Report Type:Expedited (15-DaCompany Report #2004009792
Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Adrenal Insufficiency	Foreign	Neurontin			
Hospitalization -		Blood Corticotrophin	Health	(Gabapentin)	PS		ORAL
800 MG BID,							
Initial or Prolonged		Decreased	Professional				
ORAL							
Other		Blood Cortisol Decreased		Amiodarone			
		Bradycardia		(Amiodarone)	C		
		Depressed Level Of		Acetylsalicylate			
		Consciousness		Lysine			
		Electrocardiogram Qrs		(Acetylsalicylate			
		Complex Prolonged		Lysine)	C		
		Hypertonia		Hydrocortisone			
		Hypothermia		(Hydrocortisone)	C		
		Myoclonus					

Date:02/26/04ISR Number: 4307487-3Report Type:Expedited (15-DaCompany Report #2004010454
Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Atrophy	Foreign	Neurontin			
Initial or Prolonged		Bradyarrhythmia	Health	(Gabapentin)	PS		ORAL
ORAL							
		Electroencephalogram	Professional	Furosemide			
		Abnormal		(Furosemide)	SS		ORAL
		Fall		Irbesartan			
		Headache		(Irbesartan)	SS		ORAL
		Metabolic Disorder		Venlafaxine			
		Vertigo		Hydrochloride(Venlaf			
				axine Hydrochloride)	SS		ORAL
75 MG , ORAL							
				Tramadol			
				Hydrochloride			
				(Tramadol			
				Hydrochloride)	SS		ORAL
ORAL							
				Amlodipine Besilate			
				(Amlodipine			

Besilate)	C
Acarbose (Acarbose)	C
Lansoprazole	
(Lansoprazole)	C
Diffrarel	
(Betacarotene,	
Myrtillus)	C
Paracetamol	
(Paracetamol)	C
Trimetazidine	
Hydrochloride	
(Trimetazidine	
Hydrochloride)	C

Date:02/26/04ISR Number: 4307536-2Report Type:Expedited (15-DaCompany Report #2003125029
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mycotoxycosis	Consumer	Neurontin (Gabapentin)	PS		
				Zoloft (Sertraline)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/26/04ISR Number: 4307539-8Report Type:Expedited (15-DaCompany Report #2004009909
Age:59 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged 600 MG (TID)	Blood Creatinine Increased	Health Professional	Neurontin (Gabapentin)	PS		
Other	Mental Status Changes Myoclonus Somnolence		Oxycocet (Paracetamol, Oxycodone Hydrochloride) All Other Therapeutic Products	SS SS		

Date:02/26/04ISR Number: 4307540-4Report Type:Expedited (15-DaCompany Report #2003122403
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death	Arrhythmia	Health	Neurontin			
Other	Faecal Incontinence	Professional Company Representative	(Gabapentin)	PS		

Date:02/26/04ISR Number: 4307550-7Report Type:Expedited (15-DaCompany Report #2002062761
Age:72 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged 800 MG (400 MG, BID), UNKNOWN	Hepatic Failure Necrosis	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:02/26/04ISR Number: 4307618-5Report Type:Expedited (15-DaCompany Report #2004010384
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Foreign Health	Neurontin (Gabapentin)	PS		
1600 OR 1800			Professional				
(DAILY)			Company Representative	Clonazepam (Clonazepam)	C		
				Levomepromazine (Levomepromazine)	C		

Date:02/26/04ISR Number: 4307912-8Report Type:Periodic Company Report #2003164678US
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia	Health	Bextra (Valdecoxib)			
UNK, UNK;		Difficulty In Walking	Professional	Tablet	PS		ORAL
ORAL		Flushing					
		Muscle Twitching		Celebrex (Celecoxib,			
UNKNOWN	UNK, UNK, UNK	Myalgia		Celecoxib)Capsule	SS		
		Nausea		Viread (Tenofovir)	SS		
UNKNOWN	300 UNK, UNK,	Renal Failure Acute					
UNK							
		Somnolence		Fortovase			
UNKNOWN	UNK, UNK, UNK	Vomiting		(Saquinavir)	SS		
UNKNOWN	UNK, UNK, UNK			Zerit (Stavudine)	SS		

Freedom Of Information (FOI) Report

UNKNOWN	UNK, UNK, UNK	Kaletra (Lopinavir/Ritonavir)	SS
UNKNOWN	UNK, UNK, UNK	Phenergan (Promethazine)	SS
UNKNOWN	UNK, UNK, UNK	Ambien (Zolpidem Tartrate)	SS
UNKNOWN	UNK, UNK, UNK	Ensure (Vitamins Nos)	SS
UNKNOWN	UNK, UNK, UNK	Tricor (Fenofibrate)	SS
UNKNOWN	UNK, UNK, UNK	Dulcolax (Bisacodyl)	SS
UNKNOWN	UNK, UNK, UNK	Marinol (Dronabinol)	SS
UNKNOWN	UNK, UNK, UNK	Somac (Carisoprodol)	SS
UNKNOWN	UNK, UNK, UNK	Bactrim Ds (Sulfamethoxazole, Trimethoprium)	SS
UNKNOWN	UNK, UNK, UNK	Biaxin (Clarithromycin)	SS
UNKNOWN	UNK, UNK, UNK	Protonix	SS
UNKNOWN	UNK, UNK, UNK	Oxandrin (Oxandrolone)	SS
UNKNOWN	UNK, UNK, UNK	Neurontin (Gabapentin)	SS
UNKNOWN	UNK, UNK, UNK	Colchicine (Colchicine)	SS

Date:02/27/04ISR Number: 4308650-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
Age:44 YR Gender:Female I/FU:F

Outcome PT
Hospitalization - Acute Sinusitis

Initial or Prolonged
Disability
Other

Anger
Antinuclear Antibody
Positive
Anxiety
Aphasia
Arthritis
Back Pain
Blepharitis
Blepharospasm
Brain Damage
Breast Mass
Bruxism
Chest Pain
Chromaturia
Coordination Abnormal
Depression
Diplopia
Dizziness
Drug Withdrawal Syndrome
Dysuria
Ear Pain
Eyelid Ptosis
Facial Bones Fracture
Fall
Gastrooesophageal Reflux
Disease
Head Injury
Hordeolum
Hypersensitivity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Impaired Healing Insomnia Intervertebral Disc Degeneration Keratoconjunctivitis	Report Source	Product	Role	Manufacturer	Route
1800 MG	(DAILY), ORAL	Sicca	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Laryngitis		Lamotrigine (Lamotrigine)	C		
		Loss Of Consciousness		Lithium Carbonate (Lithium Carbonate)	C		
		Lumbar Radiculopathy		Clonazepam (Clonazepam)	C		
		Menorrhagia		Methylphenidate Hydrochloride (Methylphenidate Hydrochloride)	C		
		Nystagmus		Levothyroxine Sodium (Levothyroxine Sodium)	C		
		Ovarian Cyst		Liothyronine Sodium (Liothyronine Sodium)	C		
		Pelvic Inflammatory Disease		Sertraline Hydrochloride (Sertraline Hydrochloride)	C		
		Photophobia		Pilocarpine Hydrochloride (Pilocarpine Hydrochloride)	C		
		Prescribed Overdose		Metoprolol Succinate (Metoprolol Succinate)	C		
		Restless Legs Syndrome		Lansoprazole (Lansoprazole)	C		
		Road Traffic Accident		Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C		
		Self Injurious Behaviour		Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C		
		Sensory Disturbance		Bupropion Hydrochloride			
		Speech Disorder					
		Suicidal Ideation					
		Swollen Tongue					
		Tremor					
		Vestibular Neuronitis					
		Visual Field Defect					
		White Blood Cell Count Abnormal					

(Bupropion Hydrochloride)	C
Celebrex (Celecoxib)	C
Quetiapine Fumarate (Quetiapine Fumarate)	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C
Olanzapine	

Freedom Of Information (FOI) Report

(Olanzapine)	C
Omeprazole	
(Omeprazole)	C
Nasal Preparations	C
Nefazodone	
Hydrochloride	
(Nefazodone	
Hydrochloride)	C
Totolin	
(Guaifenesin,	
Phenylpropanolamine	
Hydrochloride)	C
Meclozine	
(Meclozine)	C
Fluticasone	
Propionate	
(Fluticasone	
Propionate)	C
Trimethobenzamide	
Hydrochloride	
(Trimethobenzamide	
Hydrochloride)	C
Respaire-Sr-120	
(Guaifenesin,	
Pseudophedrine	
Hydrochloride)	C
Risperidone	
(Risperidone)	C

Date:02/27/04ISR Number: 4308885-4Report Type:Expedited (15-DaCompany Report #2004011014
 Age:57 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300MG (TID), Other ORAL	Cardiac Disorder Loss Of Consciousness	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			All Other Therapeutic Products Moduretic "Msd" (Hdyrochlorothiazide , Amiloride Hydrochloride) Acetylsalicylic Acid	C		C

(Acetylsalicylic Acid)	C
Isosorbide Mononitrate (Isosorbide Mononitrate)	C
Clopidogrel Sulfate (Clopidogrel Sulfate)	C
Escitalopram (Escitalopram)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/27/04ISR Number: 4308924-0Report Type:Expedited (15-DaCompany Report #FR-JNJFOC-20040203901

Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cerebral Atrophy Drug Interaction Fall	Foreign Health Professional	Contramal (Tramadol Hydrochloride) Unspecified			
2 IN 1 DAY, ORAL		Headache			PS		ORAL
1 IN 1 DAY		Vertigo		Furosemide (Furosemide)	SS		
2 IN 1 DAY, ORAL				Neurontin (Gabapentin)	SS		ORAL
1 IN 1 DAY, ORAL				Irbesartan (Irbesartan)	SS		ORAL
ORAL				Effexor (Venlafaxine Hydrochloride)	SS		ORAL
				Difrarel (Difrarel) Dafalgan (Paracetamol) Vastarel (Trimetazidine Hydrochloride) Amlor (Amlodipine Besilate) Glucor (Acabose) Lansoprazole (Lansoprazole)	C C C C C C		

Date:03/01/04ISR Number: 4308270-5Report Type:Periodic

Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12413969

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Euphoric Mood		Glucophage Tabs 500			

Mg	PS	Bristol-Myers Squibb	ORAL
Neurontin	SS		
Furosemide	C		
Potassium	C		
Synthroid	C		
Albuterol	C		
Hydrocodone	C		
Mom	C		
Metamucil	C		
Colace	C		
Ibuprofen	C		
Acetaminophen	C		

Date:03/01/04ISR Number: 4309453-0Report Type:Expedited (15-DaCompany Report #2004007600
Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Albumin Decreased Dyspnoea	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
1200 MG (TID), ORAL		Medication Error	Professional				
150 MG (TID), ORAL		Oedema Peripheral Pitting Oedema Swelling Face		Diclofenac (Diclofenac)	SS		
				Mexiletine			

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800 MG (QID), ORAL	Hydrochloride (Mexiletine Hydrochloride)	SS	ORAL
	Fentanyl Citrate (Fentanyl Citrate)	C	
	Misoprostol (Misoprostol)	C	
	Quetiapine Fumarate (Quetiapine Fumarate)	C	

Date:03/01/04ISR Number: 4309595-XReport Type:Expedited (15-DaCompany Report #2004012403
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2400 MG (QID), ORAL	Drug Level Decreased	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

			Lamotrigine (Lamotrigine)	C		
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Date:03/01/04ISR Number: 4309760-1Report Type:Expedited (15-DaCompany Report #2004010782
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 1200 MG (QID), ORAL	Chromatopsia Pain	Health Professional	Neurontin (Gabapentin)	PS		ORAL

All Other Therapeutic Products	SS	
Hydromorphone (Hydromorphone)	C	
Potassium Chloride (Potassium Chloride)	C	
Insulin Glargine		

(Insulin Glargine)	C
Insulin Lispro	
(Insulin Lispro)	C
Atorvastatin	
(Atorvastatin)	C
Magnesium	
(Magnesium)	C
Omeprazole	
(Omeprazole)	C
Oxycodone	
Hydrochloride	
(Oxycodone	
Hydrochloride)	C
Pentoxifylline	
(Pentoxifylline)	C
Prednisone	
(Prednisone)	C
Torasemide	
(Torasemide)	C
Calcium Carbonate	
(Calcium Carbonate)	C
Paracetamol	
(Paracetamol)	C
Ascorbic Acid	

Freedom Of Information (FOI) Report

(Ascorbic Acid)	C
Tocopherol	
(Tocopherol)	C
Warfarin (Warfarin)	C
Metolazone	
(Metolazone)	C
Cetirizine	
Hydrochloride	
(Cetirizine	
Hydrochloride)	C
Allopurinol	
(Allopurinol)	C
Darbepoetin Alfa	
(Darbepoetin Alfa)	C
Diphenhydramine	
Hydrochloride	
(Diphenhydramine	
Hydrochloride)	C
Calcitrol	
(Ergocalciferol,	
Retinol, Calcium	
Carbonate)	C
Mycophenolate	
Mofetil	
(Mycophenolate	
Mofetil)	C
Clonazepam	
(Clonazepam)	C
Colchicine	
(Colchicine)	C
Losartan Potassium	
(Losartan Potassium)	C
Folic Acid (Folic	
Acid)	C
Furosemide	
(Furosemide)	C
Amitriptyline	
(Amitriptyline)	C

Date:03/01/04ISR Number: 4309761-3Report Type:Expedited (15-DaCompany Report #2004010846
 Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arterial Occlusive	Consumer	Neurontin			

900 MG (TID),
ORAL
Disease
Weight Increased
(Gabapentin)
PS
ORAL
Amlodipine Besilate
(Amlodipine
Besilate)
Vitamins
C
C

Date:03/02/04ISR Number: 4308786-1Report Type:Expedited (15-DaCompany Report #PHNU2004DE00766
Age: Gender:Male I/FU:F

Outcome PT
Disability Abnormal Behaviour
Other Alopecia
Anosmia
Bradyphrenia
Chills

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FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	Condition	Report Source	Product	Role	Manufacturer	Route
		Condition Aggravated Depression Difficulty In Walking					
400 mg, QOD		Disturbance In Attention Drug Interaction		Tegretal	PS	Novartis Sector: Pharma	ORAL
UNKNOWN		Dyskinesia Electroencephalogram		Ergenyl "Sanofi-Synthelabo"	SS		
UNKNOWN		Abnormal		Neurontin	SS		
60mg/day		Erectile Dysfunction		Gabitril	SS		ORAL
100mg/day	549 DAY	Fall		Topamax	SS		ORAL
400 mg, QID		Fatigue		Carbabeta	SS		ORAL
400 mg, TID		Gait Disturbance		Carbabeta	SS		ORAL
62.5 mg, TID		General Physical Health		Madopar	SS		ORAL
1mg/day		Deterioration Grand Mal Convulsion		Amantadin Parkotil	SS SS		ORAL ORAL
0.20mg/day		Liver Function Test		Parkotil	SS		ORAL
200 mg, QID		Abnormal		Comtess	SS		ORAL
1DF/day		Memory Impairment Mental Impairment		Levodopa Movergan	SS SS		ORAL ORAL
UNKNOWN	3DF/Day	Motor Dysfunction		Zentropil	C		
5mg/day		Movement Disorder		Selegiline	C		
15mg/day	366 DAY	Muscle Twitching		Remergil	C		ORAL
UNKNOWN	1000-1500mg/d	Myalgia Nerve Conduction Studies		Tasmar Keppra	C C		
ay		Abnormal					
UNKNOWN		Neuroleptic Malignant Syndrome		Nacom "Dupont Pharma"	C		
		Paralysis		Phenhydan	C		

UNKNOWN	500 mg, BID	Parkinson'S Disease	Keppra	C
		Salivary Hypersecretion	Lamictal	
		Simple Partial Seizures	"Glaxosmithkline"	C
UNKNOWN		Speech Disorder		
		Tremor		

Date:03/02/04ISR Number: 4309689-9Report Type:Expedited (15-DaCompany Report #2004010558
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Disorder Leukaemia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
200 MG (BID), Other ORAL		Lymphoma	Professional				
				Estradiol (Estradiol)	C		
				Escitalopram (Escitalopram)	C		

Date:03/02/04ISR Number: 4310984-8Report Type:Direct
 Age: Gender:Male I/FU:I Company Report #CTU 213552

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening PROVIDED IN Required PROVED IN		Back Pain		Effexor Xr	PS		
Intervention to Prevent Permanent Impairment/Damage		Drug Dependence		Zyprexa	SS		
		Infection		Hydrocodone	SS		
		Mental Disorder		Neurontin	SS		
		Pain		Ambien	SS		
				Oxycontin	SS		
				Oxycodone/Apap	C		
				Duragesic	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/04ISR Number: 4311056-9Report Type:Expedited (15-DaCompany Report #2004010841
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Anxiety Condition Aggravated	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG (600 MG TID), ORAL		Delayed Recovery From Anaesthesia Depressed Mood		Topiramate (Topiramate)	SS		ORAL
ORAL		Diplopia Drug Ineffective Dyspnoea		Drug Unspecified Escitalopram (Escitalopram)	SS SS		
10 MG (DAILY),		Epileptic Aura Grand Mal Convulsion Hot Flush Liver Function Test Abnormal Loss Of Consciousness Nerve Injury Vision Blurred Weight Increased		Citalopram Hydrobromide (Citalopram Hydrobromide) Levothyroxine Sodium (Levothyroxine Sodium) Phenobarbital (Phenobarbital) Alprazolam (Alprazolam) Clonazepam (Clonazepam)	SS C C C		

Date:03/02/04ISR Number: 4311061-2Report Type:Expedited (15-DaCompany Report #2004011279
Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Amnesia Difficulty In Walking	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Loss Of Consciousness Speech Disorder Visual Disturbance		Magnesium Oxide (Magnesium Oxide) Digoxin (Digoxin) Atorvastatin	C C		

(Atorvastatin) C
 Gemfibrozil C
 (Gemfibrozil) C
 Ramipril (Ramipril) C
 Vitamins C
 Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C
 Insulin (Insulin) C

Date:03/02/04ISR Number: 4311130-7Report Type:Expedited (15-DaCompany Report #2004011149
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG ORAL		Asthenia Drug Interaction	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other 2.5 MG ORAL		Endocarditis Medication Error	Professional	Lorazepam (Lorazepam)	SS		ORAL
0.5 MG ORAL		Oxygen Saturation Decreased		Alprazolam (Alprazolam)	SS		
ORAL		Somnolence		Clonazepam (Clonazepam)	SS		ORAL
				Lactulose			

FDA - Adverse Event Reporting System (AERS)

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(Lactulose)	SS
Buflomedil	
(Buflomedil)	C
Salbutamol	
(Salbutamol)	C
Insulin Human	
(Insulin Human)	C
Salmeterol	
(Salmeterol)	C
Beclometasone	
(Beclometasone)	C
Perindopril	
(Perindopril)	C

Date:03/02/04ISR Number: 4311132-0Report Type:Expedited (15-DaCompany Report #2004011016
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Balance Disorder	Foreign	Neurontin (Tablets)			
Initial or Prolonged	Coordination Abnormal	Health	(Gabapentin)	PS		ORAL
1800 MG						
(TID), ORAL	Dizziness	Professional				
	Fall	Company	Carbamazepine			
		Representative	(Carbamazepine)	C		

Date:03/03/04ISR Number: 4310486-9Report Type:Expedited (15-DaCompany Report #PHNU2004DE00766
 Age: Gender:Male I/FU:F

Outcome	PT
Disability	Abnormal Behaviour
Other	Alopecia
	Anosmia
	Aphasia
	Bradyphrenia
	Cerebellar Syndrome
	Chills
	Cogwheel Rigidity
	Condition Aggravated
	Depression
	Difficulty In Walking
	Disturbance In Attention
	Dysarthria

Dyskinesia
Electroencephalogram
Abnormal
Erectile Dysfunction
Fall
Fatigue
Gait Disturbance
General Physical Health
Deterioration
Grand Mal Convulsion
Hypokinesia
Liver Function Test
Abnormal
Malaise
Masked Facies
Memory Impairment
Mental Impairment
Micrographia
Mobility Decreased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Motor Dysfunction Movement Disorder Muscle Spasms				
400 mg, QOD		Health Professional	Tegretal	PS	Novartis Sector: Pharma	ORAL
UNKNOWN		Nerve Conduction Studies Abnormal	Ergenyl "Sanofi-Synthelabo"	SS		
UNKNOWN		Neuroleptic Malignant Syndrome	Neurontin	SS		
60mg/day			Gabitril	SS		ORAL
100mg/day	549 DAY	Paralysis	Topamax	SS		ORAL
400 mg, QID		Parkinson'S Disease	Carbabeta	SS		ORAL
400 mg, QID		Performance Status Decreased	Carbabeta	SS		ORAL
1DF/day			Movergan	SS		ORAL
		Posture Abnormal	Levodopa	SS		ORAL
.5 mg, QID		Regressive Behaviour	Parkotil	SS		ORAL
1mg/day		Salivary Hypersecretion	Parkotil	SS		ORAL
0.20mg/day		Simple Partial Seizures	Parkotil	SS		ORAL
200 mg, QID		Speech Disorder	Comtess	SS		ORAL
50 mg, QID		Tongue Paralysis	Amantadin	SS		ORAL
62.5 mg, TID		Tremor	Levodopa	SS		ORAL
400 mg, TID			Madopar	SS		ORAL
UNKNOWN	3DF/Day		Carbabeta	SS		ORAL
UNKNOWN	1000-1500mg/d		Zentropil	C		
ay			Keppra	C		
UNKNOWN			Nacom "Dupont Pharma"	C		

15mg/day	366 DAY	Tasmar	C	
		Remergil	C	ORAL
		L-Dopa	C	
5mg/day		Levetiracetam	C	
		Selegiline	C	
		Phenhydan	C	
UNKNOWN	500 mg, BID	Keppra	C	
		Lamictal		
UNKNOWN		"Glaxosmithkline"	C	

Date:03/03/04ISR Number: 4311898-XReport Type:Expedited (15-DaCompany Report #2003118605
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 800 MG (BID),		Generalised Oedema Hepatitis Hepatosplenomegaly	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
ORAL		Infectious Mononucleosis Mass Pyrexia Toxoplasmosis		Carbamazepine (Carbamazepine) All Other Therapeutic Products Ketoprofen (Ketoprofen)	C C C C		

Date:03/03/04ISR Number: 4312279-5Report Type:Expedited (15-DaCompany Report #2004002731
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Other	Abdominal Pain Upper Arthralgia Asthenia Chest Pain

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Complex Partial Seizures Fall Loss Of Consciousness	Report Source				
2800 MG (7 TIMES A DAY), ORAL		Overdose Pain Pain In Extremity Paraesthesia Skin Discolouration Tremor Vascular Occlusion Weight Decreased	Consumer	Dilantin Suspension (Phenytoin Sodium) Neurontin (Gabapentin)	PS SS		ORAL
				Lamotrigine (Lamotrigine) Atorvastatin (Atorvastatin) Benazepril Hydrochloride (Benazepril Hydrochloride) Escitalopram (Escitalopram) All Other Therapeutic Products Carbamazepine (Carbamazepine) Atenolol (Atenolol) Warfarin (Warfarin)	C C C C C C		

Date:03/03/04ISR Number: 4312325-9Report Type:Expedited (15-DaCompany Report #2004011148
Age: Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Body Temperature Decreased Hypotension Paralysis	Consumer	Neurontin (Gabapentin)	PS		

Date:03/03/04ISR Number: 4312334-XReport Type:Expedited (15-DaCompany Report #2004011471
Age:9 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Consumer	Neurontin (Tablets)			
		Brain Neoplasm Malignant		(Gabapentin)	PS		
		Condition Aggravated		Valproate Semisodium			
		Pharmaceutical Product		(Valproate			
		Complaint		Semisodium)	C		
		Simple Partial Seizures		Desloratadine			
				(Desloratadine)	C		

Date:03/03/04ISR Number: 4312337-5Report Type:Expedited (15-DaCompany Report #2004012493

Age:51 YR Gender:Male I/FU:I

Outcome	PT
Other	Bacterial Infection
	Gastric Infection
	Graft Infection
	Hernia
	Malaise
	Post Procedural
	Complication

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
			Potassium Chloride (Potassium Chloride)	C		
			Furosemide (Furosemide)	C		
			Methadone (Methadone)	C		
			Alprazolam (Alprazolam)	C		
			Isosorbide Mononitrate (Isosorbide Mononitrate)	C		
			Fenofibrate (Fenofibrate)	C		
			Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
			Colchicine (Colchicine)	C		
			Trandolapril (Trandolapril)	C		
			Atenolol (Atenolol)	C		
			Allopurinol (Allopurinol)	C		
			Oxygen (Oxygen)	C		

Date:03/03/04ISR Number: 4312400-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
Age:44 YR Gender:Female I/FU:F

Outcome
Hospitalization - PT
Initial or Prolonged Agitation
Disability Ankle Fracture
Other Bacteraemia
Cervicitis
Decreased Appetite
Depression
Drug Hypersensitivity
Dysgeusia

Dysphagia
Facial Bones Fracture
Gastrointestinal Disorder
Gastrointestinal
Infection
Hypersensitivity
Hypersomnia
Hypoglycaemia
Laboratory Test Abnormal
Macroglossia
Nausea
Oesophageal Disorder
Ovarian Infection
Overdose
Pain
Pitting Oedema
Road Traffic Accident

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Sialoadenitis Somatisation Disorder Speech Disorder	Report Source	Product	Role	Manufacturer	Route										
1800 MG		Swollen Tongue Traumatic Brain Injury	Consumer Health	Neurontin (Gabapentin)	PS		ORAL										
DAILY, ORAL		Upper Respiratory Tract	Professional	Risperidone (Risperidone) Lamotrigine (Lamotrigine) Lithium Carbonate (Lithium Carbonate) Clonazepam (Clonazepam) Methylphenidate Hydrochloride (Methylphenidate Hydrochloride) Levothyroxine Sodium (Levothyroxine Sodium) Liothyronine Sodium (Liothyronine Sodium) Sertraline Hydrochloride (Sertraline Hydrochloride) Pilocarpine Hydrochloride (Pilocarpine Hydrochloride) Metoprolol Succinate (Metoprolol Succinate) Lansoprazole (Lansoprazole) Hyoscyamine Sulfate (Hyoscyamine Sulfate) Diltiazem Hydrochloride (Diltiazem Hydrochloride)	SS	C	C	C	C	C	C	C	C	C	C	C	C

Bupropion Hydrochloride (Bupropion Hydrochloride)	C
Celebrex (Celecoxib)	C
Quetiapine Fumarate (Quetiapine Fumarate)	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
Olanzapine (Olanzapine)	C

Freedom Of Information (FOI) Report

Omeprazole	
(Omeprazole)	C
Nasal Preparations	C
Nefazodone	
Hydrochloride	
(Nefazodone	
Hydrochloride)	C
Totolin	
(Guaifenesin,	
Phenylpropanolamine	
Hydrochloride)	C
Meclozine	
(Meclozine)	C
Fluticasone	
Propionate	
(Fluticasone	
Propionate)	C
Trimethobenzamide	
Hydrochloride	
(Trimethobenzamide	
Hydrochloride)	C
Respaire-Sr-120	
(Guaifenesin,	
Pseudoephedrine	
Hydrochloride)	C

Date:03/03/04ISR Number: 4312593-3Report Type:Expedited (15-DaCompany Report #2004001696
 Age:75 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cataract	Consumer	Neurontin			
Other		Condition Aggravated	Health	(Gabapentin)	PS		ORAL
1200 MG		Hypoaesthesia	Professional				
(TID), ORAL		Pain		Telmisartan			
		Paraesthesia		(Telmisartan)	C		
		Post Procedural		Esomeprazole			
		Complication		(Esomeprazole)	C		
				Atorvastatin			
				(Atorvastatin)	C		
				Acetylsalicylic Acid			
				(Acetylsalicylic			
				Acid)	C		

Date:03/03/04ISR Number: 4312594-5Report Type:Expedited (15-DaCompany Report #2004012957

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Injury		Neurontin			
Other		Mental Disorder		(Gabapentin)	PS		
		Pain					
		Suicide Attempt					

Date:03/04/04ISR Number: 4313064-0Report Type:Expedited (15-DaCompany Report #2003127306

Age:84 YR Gender:Female I/FU:F

Outcome	PT
Other	Cardiac Failure
	Congestive

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Freedom Of Information (FOI) Report

Dose	Duration	Cholestasis Dizziness Hypocalcaemia	Report Source	Product	Role	Manufacturer	Route
300 MG (QID), ORAL		Hypotension International Normalised Ratio Decreased	Foreign Literature Health	Neurontin (Gabapentin)	PS		ORAL
35 MG (BID), 100 MG (DAILY), ORAL		Loss Of Consciousness Sleep Disorder Treatment Noncompliance Vertigo	Professional	Amitriptyline Hydrochloride (Amitriptyline Hydrochloride) Atenolol (Atenolol)	SS SS		ORAL
30 MG (DAILY), ORAL				Nifedipine (Nifedipine)	SS		ORAL
5 MG (DAILY), ORAL				Lisinopril (Lisinopril)	SS		ORAL
				Sodium Polystyrene Sulfonate (Sodium Polystyrene Sulfonate) Cyanocobalamin (Cyanocobalamin) Thiamine Hydrochloride (Thiamine Hydrochloride) Pyridoxine (Pyridoxine)	C C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Status Epilepticus	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Professional	Zoxan Ip (Doxazosin)	SS		ORAL
ORAL				Nitrendipine (Nitrendipine)	SS		ORAL
				Acetylsalicylate Lysine (Acetylsalicylate Lysine)	C		

Date:03/05/04ISR Number: 4312541-6Report Type:Expedited (15-DaCompany Report #PHNU2004DE00766
Age: Gender:Male I/FU:F

Outcome	PT
Disability	Abnormal Behaviour
Other	Alopecia
	Anosmia
	Aphasia
	Bradyphrenia
	Cerebellar Syndrome
	Chills
	Cogwheel Rigidity
	Complex Partial Seizures
	Condition Aggravated

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Depression Difficulty In Walking Disturbance In Attention	Report Source	Product	Role	Manufacturer	Route
400 mg, QOD		Drug Interaction Dysarthria	Health Professional	Tegretal	PS	Novartis Sector: Pharma	ORAL
UNKNOWN		Dyskinesia Electroencephalogram		Ergenyl "Sanofi-Synthelabo"	SS		
UNKNOWN		Abnormal		Neurontin	SS		
60mg/day		Epilepsy		Gabitril	SS		ORAL
100mg/day	549 DAY	Erectile Dysfunction		Topamax	SS		ORAL
400 mg, QID		Fall		Carbabeta	SS		ORAL
400 mg, QID		Fatigue		Carbabeta	SS		ORAL
1DF/day		Gait Disturbance		Movergan	SS		ORAL
.5 mg, QID		General Physical Health Deterioration		Levodopa Parkotil	SS SS		ORAL ORAL
1mg/day		Grand Mal Convulsion		Parkotil	SS		ORAL
0.20mg/day		Hypokinesia		Parkotil	SS		ORAL
200 mg, QID		Liver Function Test		Comtess	SS		ORAL
50 mg, QID		Abnormal Malaise		Amantadin Levodopa	SS SS		ORAL ORAL
62.5 mg, TID		Masked Facies		Madopar	SS		ORAL
400 mg, TID		Memory Impairment		Carbabeta	SS		ORAL
UNKNOWN	3DF/Day	Mental Impairment		Zentropil	C		
UNKNOWN	1000-1500mg/d	Micrographia		Keppra	C		
ay		Mobility Decreased					
UNKNOWN		Motor Dysfunction Movement Disorder		Nacom "Dupont Pharma"	C		

15mg/day	366 DAY	Muscle Spasms	Tasmar	C	
		Muscle Twitching	Remergil	C	ORAL
		Myalgia	L-Dopa	C	
		Nerve Conduction Studies	Levetiracetam	C	
		Abnormal	Selegiline	C	
5mg/day		Neuroleptic Malignant	Phenhydan	C	
		Syndrome	Keppra	C	
UNKNOWN	500 mg, BID	Nuclear Magnetic	Lamictal		
		Resonance Imaging Brain	"Glaxosmithkline"	C	
UNKNOWN		Abnormal			
		Paralysis			
		Parkinson'S Disease			
		Performance Status			
		Decreased			
		Posture Abnormal			
		Regressive Behaviour			
		Salivary Hypersecretion			
		Simple Partial Seizures			
		Speech Disorder			
		Tongue Paralysis			
		Tremor			

Date:03/05/04ISR Number: 4313649-1Report Type:Expedited (15-DaCompany Report #2004012533
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Condition Aggravated	Consumer	Dilantin Suspension			
Initial or Prolonged		Convulsion		(Phenytoin Sodium)	PS		
Other		Hot Flush		Neurontin			
2400 MG		Muscle Twitching		(Gabapentin)	SS		
(TID),		Tremor		Phenobarbital			
				(Phenobarbital)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Valproate Semisodium
 (Valproate
 Semisodium) C
 Clonazepam
 (Clonazepam) C

Date:03/05/04ISR Number: 4314015-5Report Type:Expedited (15-DaCompany Report #2004012903
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	900 MG (TID), ORAL	Cardiac Disorder	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Lisinopril
 (Lisinopril) C
 Alfuzosin
 (Alfuzosin) C
 Zopiclone
 (Zopiclone) C

Date:03/08/04ISR Number: 4314351-2Report Type:Expedited (15-DaCompany Report #2004002253
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:03/08/04ISR Number: 4314359-7Report Type:Expedited (15-DaCompany Report #2004012237
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 50 MG, ORAL		Drug Ineffective Facial Palsy	Consumer	Sinequan (Doxepin) Navane (Capsules)	PS		ORAL

ORAL	Insomnia Tinnitus	(Tiotixene) (Thiothixene)	SS	ORAL
300 MG		Neurontin (Gabapentin) (Gabapentin)	SS	ORAL
(DAILY), ORAL		Rofecoxib (Rofexcoxib)	SS	

Date:03/08/04ISR Number: 4314423-2Report Type:Expedited (15-DaCompany Report #2004009355
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Back Disorder
Other	Back Pain
	Blood Cholesterol
	Increased
	Chest Pain
	Condition Aggravated
	Erectile Dysfunction
	Facial 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
		Feeling Cold Gastrooesophageal Reflux Disease	Consumer				
1800 MG		Glossodynia Hypersomnia	Consumer	Neurontin (Gabapentn)	PS		ORAL
(TID), ORAL		Medication Error					
20 MG		Movement Disorder Neck Pain		Lipitor (Atorvsatatin)	SS		ORAL
(DAILY), ORAL		Oesophageal Spasm					
		Oral Mucosal Disorder		Lithium (Lithium)	SS		
		Pain In Jaw		Naproxen (Naproxen)	SS		
		Sleep Disorder		Rofecoxib			
		Tooth Abscess		(Rofecoxib)	SS		
1000 MG		Tremor		Amoxicillin (Amoxicillin)	SS		ORAL
(BID), ORAL							
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Vitamins	C		
				Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C		

Date:03/08/04ISR Number: 4314519-5Report Type:Expedited (15-DaCompany Report #2004010782
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Level Decreased Pain	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 MG QID, ORAL		Procedural Complication					
		Xanthopsia		Hydromorphone (Hydromorphone)	C		
				Potassium Chloride			

(Potassium Chloride)	C
Insulin Glargne	
(Insulin Glargine)	C
Insulin Lispro	
(Insulin Lispro)	C
Atorvastatin	
(Atorvastatin)	C
Magnesium	
(Magnesium)	C
Omeprazole	
(Omeprazole)	C
Oxycodone	
Hydrochloride	
(Oxycodone	
Hydrochloride)	C
Pentoxifylline	
(Pentoxifylline)	C
Prednisone	
(Prednisone)	C
Torasemide	
(Torasemide)	C
Calcium Carbonate	
(Calcium Carbonate)	C
Paracetamol	
(Paracetamol)	C

Freedom Of Information (FOI) Report

Ascorbic Acid	
(Ascorbic Acid)	C
Tocopherol	
(Tocopherol)	C
Warfarin (Warfarin)	C
Metolazone	
(Metolazone)	C
Cetirizine	
Hydrochloride	
(Cetirizine	
Hydrochloride)	C
Allopurinol	
(Allopurinol)	C
Darbepoetin Alfa	
(Darbepoetin Alfa)	C
Diphenhydramine	
Hydrochloride	
(Diphenhydramine	
Hydrochloride)	C
Calcitrol	
(Ergocalciferol,	
Retinol, Calcium	
Carbonate)	C
Mycophenolate	
Mofetil	
(Mycophenolate	
Mofetil)	C
Clonazepam	
(Clonazepam)	C
Colchicine	
(Colchicine)	C
Losartan Potassium	
(Losartan Potassium)	C
Folic Acid (Folic	
Acid)	C
Furosemide	
(Furosemide)	C
Amitriptyline	
(Amitriptyline)	C

Date:03/09/04ISR Number: 4314573-0Report Type:Expedited (15-DaCompany Report #DSA_23975_2004
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization - 2.5 MG QD PO	Depressed Level Of Consciousness	Foreign Health	Temesta Alprazolam	PS SS	ORAL ORAL
Initial or Prolonged 0.25 MG BID	Drug Interaction	Professional			
PO	Medication Error	Other	Neurontin	SS	ORAL
800 MG TID PO	Oxygen Saturation Decreased		Neurontin Rivotril	SS SS	ORAL
5 GTT QD PO	Somnolence		Fonzylane Coversyl Duphalac Insulin Mixtard Serevent Becotide Ventoline	C C C C C C C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/09/04ISR Number: 4315698-6Report Type:Expedited (15-DaCompany Report #US-SHR-04-021606
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase Increased Aspartate	Health Professional	Betaseron (Interferon Beta-1b) Injection, 250ug	PS		
SUBCUTANEOUS	8 MIU,	EVERY Aminotransferase					
2D,		Increased					
SUBCUTANEOUS		Blood Bilirubin Increased		Zanaflex (Tizanidine Hydrochloride)	SS		ORAL
ORAL				Neurontin /Unk/(Gabapentin)	SS		ORAL
ORAL				Potassium (Potassium)	SS		ORAL

Date:03/10/04ISR Number: 4314918-1Report Type:Direct Company Report #CTU 214144
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea		Neurontin 100 Mg	PS		ORAL
1 PO TID							

Date:03/10/04ISR Number: 4315473-2Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
Age:44 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Agitation
Initial or Prolonged	Ankle Fracture
Disability	Arthralgia
Other	Bacteraemia
	Biopsy Tongue Abnormal
	Brain Damage
	Cervicitis
	Decreased Appetite

Depression
Drug Hypersensitivity
Drug Level Increased
Dysgeusia
Dyskinesia
Dysphagia
Facial Bones Fracture
Gastrointestinal Disorder
Gastrointestinal
Infection
Hypersomnia
Hypoglycaemia
Imprisonment
Joint Swelling
Macroglossia
Memory Impairment
Nausea
Oesophageal Disorder
Ovarian Infection
Pitting Oedema
Post Procedural
Complication
Road Traffic Accident
Sialoadenitis
Somatisation Disorder
Speech Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Swollen Tongue Traumatic Brain Injury Upper Respiratory Tract	Report Source	Product	Role	Manufacturer	Route
1800 MG		Infection Vomiting	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL		Weight Increased	Professional	Lamotrigine (Lamotrigine)	C		
				Lithium Carbonate (Lithium Carbonate)	C		
				Clonazepam (Clonazepam)	C		
				Methylphenidate Hydrochloride (Methylphenidate Hydrochloride)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Liothyronine Sodium (Liothyronine Sodium)	C		
				Sertraline Hydrochloride (Sertraline Hydrochloride)	C		
				Pilocarpine Hydrochloride (Pilocarpine Hydrochloride)	C		
				Metoprolol Succinate (Metoprolol Succinate)	C		
				Lansoprazole (Lansoprazole)	C		
				Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C		
				Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C		
				Bupropion Hydrochloride			

(Bupropion Hydrochloride)	C
Celebrex (Celecoxib)	C
Quetiapine Fumarate (Quetiapine Fumarate)	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C
Olanzapine	

Freedom Of Information (FOI) Report

(Olanzapine)	C
Omeprazole	
(Omeprazole)	C
Nasal Preparations	C
Nefazodone	
Hydrochloride	
(Nefazodone	
Hydrochloride)	C
Totolin	
(Guaifenesin,	
Phenylpropanolamine	
Hydrochloride)	C
Meclozine	
(Meclozine)	C
Fluticasone	
Propionate	
(Fluticasone	
Propionate)	C
Trimethobenzamide	
Hydrochloride	
(Trimethobenzamide	
Hydrochloride)	C
Respaire-Sr-120	
(Guaifenesin,	
Pseudoephedrine	
Hydrochloride)	C
Risperidone	
(Risperidone)	C

Date:03/10/04ISR Number: 4315676-7Report Type:Expedited (15-DaCompany Report #KII-2001-0004421
 Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma Completed Suicide Depression Multi-Organ Failure	Health Professional Company Representative	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet			PS
160 MG, BID		Multiple Drug Overdose Stupor		Oxyir (Oxycodone Hydrochloride, Oxycodone Hydrochloride)			SS
5 MG				Neurontin			

600 MG, AM;	(Gabapentin)	SS	
1200 MG, PM			
150 MG, HS	Desyrel (Trazodone Hydrochloride)	SS	
10 MG, SEE	Sonata (Zaleplon)	SS	ORAL
TEXT, ORAL			
1 TABLET,	Soma (Carisoprodol)	SS	ORAL
QID, ORAL			

Date:03/10/04ISR Number: 4315722-0Report Type:Expedited (15-DaCompany Report #2004013492
Age: Gender:Male I/FU:I

Outcome PT
Other Bipolar Disorder
Brain Damage
Closed Head Injury
Depressed Level Of
Consciousness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Dependence Drug Withdrawal Syndrome Fear	Report Source	Product	Role	Manufacturer	Route
		Feeling Abnormal Hallucination, Visual Pain Psychotic Disorder	Consumer	Neurontin (Gabapentin)	PS		

Date:03/10/04ISR Number: 4315725-6Report Type:Expedited (15-DaCompany Report #2004013496
Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Other		Abnormal Behaviour Anger Feeling Abnormal Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:03/10/04ISR Number: 4315735-9Report Type:Expedited (15-DaCompany Report #2004013502
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Other		Drug Ineffective Gastric Operation Intestinal Operation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:03/10/04ISR Number: 4315999-1Report Type:Expedited (15-DaCompany Report #2004012916
Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Other		Agranulocytosis Aplastic Anaemia Chills Cytomegalovirus Infection Erythropoiesis Abnormal	Foreign Health Professional	Neurontin (Gabapentin) Flucloxacillin Sodium	PS		ORAL
1.8 G DAILY ORAL							

6 GRAM DAILY	Fungal Infection	(Flucloxacillin		
	Granulocytosis	Sodium)	SS	ORAL
ORAL	Pancytopenia			
	Pseudomonas Infection	Cefadroxil		
ORAL	Pyrexia	(Cefadroxil)	SS	ORAL
		Insulin Lispro	C	
		Insulin Glargine	C	

Date:03/10/04ISR Number: 4316119-XReport Type:Expedited (15-DaCompany Report #2003018320
Age:76 YR Gender:Male I/FU:F

Outcome	PT
Death	Abnormal Behaviour
Hospitalization -	Anxiety
Initial or Prolonged	Arteriosclerosis
Other	Atrioventricular Block
	Second Degree
	Bone Marrow Depression
	Bone Marrow Disorder
	C-Reactive Protein
	Increased
	Cardiac Enzymes Increased

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Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
900 MG		Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
(DAILY), ORAL		Professional				
INTRAVENOUS	600 MG		Vfend Injection (Voriconazole)	SS		
(DAILY),						
INTRAVENOUS			Zoloft (Sertraline Hydrochloride) (Sertraline)	SS		
			Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		
			Vancomycin Hydrochloride (Vancymycin Hydrochloride)	C		
			Fluconazole (Fluconazole)	C		
			Meropenem (Meropenem)	C		
			Levofloxacin (Levofloxacin)	C		
			Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
			Clindamycin Hydrochloride (Clindamycin Hydrochloride)	C		

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Asthenia Cardiac Failure	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other ORAL			Chlamydial Infection	Professional				
ORAL			Diastolic Dysfunction Headache		Dihydroergotoxine (Dihydroergotoxine)	SS		ORAL
ORAL			Hypertrophic Cardiomyopathy		Alendronate Sodium (Alendronate Sodium)	SS		ORAL
			Inflammation Pulmonary Embolism Pulmonary Fibrosis					

Date:03/11/04ISR Number: 4316396-5Report Type:Expedited (15-DaCompany Report #2003116310
Age:59 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Asthenia Catatonia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
300 MG (THREE TIMES A DAY),ORAL			Dysarthria	Professional				
			Somnolence	Company				
			Tremor	Representative	Panadeine Co			

Freedom Of Information (FOI) Report

(Codeine Phosphate,
Paracetamol) C
Triazolam
(Triazolam) C
Loperamide
Hydrochloride
(Loperamide
Hydrochloride) C

Date:03/12/04ISR Number: 4316154-1Report Type:Expedited (15-DaCompany Report #PHBS2003CH11427
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 37440MIN	Coagulopathy Diabetic Neuropathy		Tegretol	PS	Novartis Sector: Pharma	ORAL
Initial or Prolonged UNKNOWN	Eosinophilia 27360MIN		Neurontin	SS		
UNKNOWN	Erythema		Huminsulin Basal	C		
UNKNOWN	Exanthem		Insulin Lispro	C		
UNKNOWN	Haematoma		Dafalgan	C		
	Hepatic Enzyme Increased Hepatocellular Damage Hypersensitivity Hyponatraemia Leukocytosis Lymphocyte Transformation Test Positive Pain Petechiae Polycythaemia Pyrexia Rash Maculo-Papular Shift To The Left Sinus Tachycardia Skin Oedema Skin Test Positive					

Date:03/12/04ISR Number: 4317326-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 400 MG (BID), Disability ORAL Other	Acrochordon Cardiac Pacemaker Insertion Convulsion Dermatitis Contact Diarrhoea Gallbladder Disorder Hypoaesthesia Intervertebral Disc Displacement Joint Injury Lower Gastrointestinal Haemorrhage Peptic Ulcer Protein Total Decreased Syncope Viral Pharyngitis	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin)	PS SS C C C C		ORAL

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Date:03/12/04ISR Number: 4317396-1Report Type:Expedited (15-DaCompany Report #2004015759
Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Asthenia Multiple Sclerosis Relapse Tremor	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:03/12/04ISR Number: 4317510-8Report Type:Expedited (15-DaCompany Report #2004001757
Age:14 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1500 MG Other (TID), ORAL (BOLUS)	Alanine Aminotransferase Increased Aspartate Aminotransferase Increased Electroencephalogram Abnormal Electromyogram Abnormal Labyrinthitis Liver Function Test Abnormal Nervous System Disorder Respiratory Arrest Viral Infection	Health Professional Company Representative	Neurontin (Gabapentin) Corticosteroid Nos (Corticosteroid Nos) Panadeine Co (Codeine Phosphate, Paracetamol) Amitriptyline Hydrochloride (Aqmitriptyline Hydrochloride) Meclozine Hydrochloride (Meclozine Hydrochloride) Oxaprozin (Oxaprozin)	PS SS C C C		ORAL

Date:03/12/04ISR Number: 4317551-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200295
Age:74 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death	Arrhythmia	Health	Neurontin			

Hospitalization - 2400 MG (600 Initial or Prolonged MG, QID) Other	Cardiac Arrest Cardio-Respiratory Arrest Coordination Abnormal Deep Vein Thrombosis Fall	Professional Company Representative	(Gabapentin)	PS
400 MG (100 MG, Q6H)	Peripheral Embolism		Fosphenytoin Sodium (Fosphenytoin Sodium)	SS
2000 MG (500 MG, QID)	Post Procedural Complication Post Procedural Haematoma		Metronidazole (Metronidazole)	SS
800 MG (200 MG, QID)	Pulmonary Embolism Pyrexia Subdural Haematoma		Carbamazepine (Carbamazepine)	SS
			Clavulin (Clavulanate Potassium, Amoxicillin Trihydrate)	SS

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Freedom Of Information (FOI) Report

Date:03/12/04ISR Number: 4317573-XReport Type:Expedited (15-DaCompany Report #2004014131

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3600 MG		Dizziness Loss Of Consciousness	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other (TID), ORAL		Myokymia Speech Disorder Syncope	Professional				

Date:03/12/04ISR Number: 4317574-1Report Type:Expedited (15-DaCompany Report #2003121920

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Disability 900 MG (TID), ORAL		Cellulitis Completed Suicide Deafness	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Leukoerythroblastic Anaemia Liver Disorder		Amitriptyline (Amitriptyline) Fentanyl (Fentanyl) Etoricoxib (Etoricoxib) Prednisone (Prednisone) Diazepam (Diazepam) Morphine Hydrochloride (Morphine Hydrochloride)	C C C C C		

Date:03/12/04ISR Number: 4317797-1Report Type:Expedited (15-DaCompany Report #2004014109

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Condition Aggravated	Consumer	Neurontin			

1800M MG	Headache	(Gabapentin)	PS	ORAL
(TID), ORAL	Myalgia			
	Pneumonia	Ranitidine		
	Sluggishness	Hydrochloride		
	Stomach Discomfort	(Ranitidine		
	Weight Decreased	Hydrochloride)	C	
		Clonidine		
		(Clonidine)	C	
		Phenobarbital		
		(Phenobarbital)	C	
		Atenolol (Atenolol)	C	
		Prednisone		
		(Prednisone)	C	
		Tolterodine		
		L-Tartrate		
		(Tolterodine		
		L-Tartrate)	C	
		Paroxetine		
		Hydrochloride		
		(Paroxetine		
		Hydrochloride)	C	

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Freedom Of Information (FOI) Report

Date:03/13/04ISR Number: 4317216-5Report Type:Expedited (15-DaCompany Report #2004001696
Age:75 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cataract	Consumer	Neurontin			
Other		Hypoaesthesia	Health	(Gabapentin)	PS		ORAL
1200 MG		Pain	Professional				
(TID), ORAL		Paraesthesia		Telmisartan			
		Post Procedural		(Telmisartan)	C		
		Complication		Esomeprazole			
		Spinal Operation		(Esomeprazole)	C		
				Atorvastatin			
				(Atorvastatin)	C		
				Acetylsalicylic Acid			
				(Acetylsalicylic			
				Acid)	C		

Date:03/15/04ISR Number: 4318267-7Report Type:Expedited (15-DaCompany Report #2004010848
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blood Creatinine	Health	Gabapentin			
Hospitalization -		Increased	Professional	(Gabapentin)	PS		ORAL
Initial or Prolonged		Condition Aggravated					
600M MG (200		Encephalopathy		Oxycocet			
Other		Mental Status Changes		(Paracetamol,Oxycodo			
MG TID),ORAL		Myoclonus		ne Hydrochloride)	SS		ORAL
(TWO TABLETS		Oedema Peripheral					
Q4H), ORAL		Somnolence		Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	SS		ORAL
60 MG (BID),							
ORAL				Bosentan (Bosentan)	SS		ORAL
62.5 MG							

(BID), ORAL

Warfarin Sodium	
(Warfarin Sodium)	C
Furosemide	
(Furosemide)	C
Prednisone	
(Prednisone)	C
Epoprostenol Sodium	
(Epoprostenol Sodium)	C
Pantoprazole	
(Pantoprazole)	C
Metformin	
Hydrochloride	
(Metformin Hydrochloride)	C
Bactrim	
(Sulfamethoxazole, Trimethoprim)	C
Oxygen (Oxygen)	C
Lisinopril	
(Lisinopril)	C
Metolazone	
(Metolazone)	C

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Date:03/15/04ISR Number: 4318273-2Report Type:Expedited (15-DaCompany Report #2004012493

Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bacterial Infection	Consumer	Neurontin			
Other		Condition Aggravated		(Gabapentin)	PS		ORAL
ORAL		Gastric Infection		Potassium Chloride			
		Hernia		(Potassium Chloride)	C		
		Hernia Hiatus Repair		Furosemide			
		Implant Site Infection		(Furosemide)	C		
		Malaise		Methadone			
		Neuropathy		(Methadone)	C		
		Postoperative Infection		Alprazolam			
		Vomiting		(Alprazolam)	C		
				Isosorbide			
				Mononitrate			
				(Isosorbide			
				Mononirate)	C		
				Fenofibrate			
				(Fenofibrate)	C		
				Acetylsalicylicacid			
				(Acetylsalicylic			
				Acid)	C		
				Colchicine			
				(Colchicine)	C		
				Trandolapril			
				(Trandolapril)	C		
				Atenolol (Atenolol)	C		
				Allopurinol			
				(Allopurinol)	C		
				Oxygen (Oxygen)	C		

Date:03/15/04ISR Number: 4318649-3Report Type:Expedited (15-DaCompany Report #2004012403

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Laboratory Test	Foreign	Neurontin (Tablets)			
		Interference	Health	(Gabapentin)	PS		ORAL
2400 MG			Professional				
(QID), ORAL				Lamotrigine			

Date:03/15/04ISR Number: 4318680-8Report Type:Expedited (15-DaCompany Report #2004014815

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Dissociation	Foreign	Neurontin			
Initial or Prolonged	Loss Of Consciousness	Health	(Gabapentin)	PS		ORAL
1800 MG		Professional				
(DAILY), ORAL		Company	All Other			
		Representative	Therapeutic Products	SS		

Date:03/15/04ISR Number: 4318681-XReport Type:Expedited (15-DaCompany Report #2003122796

Age: Gender:Female I/FU:I

Outcome	PT	Report Source
Other	Breast Cancer Metastatic	Foreign
	Hepatic Enzyme Increased	Health
	Vomiting	Professional

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Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
900 MG (TID)			Neurontin (Gabapentin)	PS		

Date:03/15/04ISR Number: 4318908-4Report Type:Expedited (15-DaCompany Report #2004014973
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Zoloft (Sertraline)	PS		
Disability		Coma		Celecoxib			
Other		Gangrene		(Celecoxib)	SS		
		Illiteracy		Neurontin			
		Nervous System Disorder		(Gabapentin)	SS		
		Post Procedural		Lipitor			
		Complication		(Atorvastatin)	SS		
		Reading Disorder		Zyrtec (Tablets)			
		Wheelchair User		(Cetirizine)	SS		
				Glucotrol			
				(Glipizide)	SS		

Date:03/16/04ISR Number: 4317950-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0313585A
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Lamictal	PS	Glaxosmithkline	ORAL
Other		Visual Acuity Reduced		Neurontin	SS		ORAL
300MG per day							
800MG Twice		Visual Field Defect		Sectral	C		ORAL
				Mogadon	C		ORAL
per day							

Date:03/16/04ISR Number: 4320047-3Report Type:Expedited (15-DaCompany Report #USA-2004-0013630
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression Anxiety Thinking Abnormal	Consumer Other	Oxycontin Tablets(Oxycodone Hydrochloride) Cr Tablet Xanax (Alprazolam) Neurontin (Gabapentin)	PS SS SS		

Date:03/17/04ISR Number: 4320147-8Report Type:Expedited (15-DaCompany Report #2003118605
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 800 MG BID, ORAL		Generalised Oedema Hepatitis Hepatosplenomegaly Infectious Mononucleosis Pyrexia Swelling Toxoplasmosis	Foreign Consumer	Gabapentin (Gabapentin) Carbamazepine (Carbamazepine) Ketoprofen (Ketoprofen)	PS SS C		ORAL

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Date:03/17/04ISR Number: 4320148-XReport Type:Expedited (15-DaCompany Report #2004015519

Age:54 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG DAILY, ORAL	Abnormal Behaviour Delirium Hallucination, Visual	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Becozym (Vitamin B Nos)	C
Thiamine Hydrochloride (Thiamine Hydrochloride)	C
Chloral Hydrate (Chloral Hydrate)	C
Pipamperone (Pipamperone)	C

Date:03/17/04ISR Number: 4320206-XReport Type:Expedited (15-DaCompany Report #2004015759

Age:53 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID) ORAL	Multiple Sclerosis Relapse	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Morphine Sulfate (Morphine Sulfate)	C
Paracetamol (Paracetamol)	C
Celiprolol (Celiprolol)	C
Spiroinolactone (Spiroinolactone)	C
Chondroitin Sulfate Sodium (Chondroitin Sulfate Sodium)	C

Date:03/17/04ISR Number: 4320210-1Report Type:Expedited (15-DaCompany Report #2004007396
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Condition Aggravated	Foreign	Neurontin			
Other		Convulsion	Health	(Gabapentin)	PS		ORAL
400 MG							
(DAILY), ORAL		Drug Ineffective	Professional				
		Pharmaceutical Product		Gabapentin			
UNKNOWN	UNKNOWN	Complaint		(Gabapentin)	SS		
		Pneumonia Aspiration					
		Porphyria					

Date:03/17/04ISR Number: 4320271-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200657
Age:44 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Ankle Fracture
Initial or Prolonged	Anxiety
Disability	Aphonia
Other	Asthma
	Atrial Flutter
	Atrial Tachycardia

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Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1800 MG	(DAILY), ORAL	Atrioventricular Block				
		Blood Urea Increased				
		Brain Damage				
		Chest Pain	Neurontin			
		Condition Aggravated	(Gabapentin)	PS		ORAL
		Convulsion				
		Depression	Lamotrigine			
		Diarrhoea	(Lamotrigine)	C		
		Drug Hypersensitivity	Lithium Carbonate			
		Dyskinesia	(Lithium Carbonate)	C		
		Dyspnoea	Clonazepam			
		Ecchymosis	(Clonazepam)	C		
		Facial Bones Fracture	Methylphenidate			
		Increased Upper Airway	Hydrochloride			
		Secretion	(Methylphenidate			
		Intentional Misuse	Hydrochloride)	C		
		Irritable Bowel Syndrome	Levothyroxine Sodium			
		Mean Cell Haemoglobin	(Levothyroxine			
		Increased	Sodium)	C		
		Medication Error	Liothyronine Sodium			
		Nausea	(Liothyronine			
		Nervous System Disorder	Sodium)	C		
		Pharyngitis	Sertraline			
		Propionibacterium	Hydrochloride			
		Infection	(Sertraline			
		Pruritus	Hydrochloride)	C		
		Psoriasis	Pilocarpine			
		Pyrexia	Hydrochloride			
		Red Blood Cell Count	(Pilocarpine			
		Decreased	Hydrochloride)	C		
		Rhinitis Allergic	Metoprolol Succinate			
		Road Traffic Accident	(Metoprolol			
		Skin Ulcer	Succinate)	C		
		Speech Disorder	Lansoprazole			
		Stress	(Lansoprazole)	C		
		Supraventricular	Hyoscyamine Sulfate			
		Tachycardia	(Hyoscyamine			
		Swollen Tongue	Sulfate)	C		
		Syncope	Diltiazem			
		Thermal Burn	Hydrochloride			
		Viral Sinusitis	(Diltiazem			
		White Blood Cell Count	Hydrochloride)	C		
		Decreased	Bupropion			
		Wrist Fracture	Hydrochloride			

(Bupropion Hydrochloride)	C
Celebrex (Celecoxib)	C
Quetiapine Fumarate (Quetiapine Fumarate)	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C
Olanzapine	

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(Olanzapine)	C
Omeprazole	
(Omeprazole)	C
Nasal Preparations	C
Nefazodone	
Hydrochloride	
(Nefazodone	
Hydrochloride)	C
Totolin	
(Guaifenesin,	
Phenylpropanolamine	
Hydrochloride)	C
Meclozine	
(Meclozine)	C
Fluticasone	
Propionate	
(Fluticasone	
Propionate)	C
Trimethobenzamide	
Hydrochloride	
(Trimethobenzamide	
Hydrochloride)	C
Respaire-Sr-120	
(Guaifenesin,	
Pseudoephedrine	
Hydrochloride)	C
Risperidone	
(Risperidone)	C

Date:03/17/04ISR Number: 4320428-8Report Type:Expedited (15-DaCompany Report #2004015518
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bladder Disorder	Consumer	Neurontin			
Other		Chronic Obstructive		(Gabapentin)	PS		ORAL
600 MG		Pulmonary Disease					
(DAILY), ORAL		Dyspepsia		Diuretics	C		
		Emphysema		All Other			
		Gastrointestinal Disorder		Therapeutic Products	C		
		Lung Neoplasm Malignant					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Bradyarrhythmia Electroencephalogram Abnormal Fall Headache	Health Professional Other	Effexor Ip (Venlafaxine Hydrochloride, Capsule, Extended Release, 0)	PS		ORAL
37.5 MG 2X PER 1 DAY; ORAL	Metabolic Disorder Nervous System Disorder					
300 MG 1X PER 1 DAY; ORAL	Vertigo		Aprovel (Irbesartan, , 0)	SS		ORAL
2 TABLET 1X PER 1 DAY; ORAL			Contramal (Tramadol Hydrochloride, , 0)	SS		ORAL
			Lasilix (Furosemide,			

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1 DOSE 1X PER		, 0)	SS	ORAL
1 DAY; ORAL				
2 DOSE 1X PER		Neurontin (Gabapentin, , 0)	SS	ORAL
1 DAY; ORAL				
		Amlor (Amlodipine Besilate)	C	
		Glucor (Acarbose)	C	
		Ogast (Lansoprazole)	C	
		Difrarel (Betacarotene/Myrtil lus)	C	
		Dafalgan (Paracetamol)	C	
		Vastarel (Trimetazidine Hydrochloride)	C	

Date:03/19/04ISR Number: 4322535-2Report Type:Expedited (15-DaCompany Report #2004016481
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Acute Psychosis Aggression	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Confusional State Drug Withdrawal Syndrome	Professional	All Other Therapeutic Products	C		

Date:03/19/04ISR Number: 4322536-4Report Type:Expedited (15-DaCompany Report #2004016094
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (DAILY), ORAL		Retinal Detachment	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Bradycardia		Lasilix Faible	PS	Aventis	
Initial or Prolonged	Dizziness				Pharmaceuticals Inc.	ORAL
	Fall		Neurontin	SS		ORAL
Dose unit:						
units	Headache					
	Metabolic Disorder		Aprovel	SS		ORAL
Dose unit:	Nervous System Disorder		Contramal	SS		ORAL
units						
			Effexor	SS		ORAL
			Amlor	C		ORAL
			Glucor	C		ORAL
			Ogast	C		ORAL
			Difrarel	C		ORAL
			Dafalgan	C		ORAL
			Vastarel 20 Mg	C		ORAL

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Date:03/22/04ISR Number: 4323031-9Report Type:Expedited (15-DaCompany Report #001-0945-M0200657

Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG Disability (DAILY), ORAL Other	Ankle Fracture Anxiety Arrhythmia Supraventricular Asthma Atrial Flutter Blood Urea Decreased Chest Pain Convulsion Deafness Depression Diarrhoea Disease Recurrence Dissociation Drug Hypersensitivity Dyskinesia Dyspnoea Ecchymosis Facial Bones Fracture Fatigue Hand Fracture Hypothyroidism Increased Upper Airway Secretion Irritable Bowel Syndrome Loss Of Consciousness Mean Cell Haemoglobin Increased Multiple Fractures Nausea Nervous System Disorder Pharyngitis Propionibacterium Infection Pruritus Psoriasis Red Blood Cell Count Decreased Rhinitis Allergic Road Traffic Accident	Consumer Health Professional	Neurontin (Gabapentin) Lamotrigine (Lamotrigine) Lithium Carbonate (Lithium Carbonate) Clonazepam Methylphenidate Hydrochloride (Methylphenidate Hydrochloride) Levothyroxine Sodum (Levothyroxine Sodium) Liothyronine Sodium (Liothyronine Sodium) Sertraline Hydrochloride (Sertraline Hydrochloride) Pilocarpine Hydrochloride (Pilocaprine Hydrochloride) Metoprolol Succinate (Metoprolol Succinate) Lansoprazole (Lansoprazole) Hyoscyamine Suflate (Hyoscyamine Sulfate) Diltiazem Hydrochloride (Diltiazem Hydrochloride) Bupropion Hydrochloride	PS C C C C C C C C C C C C C C C C		ORAL

Self-Medication	(Bupropion	
Skin Ulcer	Hydrochloride)	C
Speech Disorder	Celebrex (Celecoxib)	C
Stress	Quetiapine Fumarate	
Swollen Tongue	(Quetiapine	
Syncope	Fumarate)	C
Thermal Burn	Metoprolol Tartrate	
Tinnitus	(Metoprolol	
Viral Sinusitis	Tartrate)	C
White Blood Cell Count	Citalopram	
Decreased	Hydrobromide	
Wrist Fracture	(Citalopram	
	Hydrobromide)	C
	Hyoscyamine Sulfate	
	(Hyoscyamine	
	Sulfate)	C
	Olanzapine	
	(Olanzapine)	C

Freedom Of Information (FOI) Report

Omeprazole
 (Omeprazole) C
 Nasal Preparations C
 Nefazodone
 Hydrochloride
 (Nefazodone
 Hydrochloride) C
 Totolin
 (Guaifenesin,
 Phenylpropanolamine
 Hydrochloride) C
 Meclozine (Meclozine) C
 Fluticasone
 Propionate
 (Fluticasone
 Propionate) C
 Trimethobenzamide
 Hydrochloride
 (Trimethobenzamide
 Hydrochloride) C
 Respaire-Sr-120
 (Guaifenesin,
 Pseudoephedrine
 Hydrochloride) C
 Risperidone
 (Risperidone) C

Date:03/22/04ISR Number: 4323082-4Report Type:Expedited (15-DaCompany Report #2004016474
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour	Consumer	Neurontin			
Other		Blood Cholesterol		(Gabapentin)	PS		ORAL
1800 MG		Increased					
(TID); ORAL		Cerebrovascular Accident		Lipitor			
		Chest Pain		(Atorvastatin)	SS		
UNKNOWN	UNKNOWN	Confusional State					
(UNKNOWN),		Depression					
UNKNOWN		Dysarthria		Clopidogrel Sulfate	C		

Insomnia
Speech Disorder
Tremor
Weight Decreased

Date:03/22/04ISR Number: 4323086-1Report Type:Expedited (15-DaCompany Report #2004016191

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged UNKNOWN Other (UNKNOWN); ORAL	Crying Diabetes Mellitus Fatigue Hepatic Cirrhosis Hepatic Steatosis Hepatomegaly Impaired Work Ability Intestinal Obstruction Liver Disorder Tremor	Health Professional	Neurontin (Tablets) (Gabapentin) Diltiazem Hydrochloride Benazepril Hydrochloride Hydrochlorothiazide	PS C C C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/22/04ISR Number: 4323184-2Report Type:Expedited (15-DaCompany Report #2003113622
 Age:19 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 1200 MG, TID, Initial or Prolonged ORAL Disability ORAL	Coagulopathy Demyelination Dermatitis Exfoliative Diabetic Neuropathy Eosinophilia Exanthem Generalised Oedema Haematoma Hepatocellular Damage Hypersensitivity Hyponatraemia International Normalised Ratio Increased Leukocytosis Lymphocyte Transformation Test Positive Petechiae Platelet Count Increased Pyrexia Rash Maculo-Papular Shift To The Left Sinus Tachycardia Skin Test Positive Swelling	Foreign Health Professional	Neurontin (Gabapentin) Carbamazepine Insulin Human Insulin Lispro Paracetamol	PS SS C C C		ORAL ORAL

Date:03/22/04ISR Number: 4323221-5Report Type:Expedited (15-DaCompany Report #2004002729
 Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (SIX TIMES A DAY), ORAL	Hunger Hypoglycaemia Unawareness Paraesthesia Tremor	Foreign Consumer Company Representative	Neurontin (Tablets) (Gabapentin)	PS		ORAL

Lansoprazole
 (Lansoprazole) C
 Losartan Potassium
 (Losartan Potassium) C

Date:03/23/04ISR Number: 4324048-0Report Type:Expedited (15-DaCompany Report #2004005466

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Asthenia	Consumer	Dilantin Kapseals (Phenytoin Sodium)	PS		ORAL
ORAL Other		Chest Discomfort Chest Pain		Neurontin (Gabapentin)	SS		ORAL
1200 MG BID		Convulsion					
ORAL		Dementia Fatigue		Phenytoin (Phenytoin)	SS		
300 MG DAILY		Insomnia Loss Of Consciousness		Phenobarbital (Phenobarbital)	SS		
30 MG DAILY		Paraesthesia Vitamin B Complex Deficiency					

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Freedom Of Information (FOI) Report

Date:03/23/04ISR Number: 4324591-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 400 MG (BID), Disability ORAL Other	Acrochordon Angina Unstable Back Pain Bladder Constriction Bone Disorder Brain Neoplasm Cardiac Arrest Cognitive Disorder Conversion Disorder Convulsion Coronary Artery Disease Disease Recurrence Dizziness Dysphemia Electrocardiogram Q Waves Electrocardiogram Qrs Complex Shortened Excoriation Exposure To Toxic Agent Fatigue Gallbladder Disorder Hypoaesthesia Intervertebral Disc Displacement Lower Gastrointestinal Haemorrhage Lymphocyte Count Decreased Memory Impairment Mobility Decreased Nervous System Disorder Peptic Ulcer Protein Total Decreased Pruritus Red Blood Cell Sedimentation Rate Increased Scoliosis Skin Lesion Somnolence	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin)	PS SS C C C C		ORAL

Speech Disorder
Syncope
Tremor

Date:03/23/04ISR Number: 4324855-4Report Type:Expedited (15-DaCompany Report #2003121737

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other 300 MG (DAILY), ORAL	Carotid Artery Disease Epilepsy Haematocrit Decreased Haemoglobin Decreased Red Blood Cell Count Decreased Somnolence	Foreign Health Professional Company Representative	Neurontin (Tablets)(Gabapentin) Clomipramine (Clomipramine) Oxybutynin (Oxybutynin) Baclofen (Baclofen)	 PS C C C		 ORAL

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Freedom Of Information (FOI) Report

Acetylsalicylate
 Lysine
 (Acetylsalicylate
 Lysine) C
 Lormetazepam
 (Lormetazepam) C
 Nulytely (Sodium
 Bicarbonate,
 Potassium Chloride,
 Sodium Chloride,
 Macrogol) C
 Risedronate Sodium
 (Risedronate Sodium) C
 Tramadol C

Date:03/23/04ISR Number: 4324858-XReport Type:Expedited (15-DaCompany Report #2004006431
 Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Foreign	Neurontin			
Other		Cardiomegaly	Health	(Gabapentin)	PS		ORAL
200 MG (BID),		Headache	Professional				
ORAL		Musculoskeletal Stiffness	Company				
		Myalgia	Representative				
		Neck Pain					
		Pyrexia					
		Red Blood Cell					
		Sedimentation Rate					
		Increased					
		Temporal Arteritis					

Date:03/23/04ISR Number: 4325892-6Report Type:Expedited (15-DaCompany Report #2012272
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Consumer	Oxycontin			
Hospitalization -		Asthenia	Health	Tablets(Oxycodon			
Initial or Prolonged		Back Pain	Professional	Hydrochloride) Cr			

Other
10 & 20 & 40

Drug Abuser

Other

Tablet

PS

Drug Toxicity

MG

Headache

Iron Deficiency Anaemia

Mallory-Weiss Syndrome

Neck Pain

Respiratory Arrest

Tachycardia

Diazepam(Diazepam) SS

Oxazepam(Oxazepam) SS

Temazepam(Temazepam) SS

Lorazepam(Lorazepam) SS

Cannabnoids(Cannabis
) SS

Diphenhydramine
Hydrochloride(Diphen
hydramine

Hydrochloride) SS

Gabapentin(Gabapenti
n) SS

Claritin(Loratadine) C

Zoloft (Sertraline
Hydrochloride) C

Altace(Ramipril) C

Allopurinol
(Allopurinol) C

Depakote (Valproate

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Freedom Of Information (FOI) Report

Semisodium) C
 Soma (Carisoprodol) C
 Relafen (Nabumetone) C

Date:03/24/04ISR Number: 4325533-8Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
 Age:44 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Abdominal Pain	Consumer	Neurontin			
Initial or Prolonged	Ankle Fracture	Health	(Gabapentin)	PS		ORAL
1800 MG						
Disability	Anxiety	Professional				
(DAILY), ORAL						
Other	Aphonia		Lamotrigine			
	Asthma		(Lamotrigine)	C		
	Atrial Flutter		Lithium Carbonate			
	Atrial Tachycardia		(Lithium Carbonate)	C		
	Atrioventricular Block		Clonazepam			
	Blood Urea Increased		(Clonazepam)	C		
	Brain Damage		Methylphenidate			
	Chest Pain		Hydrochloride			
	Convulsion		(Methylphenidate			
	Deafness		Hydrochloride)	C		
	Depression		Levothyroxine Sodium			
	Diarrhoea		(Levothyroxine			
	Drug Hypersensitivity		Sodium)	C		
	Dyspnoea		Liothyronine Sodium			
	Ecchymosis		(Liothyronine			
	Facial Bones Fracture		Sodium)	C		
	Fatigue		Sertraline			
	Hand Fracture		Hydrochloride			
	Increased Upper Airway		(Sertraline			
	Secretion		Hydrochloride)	C		
	Irritable Bowel Syndrome		Pilocarpine			
	Loss Of Consciousness		Hydrochloride			
	Mean Cell Haemoglobin		(Pilocarpine			
	Mental Disorder		Hydrochloride)	C		
	Nausea		Metoprolol Succinate			
	Oedema Peripheral		(Metoprolol			
	Overdose		Succinate)	C		
	Pharyngitis		Lansoprazole			
	Propionibacterium		(Lansoprazole)	C		
	Infection		Hyoscyamine Sulfate			
	Pruritus		(Hyoscyamine			
	Psoriasis		Sulfate)	C		

Red Blood Cell Count	Diltiazem	
Decreased	Hydrochloride	
Rhinitis Allergic	(Diltiazem	
Road Traffic Accident	Hydrochloride)	C
Skin Ulcer	Bupropion	
Speech Disorder	Hydrochloride	
Stress	(Bupropion	
Swollen Tongue	Hydrochloride)	C
Syncope	Celebrex (Celecoxib)	C
Thermal Burn	Quetiapine Fumarate	
Tinnitus	(Quetiapine	
Viral Sinusitis	Fumarate)	C
White Blood Cell Count	Metoprolol Tartrate	
Decreased	(Metoprolol	
Wrist Fracture	Tartrate)	C
	Citalopram	
	Hydrobromide	
	(Citalopram	

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Freedom Of Information (FOI) Report

Hydrobromide)	C
Hyoscyamine Sulfate	
(Hyoscyamine Sulfate)	C
Olanzapine	
(Olanzapine)	C
Omeprazole	
(Omeprazole)	C
Nasal Preparations	C
Nefazodone	
Hydrochloride	
(Nefazodone Hydrochloride)	C
Totolin	
(Guaifenesin, Phenylpropanolamine Hydrochloride)	C
Meclozine	
(Meclozine)	C
Fluticasone Propionate	
(Fluticasone Propionate)	C
Trimethobenzamide Hydrochloride	
(Trimethobenzamide Hydrochloride)	C
Respaire-Sr-120	
(Guaifenesin, Pseudoephedrine Hydrochloride)	C
Risperidone	
(Risperidone)	C

Date:03/24/04ISR Number: 4325941-5Report Type:Expedited (15-DaCompany Report #2004015518

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bladder Disorder	Consumer	Neurontin			
Other		Chronic Obstructive		(Gabapentin)	PS		ORAL
600 MG		Pulmonary Disease					
(DAILY), ORAL		Dyspepsia		Diuretics	C		
		Emphysema		All Other			

Gastric Disorder
Lung Neoplasm Malignant
Pain
Pruritus

Therapeutic Products C

Date:03/24/04ISR Number: 4325945-2Report Type:Expedited (15-DaCompany Report #2004009355

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Back Disorder
Other	Back Pain
	Blood Cholesterol
	Increased
	Cardiac Disorder
	Chest Pain
	Drug Ineffective

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Intolerance Drug Level Increased Erectile Dysfunction	Report Source				
1800 MG		Facial Pain Feeling Cold	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
(TID), ORAL		Gastrointestinal Disorder	Professional				
20 MG		Gastrooesophageal Reflux Disease		Lipitor (Atorvastatin)	SS		ORAL
(DAILY), ORAL		Glossodynia					
		Hypersomnia		Lithium (Lithium)			
		Neck Pain		(Lithium)	SS		
		Oesophageal Spasm		Naproxen (Naproxen)	SS		
		Pain In Jaw		Rofecoxib			
		Paralysis		(Rofecoxib)	SS		
1000 MG		Post Procedural Pain		Amoxicillin			
(BID), ORAL		Sleep Disorder		(Amoxicillin)	SS		ORAL
		Tooth Abscess					
		Treatment Noncompliance		All Other			
		Tremor		Therapeutic Products	C		
				Levothyroxine Sodium			
				(Levothyroxine			
				Sodium)	C		
				Vitamins	C		
				Diltiazem			
				Hydrochloride			
				(Diltiazem			
				Hydrochloride)	C		

Date:03/24/04ISR Number: 4326143-9Report Type:Expedited (15-DaCompany Report #2004018059

Age: Gender:Male I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
800 MG		Hepatitis Cholestatic Hepatitis Toxic	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL			Professional				

Morphine Sulfate
 (Morphine Sulfate) C
 All Other
 Therapeutic Products C
 Piretanide Sodium
 (Piretanide Sodium) C

Date:03/25/04ISR Number: 4323789-9Report Type:Direct
 Age:56 YR Gender:Female I/FU:I

Company Report #CTU 215238

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG , ONE Required TID		Obstructive Airways Disorder		Neurontin 300 Mg	PS		
Intervention to Prevent Permanent Impairment/Damage		Swollen Tongue		Doxepin Levoxyl Celexa Morphine Ir/Morphine Msir Fosamax	C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/25/04ISR Number: 4324590-2Report Type:Direct
Age:35 YR Gender:Male I/FU:I

Company Report #CTU 215242

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 600 MG TID PO Initial or Prolonged	Pancreatitis		Gabapentin 600 Mg	PS		ORAL

Date:03/25/04ISR Number: 4327530-5Report Type:Expedited (15-DaCompany Report #FRWYE593019FEB04
Age:73 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Bradyarrhythmia Cerebral Atrophy Eczema Electroencephalogram Abnormal	Health Professional Other	Effexor Lp (Venlafaxine Hydrochloride, Capsule, Extended Release, 0)	PS		ORAL
37.5 MG 2X PER 1 DAY, ORAL	Fall Headache					
300MG 1X PER 1 DAY, ORAL	Metabolic Disorder Nervous System Disorder Pruritus		Aprovel (Irbesartan, 0)	SS		ORAL
2 TABLET 1X PER 1 DAY, ORAL	Self-Medication Toxic Skin Eruption Urticaria Vertigo		Contramal (Tramadol Hydrochloride, 0)	SS		ORAL
100 MG 3X PER 1 DAY, ORAL			Glucor (Acarbose, 0)	SS		ORAL
1 DOSE 1X PER 1 DAY, ORAL			Lasilix (Furosemide, 0)	SS		ORAL

2 DOSE 1X PER

1 DAY, ORAL

Neurontin (Gabapentin, 0) SS ORAL

Amlor (Amlodipine Besilate) C
Ogast (Lansoprazole) C
Difrarel (Betacarotene/Myrtilus) C
Dafalgan (Paracetamol) C
Vastarel (Trimetazidine Hydrochloride) C

Date:03/25/04ISR Number: 4327983-2Report Type:Expedited (15-DaCompany Report #2004017640

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 3200 MG (QID) Other , ORAL	Diffuse Alveolar Damage Lung Injury Respiratory Failure	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Oxycodone Hydrochloride(Oxycodone Hydrochloride)	C		

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FDA - Adverse Event Reporting System (AERS)

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Date:03/25/04ISR Number: 4327984-4Report Type:Expedited (15-DaCompany Report #2004017478

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Deformity	Consumer	Neurontin			
Other		Hypoaesthesia		(Gabapentin)	PS		ORAL
600 MG							
(DAILY), ORAL				All Other			
				Therapeutic Product	C		
				Potassium			
				(Potassium)	C		
				Diazepam (Diazepam)	C		
				Alprazolam			
				(Alprazolam)	C		
				Vicodin			
				(Paracetamol,			
				Hydrocodone			
				Bitartrate)	C		
				Methadone			
				(Methadone)	C		

Date:03/25/04ISR Number: 4328112-1Report Type:Expedited (15-DaCompany Report #HQWYE540423JAN04

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Intentional Misuse	Health	Effexor Xr			
		Overdose	Professional	(Venlafaxine			
		Sinus Tachycardia		Hydrochloride,			
		Sudden Onset Of Sleep		Capsule, Extended			
		Tonic Clonic Movements		Release)	PS		ORAL
600 MG THREE							
		Tonic Convulsion					
TO FOUR TIMES							
DAILY, ORAL							
				Neurontin			
				(Gabapentin,)	SS		
400 MG DOSE,							
"UP TO 10,000							

Date:03/29/04ISR Number: 4325658-7Report Type:Expedited (15-DaCompany Report #PHNU2004DE00766
Age: Gender:Male I/FU:F

Outcome	PT
Disability	Abnormal Behaviour
Other	Alopecia
	Anosmia
	Aphasia
	Bradyphrenia
	Cerebellar Syndrome
	Chills
	Cogwheel Rigidity
	Condition Aggravated
	Depression
	Difficulty In Walking
	Disturbance In Attention
	Dysarthria
	Dyskinesia
	Electroencephalogram
	Abnormal
	Erectile Dysfunction
	Fall
	Fatigue

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Gait Disturbance General Physical Health Deterioration				
400 mg, QOD		Grand Mal Convulsion Hypokinesia	Tegretal	PS	Novartis Sector: Pharma	ORAL
UNKNOWN		Liver Function Test Abnormal	Ergenyl "Sanofi-Synthelabo"	SS		
UNKNOWN		Malaise	Neurontin	SS		
60mg/day		Masked Facies	Gabitril	SS		ORAL
100mg/day	549 DAY	Memory Impairment	Topamax	SS		ORAL
400 mg, QID		Mental Impairment	Carbabeta	SS		ORAL
400 mg, QID		Micrographia	Carbabeta	SS		ORAL
1DF/day		Mobility Decreased	Movergan	SS		ORAL
		Motor Dysfunction	Levodopa	SS		ORAL
.5 mg, QID		Movement Disorder	Parkotil	SS		ORAL
1mg/day		Muscle Spasms	Parkotil	SS		ORAL
0.20mg/day		Muscle Twitching	Parkotil	SS		ORAL
200 mg, QID		Myalgia	Comtess	SS		ORAL
		Nerve Conduction Studies Abnormal	Amantadin Levodopa	SS SS		ORAL ORAL
50 mg, QID		Neuroleptic Malignant Syndrome	Madopar	SS		ORAL
62.5 mg, TID			Carbabeta	SS		ORAL
400 mg, TID						
UNKNOWN	3DF/Day	Nuclear Magnetic Resonance Imaging Brain	Zentropil	C		
UNKNOWN	1000-1500mg/d	Abnormal	Keppra	C		
ay						
UNKNOWN		Paralysis Parkinson'S Disease	Nacom "Dupont Pharma"	C		

		Performance Status	Tasmar	C	
		Decreased	Remergil	C	ORAL
15mg/day	366 DAY	Posture Abnormal	L-Dopa	C	
		Regressive Behaviour	Levetiracetam	C	
		Salivary Hypersecretion	Selegiline	C	
5mg/day		Simple Partial Seizures	Phenhydan	C	
		Speech Disorder	Keppra	C	
UNKNOWN	500 mg, BID	Tongue Paralysis	Lamictal		
		Tremor	"Glaxosmithkline"	C	
UNKNOWN					

Date:03/29/04ISR Number: 4330273-5Report Type:Expedited (15-DaCompany Report #2004018058
Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (TID),		Chronic Fatigue Syndrome Syncope	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Digoxin (Digoxin)	C		
				Furosemide (Furosemide)	C		
				Omeprazole (Omeprazole)	C		

Date:03/29/04ISR Number: 4330341-8Report Type:Expedited (15-DaCompany Report #200410158BFR
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 75 MG, TOTAL Initial or Prolonged DAILY, ORAL		Eczema	Foreign Health	Glucor (Acarbose)	PS		ORAL
ONCE, ORAL		Pruritus	Professional	Furosemide	SS		ORAL
BID, ORAL		Toxic Skin Eruption Urticaria	Other	Neurontin (Gabapentin)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

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ONCE, ORAL	Aprovel (Irbesartan)	SS	ORAL
BID, ORAL	Contramal (Tramadol Hydrochloride)	SS	ORAL
75 MG, TOTAL	Effexor (Venlafaxine Hydrochloride)	SS	ORAL
DAILY, ORAL	Ogast	C	
	Difrarel	C	
	Dafalgan	C	
	Amlor	C	
	Vastarel	C	

Date:03/29/04ISR Number: 4330345-5Report Type:Expedited (15-DaCompany Report #2004018059
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 800 MG DAILY, ORAL		General Physical Health Deterioration Hepatitis Cholestatic	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Piretanide Sodium (Piretanide Sodium)	SS		
				Morphine Sulfate (Morphine Sulfate)	C		
				Distigmine Bromide (Distigmine Bromide)	C		

Date:03/29/04ISR Number: 4330349-2Report Type:Expedited (15-DaCompany Report #2004018700
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG DAILY, ORAL		Abasia Tremor	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL

Amitriptyline(Amitriptyline) C
 Bromazepam (Bromazepam) C
 Methadone (Methadone) C

Date:03/29/04ISR Number: 4330350-9Report Type:Expedited (15-DaCompany Report #2004014437
 Age:68 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening	Asthma	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Health Professional	Cortisone (Cortisone)	C		

Date:03/29/04ISR Number: 4330351-0Report Type:Expedited (15-DaCompany Report #2004015525
 Age:77 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Malaise Palpitations	Foreign Health Professional Company

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
		Neurontin (Gabapentin)	PS		
		Piretanide (Piretanide)	C		
		Metoprolol Tartrate (Metoprolol Tartrate)	C		
		Glyceryl Trinitrate (Glyceryl Trinitrate)	C		
		Molsidomine (Molsidomine)	C		
		Clopidogrel Sulfate (Clopidogrel Sulfate)	C		

Date:03/29/04ISR Number: 4330529-6Report Type:Expedited (15-DaCompany Report #2004017254
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other 600 MG		Spinal Fracture	Consumer	Neurontin (Gabapentin)	PS		

Date:03/29/04ISR Number: 4331204-4Report Type:Expedited (15-DaCompany Report #2004007885
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG (DAILY), ORAL		Complex Partial Seizures Metrorrhagia Pain In Extremity Vaginal Haemorrhage	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Levetiracetam (Levetiracetam)	C		
				Fluoxetine (Fluoxetine)	C		

Date:03/29/04ISR Number: 4331227-5Report Type:Expedited (15-DaCompany Report #2004018051
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pancreas Transplant	Health	Dilantin Suspension			
		Renal Transplant	Professional	(Phenytoin Sodium)	PS		
		Treatment Noncompliance		Neurontin			
		Unintended Pregnancy		(Gabapentin)	SS		

Date:03/29/04ISR Number: 4331406-7Report Type:Expedited (15-DaCompany Report #2004018049
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bedridden	Consumer	Neurontin			
		Blindness		(Gabapentin)	PS		ORAL
ORAL		Drug Interaction		Aetylslicylic Acid			
		Neuralgia		(Acetylsalicylic			
		Sexual Assault Victim		Acid)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Oxazepam C
 Clonazepam
 (Clonazepam) C
 Analgesics C
 Levothyroxine Sodium
 (Levothyroxine
 Sodium) C

Date:03/29/04ISR Number: 4331409-2Report Type:Expedited (15-DaCompany Report #2004017642
 Age:71 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2400 MG Other (QID), ORAL	Catheter Related Infection Cystitis Dry Mouth Spinal Fusion Surgery Urinary Hesitation	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL

Date:03/29/04ISR Number: 4331428-6Report Type:Expedited (15-DaCompany Report #KII-2004-0008228
 Age:69 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other ORAL	Acidosis Blood Potassium Decreased Blood Pressure Diastolic Decreased	Study Health Professional Other	Morphine Sulfate (Similar To Nda 19-516) (Morphine Sulfate) Unknown	PS		ORAL
ORAL	Blood Pressure Systolic Increased		Neurontin (Gabapentin)	SS		ORAL
ORAL	Blood Urea Increased Coma		Nortriptyline (Nortriptyline)	SS		ORAL
ORAL	Drug Screen Positive Electrocardiogram Abnormal Haematocrit Decreased		Acetaminophen W/Oxycodone (Oxycodone Hydrochloride,			

ORAL	Haemoglobin Decreased	Paracetamol)	SS	ORAL
ORAL	Lethargy Nervous System Disorder	Amitriptyline (Amitriptyline)	SS	ORAL
ORAL	Pneumonia Aspiration Po2 Decreased	Benzodiazepine Derivatives ()	SS	ORAL
ORAL	Pulmonary Oedema Sedation	Beta Blocking Agents ()	SS	ORAL
ORAL	Sinus Tachycardia	Digoxin (Digoxin)	SS	ORAL
ORAL		Antiepileptics () Tricyclic Antidepressants ()	SS SS	ORAL
ORAL		Tetrahydrocannabinol (Tetrahydrocannabinol)	SS	

Date:03/29/04ISR Number: 4332244-1Report Type:Expedited (15-DaCompany Report #2004007885
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Vaginal Haemorrhage	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
1200 MG, (TID), ORAL			Professional	Levetiracetam			

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Freedom Of Information (FOI) Report

(Levetiracetam) C
 Fluoxetine
 (Fluoxetine) C
 Phenobarbital
 (Phenobarbital) C

Date:03/29/04ISR Number: 4332252-0Report Type:Expedited (15-DaCompany Report #2004019174
 Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Neurontin			
		Hip Fracture		(Gabapentin)	PS		
1600 MG		Neuralgia					
(DAILY)		Post Procedural		Neurontin			
		Complication		(Gabapentin)	SS		
1600 MG							
(DAILY)							
				Calcitonin, Salmon			
				(Calcitonin, Salmon)	C		
				Alendronate Sodium			
				(Alendronate Sodium)	C		
				Risedronate Sodium			
				(Risedronate Sodium)	C		

Date:03/29/04ISR Number: 4332833-4Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040301432
 Age:22 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Agonal Death Struggle	Health	Duragesic (Fentanyl)			
		Cardiac Failure	Professional	Patch	PS		
TRANSDERMAL	75 UG, 2 IN	Cardio-Respiratory Arrest					
72 HOUR,		Cardiomegaly					
TRANSDERMAL		Drug Toxicity		Diazepam (Diazepam)	SS		
		Hepatic Congestion		Oxycodone			
		Mucous Membrane Disorder		(Oxycodone)	SS		
		Nervous System Disorder		Morphine (Morphine)	SS		

Obesity
 Oesophagitis
 Pharmaceutical Product
 Complaint
 Pulmonary Congestion
 Pulmonary Oedema
 Respiratory Failure
 Scoliosis

Gabapentin
 (Gabapentin) SS

Date:03/29/04ISR Number: 4335207-5Report Type:Periodic Company Report #03P-163-0243402-00
 Age:48 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Arthralgia Hypotrichosis Photopsia Visual Disturbance	Consumer	Humira 40 Mg/ 0.8 Ml Pre-Filled Syringe (Humira) (Adalimumab) (Adalimumab)			
SUBCUTANEOUS	40 MG, 1 IN 2				PS		
WK,							
SUBCUTANEOUS				Gabapentin	SS		ORAL
PER ORAL				Prednisone	C		
				Esomeprazole	C		
				Oxycodone			

Freedom Of Information (FOI) Report

Hydrochloride C
Clonazepam C

Date:03/31/04ISR Number: 4330896-3Report Type:Expedited (15-DaCompany Report #2004019098
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Discomfort Abdominal Distension	Consumer	Lipitor (Atorvastatin)	PS		ORAL
80 MG		Accident At Work					
(DAILY), ORAL		Ankle Fracture Anorexia		Neurontin (Tablets) (Gabapentin)	SS		ORAL
2400 MG		Back Pain					
(TID), ORAL		Complex Regional Pain Syndrome		Topiramate (Topiramate)	C		
		Crying		Ezetimibe			
		Depression		(Ezetimibe)	C		
		Diarrhoea					
		Drug Ineffective					
		Fall					
		Fatigue					
		Feeling Abnormal					
		Hyperhidrosis					
		Insomnia					
		Intervertebral Disc Protrusion					
		Irritability					
		Malaise					
		Muscle Spasms					
		Road Traffic Accident					
		Treatment Noncompliance					

Date:03/31/04ISR Number: 4331005-7Report Type:Expedited (15-DaCompany Report #2004014973
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Coma	Consumer	Zoloft (Sertraline)	PS		
Other		Dysgraphia		Celecoxib			

Gangrene	(Celecoxib)	SS
Mental Impairment	Neurontin	
Post Procedural	(Gabapentin)	SS
Complication	Lipitor	
Reading Disorder	(Atorvastatin)	SS
Umbilical Hernia	Zyrtec (Tablets)	
	(Cetirizine)	SS
	Glucotrol	
	(Glipizide)	SS
	Valdecoxib	
	(Valdecoxib)	C

Date:03/31/04ISR Number: 4331009-4Report Type:Expedited (15-DaCompany Report #2004018498

Age:44 YR Gender:Female I/FU:I

Outcome	PT
Other	Affective Disorder
	Arthralgia
	Disease Recurrence
	Insomnia
	Tongue Neoplasm Malignant

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Freedom Of Information (FOI) Report

Stage Unspecified

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
200 MG	(DAILY), ORAL	Health Professional	Zoloft (Sertraline)	PS		ORAL
1200 MG	(DAILY), ORAL		Neurontin (Tablets) (Gabapentin)	SS		ORAL
			Antiflammatory/Antir heumatic Products	C		

Date:03/31/04ISR Number: 4331016-1Report Type:Expedited (15-DaCompany Report #2003037220
Age:74 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Diplopia Ivth Nerve Paresis	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
900 MG TID, ORAL		Strabismus	Professional	Levothyroxine (Levothyroxine)	C		
				Ezetimibe (Ezetimibe)	C		
				Cefalexin (Cefalexin)	C		
				Etodolac (Etodolac)	C		
				Pethidine Hydrochloride (Pethidine Hydrochloride)	C		

Date:03/31/04ISR Number: 4331018-5Report Type:Expedited (15-DaCompany Report #2004019965
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -	Abnormal Behaviour	Health	Neurontin	
Initial or Prolonged	Arthritis Bacterial	Professional	(Gabapentin)	PS
Other	Drug Clearance Decreased	Company		
	Renal Failure Acute	Representative		

Date:03/31/04ISR Number: 4331023-9Report Type:Expedited (15-DaCompany Report #2004005100
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Extraocular Muscle Paresis	Health Professional	Neurontin (Gabapentin)	PS		
1500 MG (BID)				Acetylsalicylic Acid	C		
				Diltiazem			
				Hydrochloride	C		
				Folic Acid	C		
				Atrovastatin	C		
				Budesonide	C		
				Salbutamol	C		
				Cetirizine			
				Hydrochloride	C		
				Mometasone Furoate	C		
				Cyclobenzaprine			
				Hydrochloride	C		
				Vicodin			

Freedom Of Information (FOI) Report

(Paracetamol,
Hydrocodone
Bitartrate) C
Nortriptyline
Hydrochloride C
Metaxalone C
Ranitidine
Hydrochloride C
Oxycocet
(Paracetamol,
Oxycodone
Hydrochloride) C

Date:03/31/04ISR Number: 4331113-0Report Type:Expedited (15-DaCompany Report #2004019177
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hospitalisation	Consumer	Neurontin (Gabapentin)	PS		

Date:03/31/04ISR Number: 4331170-1Report Type:Expedited (15-DaCompany Report #2004021304
Age: Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hepatotoxicity	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:03/31/04ISR Number: 4331174-9Report Type:Expedited (15-DaCompany Report #2004021300
Age: Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hepatotoxicity	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Bradyarrhythmia Cerebral Atrophy	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other ORAL	Electroencephalogram Abnormal	Professional	Furosemide (Furosemide)	SS		ORAL
ORAL	Fall Headache		Irbesartan (Irbesartan)	SS		ORAL
ORAL	Mental Disorder Metabolic Disorder Toxic Skin Eruption Urticaria		Tramadol Hydrochloride (Tramadol Hydrochloride)	SS		ORAL
75 MG, ORAL	Vertigo		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS		ORAL
300 MG, ORAL			Acarbose (Acarbose)	SS		ORAL
			Amlodipine Besilate (Amlodipine Besilate)	C		

Freedom Of Information (FOI) Report

Lansoprazole (Lansoprazole)	C
Diffrarel (Betacarotene, Myrtillus)	C
Paracetamol (Paracetamol)	C
Trimetazidine Hydrochloride (Trimetazidine Hydrochloride)	C
Betacarotene (Betacarotene)	C

Date:03/31/04ISR Number: 4332193-9Report Type:Expedited (15-DaCompany Report #2004018637
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Condition Aggravated Difficulty In Walking	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL		Polyneuropathy	Professional				

Date:03/31/04ISR Number: 4332217-9Report Type:Expedited (15-DaCompany Report #2002052306
Age:83 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (THREE TIMES DAILY), ORAL		Arrhythmia Asthenia Cardiac Failure Drug Toxicity Dyspnoea Generalised Oedema	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
				Amitriptyline (Amitriptyline)	C		
				Ginkgo Biloba (Ginkgo Biloba)	C		
				Omeprazole (Omeprazole)	C		

Verapamil (Verapamil)	C
Sprionolactone (Spironolactone)	C
Methadone Hydrochloride (Methadone Hydrochloride)	C
Meloxicam (Meloxicam)	C
Homeopathic Drug	C

Date:04/01/04ISR Number: 4332381-1Report Type:Expedited (15-DaCompany Report #2004019964
Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3600 MG		Biliary Tract Disorder Cholestasis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other (DAILY), ORAL		Diverticulum Hepatocellular Damage Nausea Vomiting	Professional				

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Date:04/01/04ISR Number: 4332382-3Report Type:Expedited (15-DaCompany Report #2004019953
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness	Foreign	Neurontin			
3600 MG		Hypoaesthesia	Consumer	(Gabapentin)	PS		ORAL
(DAILY), ORAL		Sexual Dysfunction					
		Somnolence					
		Weight Increased					

Date:04/01/04ISR Number: 4332384-7Report Type:Expedited (15-DaCompany Report #2004019966
Age:91 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cerebrovascular Accident	Foreign	Neurontin			
			Health	(Gabapentin)	PS		
			Professional				
			Company				
			Representative				

Date:04/01/04ISR Number: 4333379-XReport Type:Expedited (15-DaCompany Report #2004019975
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Unevaluable Event	Consumer	Neurontin			
				(Gabapentin)	PS		

Date:04/01/04ISR Number: 4333598-2Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Acrochordon
Initial or Prolonged	Angina Unstable
Disability	Arteriovenous Fistula
Other	Bladder Constriction
	Blood Pressure Increased

Brain Neoplasm
Cardiac Arrest
Convulsion
Dermatitis Contact
Dermatologic Examination
Abnormal
Electrocardiogram Qrs
Complex Shortened
Enchondromatosis
Excoriation
Gallbladder Disorder
Hypoaesthesia
Impaired Driving Ability
Intervertebral Disc
Disorder
Intervertebral Disc
Displacement
Joint Injury
Lower Gastrointestinal
Haemorrhage
Lumbar Radiculopathy
Lymphocyte Percentage
Decreased

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Marital Problem Nervous System Disorder Pain					
400 MG (BID),		Peptic Ulcer Protein Total Decreased	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Pruritus	Professional				
		Red Blood Cell Sedimentation Rate Increased Skin Irritation Stress Syncope	Company Representative	Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin)	SS C C C C		

Date:04/02/04ISR Number: 4330604-6Report Type:Expedited (15-DaCompany Report #FR-BRISTOL-MYERS SQUIBB COMPANY-12511176
Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged tablets		Cerebral Atrophy Electroencephalogram	Health Professional	Aprovel Tabs	PS	Bristol-Myers Squibb Company	ORAL
1 tablet		Abnormal		Lasilix	SS		ORAL
daily		Fall					
		Headache Vertigo		Neurontin Contramal Effexor Amlor Glucor Ogast Difrarel Dafalgan Vastarel	SS SS SS C C C C C C C		ORAL ORAL ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Abasia	Consumer	Neurontin			
Initial or Prolonged	Dysstasia		(Gabapentin)	PS		ORAL
ORAL						
Other	Pain		Losartan Potassium	C		
	Pain In Extremity		Insulin	C		
	Pulmonary Oedema		Amlodipine Besilate	C		
			All Other			
			Therapeutic Products	C		
			Diuretics	C		
			Amitriptyline	C		
			Paracetamol	C		

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Freedom Of Information (FOI) Report

Date:04/02/04ISR Number: 4333744-0Report Type:Expedited (15-DaCompany Report #2004020133
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Impaired Driving Ability Visual Acuity Reduced Visual Disturbance	Consumer	Neurontin (Gabapentin)	PS		

Date:04/02/04ISR Number: 4333746-4Report Type:Expedited (15-DaCompany Report #2004020197
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:04/02/04ISR Number: 4333747-6Report Type:Expedited (15-DaCompany Report #2004020149
Age:59 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 2700 MG		Gout Laboratory Test Abnormal	Health Professional	Neurontin (Gabapentin)	PS		ORAL
(BID), ORAL				All Other Therapeutic Products	C		

Date:04/02/04ISR Number: 4333834-2Report Type:Expedited (15-DaCompany Report #2004019978
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged ORAL		Asthma Cardiac Disorder	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other		Dizziness					

Drug Ineffective
Dyspepsia
Pain
Pharmaceutical Product
Complaint

Date:04/02/04ISR Number: 4333838-XReport Type:Expedited (15-DaCompany Report #2004019974

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Neurontin			
1800 MG		Anaemia Vitamin B12		(Gabapentin)	PS		ORAL
(TID), ORAL		Deficiency					
		Herpes Zoster		Panadeine (Codeine			
				Phsophate,	C		
				Paracetamol)	C		
				Benadryl			
				Albuterol			
				(Salbutamol)	C		

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Date:04/02/04ISR Number: 4333843-3Report Type:Expedited (15-DaCompany Report #KII-2002-0000001
Age:27 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Infarction	Health Professional Company Representative	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		
60 MG, BID				Remeron (Mirtazapine)	SS		
45 MG, HS				Neurontin (Gabapentin)	SS		
600 MG, TID				Valium (Diazepam)	SS		
10 MG, BID				Paxil (Paroxetine Hydrochloride)	SS		
25 MG, QD, PRN							

Date:04/02/04ISR Number: 4333900-1Report Type:Expedited (15-DaCompany Report #KII-2000-0002070
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Clonic Convulsion	Health Professional Company Representative	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		
				Elavil (Amitriptyline Hydrochloride)	SS		
				Neurontin (Gabapentin)	SS		

Date:04/02/04ISR Number: 4333964-5Report Type:Expedited (15-DaCompany Report #2003119698
Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Blindness Unilateral	Consumer	Neurontin			
Other		Body Height Decreased	Health	(Gabapentin)	PS		ORAL
600 MG							
(UNKNOWN);		Cataract	Professional				
ORAL (SEE		Choroidal					
IMAGE)		Neovascularisation					
		Condition Aggravated		Piroxicam	C		
		Drug Ineffective		Atorvastatin	C		
		Retinal Oedema		Lisinopril	C		
		Somnolence		Verapamil	C		
				Furosemide	C		
				Potassium	C		

Date:04/02/04ISR Number: 4333966-9Report Type:Expedited (15-DaCompany Report #2004019968
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation	Consumer	Neurontin			
400 MG		Cerebrovascular Accident		(Gabapentin)	PS		ORAL
(UNKNOWN);		Confusional State					
ORAL		Convulsion					
		Fall		Clopidogrel Sulfate	C		
		Headache		Vicodin			
		Loss Of Consciousness		(Paracetamol,			
		Migraine		Hydrocodone			

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Freedom Of Information (FOI) Report

Bitartrate) C
 Diazepam C
 Fexofenadine
 Hydrochloride C
 Budesonide C
 Furosemide C
 Potassium C

Date:04/02/04ISR Number: 4334053-6Report Type:Expedited (15-DaCompany Report #2004021348
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State	Foreign	Neurontin (Tablets)			
		Feeling Abnormal	Consumer	(Gabapentin)	PS		ORAL
1200 MG		Pyrexia					
(TID), ORAL		Sleep Talking		Lamotrigine			
		Somnolence		(Lamotrigine)	C		
		Vomiting		Theophylline			
		Wheelchair User		(Theophylline)	C		
				Budesonide			
				(Budesonide)	C		
				Salbutamol			
				(Salbutamol)	C		
				Citalopram			
				Hydrobromide			
				(Citalopram			
				Hydrobromide)	C		
				Warfarin Sodium			
				(Warfarin Sodium)	C		
				Trimethoprim			
				(Trimethoprim)	C		
				Ciprofloxacin			
				(Ciprofloxacin)	C		
				Potassium Chloride			
				(Potassium Chloride)	C		

Date:04/02/04ISR Number: 4334065-2Report Type:Expedited (15-DaCompany Report #200401867
 Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other	Convulsion	Foreign	Neurontin		
300 MG	Impaired Driving Ability	Health	(Gabapentin)	PS	ORAL
(DAILY), ORAL	Loss Of Consciousness	Professional			
	Somnolence	Company			
		Representative			

Date:04/02/04ISR Number: 4334068-8Report Type:Expedited (15-DaCompany Report #2004020054
Age:60 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Cholecystitis
Initial or Prolonged	Cholelithiasis
Other	Cholestasis
	Hepatitis
	Leukocytosis
	Neutrophilia
	Pain
	Prothrombin Level
	Decreased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Pyrexia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2400 MG (QID)		Foreign Literature	Neurontin (Tablets) (Gabapentin)	PS		
(DAILY)		Health Professional	Clonazepam (Clonazepam)	SS		

Date:04/02/04ISR Number: 4334078-0Report Type:Expedited (15-DaCompany Report #2004019973
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 800 MG		Fall Hypertension	Foreign Consumer	Gabapentin (Tablets) (Gabapentin)	PS		ORAL
(DAILY), ORAL		Post-Traumatic Amnestic Disorder		Phenytoin (Phenytoin Sodium) Clonazepam (Clonazepam) Ginkgo Biloba (Ginkgo Biloba)	C C C		

Date:04/02/04ISR Number: 4334127-XReport Type:Expedited (15-DaCompany Report #2004020473
Age:91 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - Initial or Prolonged		Cerebrovascular Accident	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:04/05/04ISR Number: 4332960-1Report Type:Direct Company Report #CTU 215971
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	40 MGS A DAY	Drug Ineffective		Celexa 40 Mgs Forest	PS	Forest	
Hospitalization - Initial or Prolonged Other Required Intervention to Prevent Permanent Impairment/Damage				Neurontin 2700 Mg Pfizer/Parkedavis Neurontin Seroquel Clonazepam Thyroid Celexa ...	SS C C C C C	Pfizer/Parkedavis	

Date:04/06/04ISR Number: 4333037-1Report Type:Expedited (15-DaCompany Report #FR-ROCHE-348810
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	18 DAY	Sudden Death		Valium	PS	Roche	ORAL
	56 DAY			Neurontin	SS		ORAL
	56 DAY			Deroxat	SS		ORAL
	120 DAY			Gardenal	SS		ORAL
INTRAMUSCULAR		1 DAY		Tercian	SS		
	18 DAY			Noctamide	SS		ORAL

Freedom Of Information (FOI) Report

Date:04/06/04ISR Number: 4335515-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000697

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 400 MG (BID), Disability ORAL Other	Acrochordon Angina Unstable Back Pain Cardiac Arrest Cardiac Telemetry Abnormal Cognitive Disorder Conversion Disorder Convulsion Coronary Artery Disease Dermatitis Contact Dissociative Disorder Dysphemia Electrocardiogram Q Waves Electrocardiogram Qrs Complex Shortened Enchondromatosis Ependymoma Excoriation Fatigue Gallbladder Disorder Hypoaesthesia Impaired Driving Ability Intervertebral Disc Disorder Intervertebral Disc Displacement Joint Injury Lower Gastrointestinal Haemorrhage Lumbar Radiculopathy Lymphocyte Count Decreased Marital Problem Memory Impairment Mobility Decreased Myocardial Infarction Nervous System Disorder Pain Peptic Ulcer	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Valproate Sodium (Valproate Sodim) Buspirone Hydrochloride Fluoxetine Hydrochloride Omeprazole Capsaicin	PS SS C C C C		ORAL

Post Procedural
Complication
Protein Total Decreased
Pruritus
Skin Irritation
Somnolence
Stress
Syncope

Date:04/06/04ISR Number: 4335545-6Report Type:Expedited (15-DaCompany Report #2004021454
Age:80 YR Gender:Male I/FU:I

Outcome PT
Other Abasia
Back Pain
Body Height Decreased
Coronary Artery Surgery

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Ineffective Exostosis Fall	Report Source	Product	Role	Manufacturer	Route
10 MG	(DAILY), ORAL	Nerve Compression Neuropathy	Health Professional	Lipitor (Atorvastatin)	PS		ORAL
300 MG (TID), ORAL		Osteoarthritis Pernicious Anaemia Post Procedural Complication		Neurontin (Gabapentin)	SS		ORAL
		Sciatica Somnolence		Diltiazem Hydrochloride	C		
				Levothyroxine Sodium	C		
				Acetylsalicylic Acid	C		
				Calcium	C		
				Tocopherol	C		

Date:04/06/04ISR Number: 4335782-0Report Type:Expedited (15-DaCompany Report #2004021645
Age: Gender:Male I/FU:I

Outcome Dose Other	Duration	PT Drug Dependence	Report Source	Product	Role	Manufacturer	Route
			Health Professional	Neurontin (Gabapentin)	PS		

Date:04/06/04ISR Number: 4335858-8Report Type:Expedited (15-DaCompany Report #2004021639
Age:47 YR Gender:Male I/FU:I

Outcome Dose Other	Duration	PT Drug Dependence	Report Source	Product	Role	Manufacturer	Route
			Health Professional	Neurontin (Gabapentin)	PS		
				Alprazolam (Alprazolam)	C		
				Carisoprodol (Carisoprodol)	C		
				Clonazepam (Clonazepam)	C		
				Hydrocodone			

(Hydrocodone)

C

Date:04/06/04ISR Number: 4335865-5Report Type:Expedited (15-DaCompany Report #2004012534
Age:92 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG DAILY, ORAL	Depressed Level Of Consciousness Status Epilepticus	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL			Zoxan Lp (Doxazosin)	SS		ORAL
ORAL			Nitrendipine (Nitrendipine)	SS		ORAL
ORAL			Acetylsalicylate Lysin (Acetylsalicylate Lysine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/07/04ISR Number: 4336226-5Report Type:Expedited (15-DaCompany Report #2004018059

Age:61 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 800 MG DAILY, ORAL	Hepatitis Cholestatic Hepatitis Toxic Hypertension	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
6 MG DAILY, ORAL			Piretanide Sodium (Piretanide Sodium)	SS		ORAL
			Morphine Sulfate (Morphine Sulfate) Distigmine Bromide (Distigmine Bromide)	C C		

Date:04/07/04ISR Number: 4336913-9Report Type:Expedited (15-DaCompany Report #2004021457

Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 800 MG (DAILY), ORAL	Anxiety Blood Triglycerides Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL	Depressed Level Of Consciousness Depression Hot Flush		Pseudoephedrine Hydrochloride (Pseudoephedrine Hydrochloride)	SS		ORAL
ORAL	Hypertension Insomnia Menopause Muscle Spasms		Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL
ORAL	Night Cramps Pain Panic Attack Somnolence		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS		ORAL

Spinal Laminectomy

Date:04/07/04ISR Number: 4336915-2Report Type:Expedited (15-DaCompany Report #2004007885
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Partial Seizures	Consumer	Neurontin			
1200 MG		Vaginal Haemorrhage	Health	(Gabapentin)	PS		ORAL
(TID), ORAL			Professional				
				Levetiracetam			
				(Levetiracetam)	C		
				Fluoxetine			
				(Fluoxetine)	C		
				Phenobarbital			
				(Phenobarbital)	C		

Date:04/08/04ISR Number: 4336732-3Report Type:Expedited (15-DaCompany Report #2004021469
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysgeusia	Consumer	Neurontin (Tablets)			
3200 MG		Ovarian Cancer		(Gabapentin)	PS		ORAL
(QID), ORAL		Pharmaceutical Product					
		Complaint		Lisinopril			
				(Lisinopril)	C		

Freedom Of Information (FOI) Report

Glibenclamide
(Glibenclamide) C

Date:04/08/04ISR Number: 4336984-XReport Type:Expedited (15-DaCompany Report #US-SHR-04-021606
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Alanine Aminotransferase Increased Aspartate Aminotransferase	Health Professional	Betaseron (Interferon Beta-1b) Injection, 250ug	PS		
SUBCUTANEOUS	8 MIU,	EVERY					
2D,		Increased					
SUBCUTANEOUS		Blood Bilirubin Increased Liver Function Test		Zanaflex (Tizanidine Hydrochloride)	SS		ORAL
ORAL		Abnormal		Neurontin /Unk/ (Gabapentin)	SS		ORAL
ORAL				Potassium (Potassium)	SS		ORAL

Date:04/08/04ISR Number: 4337317-5Report Type:Expedited (15-DaCompany Report #2004021476
Age:85 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (TID)		Malaise Oxygen Saturation Decreased Syncope Vasovagal	Foreign Health Professional Company Representative	Neurontin (Tablets) (Gabapentin) Acetylsalicylate Lysine	PS C		

Date:04/08/04ISR Number: 4337377-1Report Type:Expedited (15-DaCompany Report #2004021675
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Caesarean Section	Foreign	Neurontin			
Other		Failure To Thrive	Health	(Gabapentin)	PS		
PLACENTAL		Maternal Drugs Affecting Foetus	Professional	Clonazepam (Clonazepam)	C		
		Premature Baby		Dosulepin			
		Small For Dates Baby		(Dosulepin)	C		

Date:04/08/04ISR Number: 4337394-1Report Type:Expedited (15-DaCompany Report #2004021473
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Effect Decreased	Foreign	Gabapentin			
Other		Drug Interaction	Health	(Gabapentin)	PS		ORAL
300 MG		International Normalised	Professional				
(DAILY), ORAL		Ratio Decreased		Warfarin (Warfarin)	SS		
				Glyceryl Trinitrate (Glyceryl Trinitrate)	C		
				Fexofenadine (Fexofenadine)	C		
				Beclometasone (Beclometasone)	C		
				Salbutamol (Salbutamol)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Doxazosin
(Doxazosin) C

Date:04/08/04ISR Number: 4337401-6Report Type:Expedited (15-DaCompany Report #2004021671
Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Caesarean Section	Foreign Health	Neurontin (Gabapentin)	PS		
		Complications Of Maternal Exposure To Therapeutic Drugs	Professional	Clonazepam (Clonazepam)	C		
		Pregnancy		Dosulepin (Dosulepin)	C		
		Premature Baby					

Date:04/09/04ISR Number: 4338378-XReport Type:Expedited (15-DaCompany Report #2004021934
Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1800 MG (TID)						
		Abnormal Behaviour Anxiety	Foreign Health	Neurontin (Gabapentin)	PS		
		Insomnia	Professional Company Representative	Indapamide (Indapamide)	C		
		Suicidal Ideation		Trimetazidine (Trimetazidine)	C		

Date:04/09/04ISR Number: 4338379-1Report Type:Expedited (15-DaCompany Report #2004022191
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death							
Other		Completed Suicide	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (DAILY), ORAL		Balance Disorder Condition Aggravated Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL
20 MG (DAILY),		Drug Interaction Fall Migraine Nightmare Suicide Attempt		Escitalopram (Escitalopram)	SS		
				Allegra-D (Fexofenadine, Pseudoephedrine Hydrochloride) Propacet (Dextropropoxyphene Napsilate, Paracetamol) Pantoprazole (Pantoprazole) Docusate (Docusate)	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/09/04ISR Number: 4338880-0Report Type:Expedited (15-DaCompany Report #2004021849

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder	Consumer	Nardil (Phenelzine Sulfate)	PS		ORAL
	45 MG (TID),	Breast Cancer					
	ORAL	Carpal Tunnel Syndrome					
		Drug Effect Decreased		Neurontin (Gabapentin)	SS		ORAL
	600 MG (BID),	Electroconvulsive Therapy					
	ORAL	Feeling Abnormal					
		Mastectomy		Alprazolam	C		
		Treatment Noncompliance		Triobe (Cyanocobalamin, Folic Acid, Pyridoxine) Vitamins	C		

Date:04/09/04ISR Number: 4338971-4Report Type:Expedited (15-DaCompany Report #2003125029

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cognitive Disorder	Consumer	Neurontin (Gabapentin)	PS		
	400 MG (TID)	Feeling Abnormal	Health				
		Somnolence	Professional	Zoloft (Sertraline)	SS		
	200 MG DAILY)			Escitalopram (Escitalopram)	C		
				Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		
				Clonazepam (Clonazepam)	C		

Date:04/12/04ISR Number: 4339202-1Report Type:Expedited (15-DaCompany Report #2004021469

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysgeusia Ovarian Cancer	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
3200 MG		Pharmaceutical Product					
(QID), ORAL		Complaint		Lisinopril (Lisinopril)	C		
				Glibenclamice (Glibenclamide)	C		

Date:04/12/04ISR Number: 4339395-6Report Type:Expedited (15-DaCompany Report #2004011148
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Body Temperature Decreased Decreased Appetite Hypotension Insomnia Paralysis Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/12/04ISR Number: 4339396-8Report Type:Expedited (15-DaCompany Report #2004021959

Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anger	Consumer	Neurontin (Tablets)			
ORAL		Blindness		(Gabapentin)	PS		ORAL
		Eye Injury		Colchicine			
		Headache		(Colchicine)	C		
		Skin Discolouration		Estrogens Conjugated			
		Visual Disturbance		(Estrogens			
				Conjugated)	C		
				Vicoprofen			
				(Hydrocodone			
				Bitartrate,			
				Ibuprofen)	C		
				All Other			
				Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	C		

Date:04/13/04ISR Number: 4337529-0Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0313585A

Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Optic Neuropathy	Health	Lamictal	PS	Glaxosmithkline	ORAL
300MG per day		Visual Acuity Reduced	Professional	Neurontin	SS		ORAL
800MG Twice		Visual Field Defect					
per day				Sectral	C		ORAL
				Mogadon	C		ORAL

Date:04/13/04ISR Number: 4340657-7Report Type:Expedited (15-DaCompany Report #2004022257

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin			

(Gabapentin)

PS

Date:04/13/04ISR Number: 4340658-9Report Type:Expedited (15-DaCompany Report #2004022259
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin	PS		

Date:04/13/04ISR Number: 4340659-0Report Type:Expedited (15-DaCompany Report #2004022286
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:04/13/04ISR Number: 4340808-4Report Type:Expedited (15-DaCompany Report #2004022537
Age: Gender:Male I/FU:I

Outcome
Hospitalization -
Initial or Prolonged

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Freedom Of Information (FOI) Report

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
3600 MG (7 TIMES DAILY), ORAL		Amnesia Anticonvulsant Drug Level Decreased Anticonvulsant Drug Level	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
500 MG, ORAL		Increased Aphasia Balance Disorder Breast Mass Breast Pain Convulsion Decreased Appetite Depressed Level Of Consciousness Drug Toxicity Dysuria Fall Gynaecomastia Headache Hypoaesthesia Impatience Insomnia Liver Function Test Abnormal Mood Swings Nausea Oral Intake Reduced Personality Change Pruritus Generalised Thermal Burn Tooth Disorder Tooth Injury Tooth Loss Vomiting Weight Decreased Weight Increased		Dilantin Suspension (Phenytoin Sodium) Protein Supplements (Protein Supplements) Phenobarbital (Phenobarbital)	SS C C		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Arthritis Bacterial Balance Disorder	Consumer	Neurontin (Gabapentin)	PS		
100 MG (DAILY),		Blood Pressure Increased					
2.5 MG (DAILY), ORAL		Musculoskeletal Stiffness Neuropathy Peripheral Pain Vestibular Disorder		Norvasc (Amlodipine)	SS		ORAL
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Glyceryl Trinitrate (Glyceryl Trinitrate)	C		
				Metoprolol (Metoprolol)	C		
				Ezetimibe (Ezetimibe)	C		
				Alendronate Sodium (Alendronate Sodium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Terazosin
 Hydrochloride
 (Terazosin
 Hydrochloride) C
 Finasteride
 (Finasteride) C
 Simvastatin
 (Simvastatin) C
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C

Date:04/13/04ISR Number: 4340849-7Report Type:Expedited (15-DaCompany Report #2004022216
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dysphagia Hypoaesthesia	Consumer	Neurontin (Gabapentin)	PS		
1) UNKNOWN;		Multiple Sclerosis					
2) 1800 MG (600, TID),		Pharmaceutical Product Complaint Visual Acuity Reduced		All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:04/13/04ISR Number: 4340862-XReport Type:Expedited (15-DaCompany Report #2004022253
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:04/13/04ISR Number: 4340864-3Report Type:Expedited (15-DaCompany Report #2004022252

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:04/13/04ISR Number: 4340865-5Report Type:Expedited (15-DaCompany Report #2004022254

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:04/13/04ISR Number: 4340957-0Report Type:Expedited (15-DaCompany Report #2004022255

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/13/04ISR Number: 4340959-4Report Type:Expedited (15-DaCompany Report #2004022256

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:04/14/04ISR Number: 4338495-4Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0311384A

Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation	Consumer	Deroxat	PS	Glaxosmithkline	ORAL
Life-Threatening		Fall		Noctamid	SS		ORAL
Sudden Death				Valium	SS		ORAL
				Tercian	SS		
	INTRAMUSCULAR			Neurontin	SS		ORAL
				Gardenal	SS		ORAL

Date:04/14/04ISR Number: 4339954-0Report Type:Expedited (15-DaCompany Report #2004022432

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Astigmatism Dysarthria	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Endometriosis Fatigue Libido Decreased Muscle Twitching Visual Disturbance		Propacet (Dextropropoxyphene Napsilate, Paracetamol) Estratest Hs (Estrogens Esterified, Methyltestosterone)	C C		

Paroxetine
 Hydrochloride
 (Paroxetine
 Hydrochloride) C
 Nortriptyline
 (Nortriptyline) C
 Buspirone
 Hydrochloride
 (Buspirone
 Hydrochloride) C
 Clonazepam
 (Clonazepam) C

Date:04/14/04ISR Number: 4340537-7Report Type:Expedited (15-DaCompany Report #RENA-10959
 Age:57 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 2.4 G DAILY	Scleroderma	Foreign	Renagel	PS		ORAL
Initial or Prolonged PO		Health				
		Professional Company	Beloc Zok Neurontin	SS SS		
300 MG QD		Representative Other	Phos-Ex Rohypnol Marcumar Restex Einsalpha	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vigantoletten C

Date:04/14/04ISR Number: 4340600-0Report Type:Expedited (15-DaCompany Report #2004019966
Age:91 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:04/15/04ISR Number: 4339364-6Report Type:Expedited (15-DaCompany Report #FR-BRISTOL-MYERS SQUIBB COMPANY-12511176
Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged tablets		Bradyarrhythmia Cerebral Atrophy		Aprovel Tabs	PS	Bristol-Myers Squibb Company	ORAL
1 tablet		Fall		Lasilix	SS		ORAL
daily		Headache					
		Pruritus		Neurontin	SS		ORAL
		Rash		Contramal	SS		ORAL
		Self-Medication		Effexor	SS		ORAL
		Toxic Skin Eruption		Amlor	C		
		Urticaria		Glucor	C		
		Vertigo		Ogast	C		
				Difrarel	C		
				Dafalgan	C		
				Vastarel	C		

Date:04/15/04ISR Number: 4341455-0Report Type:Expedited (15-DaCompany Report #200410158BFR
Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - 300 MG, TOTAL Initial or Prolonged DAILY, ORAL	Blister Eczema Erythema Fall Rash Pruritic Self-Medication Toxic Skin Eruption Urticaria	Foreign Health Professional Other	Glucor (Acarbose) Furosemide Neurontin (Gabapentin) Aprovel (Irbesartan) Contramal (Tramadol Hydrochloride) Effexor (Venlafaxine Hydrochloride) Ogast (Lansoprazole) Difrarel Dafalgan (Paracetamol) Amlor (Amlodipine Besilate)	PS SS SS SS SS SS SS SS SS SS	ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL
---	---	--	--	--	--

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

60 MG, TOTAL
 DAILY, ORAL

Vastarel
 (Trimetazidine
 Hydrochloride) SS ORAL

Date:04/15/04ISR Number: 4341500-2Report Type:Expedited (15-DaCompany Report #2004022888
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction Hypercoagulation	Foreign Health	Neurontin (Gabapentin)	PS		
2400 MG (TID)			Professional Company Representative	Fluindione (Fluindione) All Other Therapeutic Products	SS C		

Date:04/15/04ISR Number: 4342833-6Report Type:Expedited (15-DaCompany Report #2004022777
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Distension Burning Sensation	Consumer	Neurontin (Solution) (Gabapentin)	PS		ORAL
100 MG (DAILY AT BEDTIME), ORAL		Colitis Condition Aggravated					
		Eye Disorder Eye Pain Feeling Abnormal Gastritis Erosive Hot Flush		Sertraline Hydrochloride (Sertraline Hydrochloride) Diazepam (Diazepam) Dimeticone, Activated (Simeticone) Irbesartan (Irebesartan) Atenolol (Atenolol)	C C C C		

Levothyroxine Sodium
(Levothyroxine Sodium) C
Acetylsalicylic Acid
(Acetylsalicylic Acid) C
Zolpidem Tartrate
(Zolpidem Tartrate) C
Meclozine
Hydrochloride
(Meclozine Hydrochloride) C
Hyoscyamine Sulfate
(Hyoscyamine Sulfate) C

Date:04/16/04ISR Number: 4341816-XReport Type:Expedited (15-DaCompany Report #2004UW02477
Age:43 YR Gender:Male I/FU:F

Outcome PT
Death Coma
Coronary Artery
Atherosclerosis
Therapeutic Agent

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Toxicity

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Health Professional	Seroquel	PS		
			Neurontin	SS		
			Lexapro	SS		
			Clonazepam	SS		
			Oxycontin	C		

Date:04/16/04ISR Number: 4342357-6Report Type:Expedited (15-DaCompany Report #2004018051
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health Professional	Dilatin Suspension (Phenytoin Sodium)	PS		
Other		Pancreas Transplant Renal Transplant Unintended Pregnancy		Neurontin (Gabapentin)	SS		

Date:04/16/04ISR Number: 4342359-XReport Type:Expedited (15-DaCompany Report #2004023523
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Deafness Nuclear Magnetic Resonance Imaging Brain					
900 MG (TID)				Levothyroxine Sodium	C		
ORAL		Abnormal		Lidocaine Hydrochloride	C		

Date:04/16/04ISR Number: 4342361-8Report Type:Expedited (15-DaCompany Report #2004023410
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Blood Sodium Decreased	Consumer	Neurontin			

Initial or Prolonged 600 MG (BID) Other ORAL	Compression Fracture Convulsion Muscle Spasms Neuralgia	(Gabapentin)	PS	ORAL
		Ibuprofen Lotrel (Amlodipine, Benazepril Hydrochloride) Alendronate Sodium Provella-14 (Estrogens Conjugated, Medroxyprogesterone Acetate)	C C C C	

Date:04/16/04ISR Number: 4342367-9Report Type:Expedited (15-DaCompany Report #2004023332
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Intervertebral Disc Protrusion Pain	Consumer	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/16/04ISR Number: 4342423-5Report Type:Expedited (15-DaCompany Report #2004023520
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other Required Intervention to Prevent Permanent Impairment/Damage		Angioneurotic Oedema Contusion Dermatitis Exfoliative Dyspnoea Multiple Allergies Oedema Peripheral Pharmaceutical Product Complaint Pituitary Tumour	Consumer	Neurontin (Gabapentin) Atarax (Tablet) (Hydroxyzine Hydrochloride) Hydroxyzine Hydrochloride (Hydroxyzine Hydrozylchloride)	PS SS SS		
300 MG (25MG INCREMENTS IN DIVIDED DOSES.), UNKNOWN		Sinusitis Urticaria					
				Metropol (Metoprolol) All Other Therapeutic Products (All Other Therapeutic Products)	SS SS		

Date:04/16/04ISR Number: 4342428-4Report Type:Expedited (15-DaCompany Report #2004022216
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Choking Dysphagia	Consumer	Neurontin (Gabapentin)	PS		
1800 MG (600,TID), UNKNOWN		Hypoaesthesia Multiple Sclerosis Pharmaceutical Product Complaint		All Other Therapeutic Products			

Visual Acuity Reduced

(All Other
Therapeutic
Products)

C

Date:04/16/04ISR Number: 4342444-2Report Type:Expedited (15-DaCompany Report #2004012493
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bacterial Infection Condition Aggravated	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Gastric Infection Hernia Hiatus Hernia Implant Site Infection Neuropathy	Professional	Potassium Chloride (Potassium Chloride) Furosemide (Furosemide) Methadone (Methadone) Alprazolam (Alprazolam) Isosorbide Mononitrate (Isosorbide Mononitrate) Fenofibrate (Fenofibrate) Acetylsalicylic Acid	C C C C C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Acetylsalicylic Acid) C
 Colchicine (Colchicine) C
 Trandolapril (Trandolapril) C
 Atenolol (Atenolol) C
 Allopurinol (Allopurinol) C
 Oxygen (Oxygen) C

Date:04/16/04ISR Number: 4342556-3Report Type:Expedited (15-DaCompany Report #2004010454
 Age:73 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG Other (DAILY), ORAL	Brain Scan Abnormal Cerebral Atrophy Electroencephalogram	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
20 MG (DAILY), ORAL	Abnormal Fall Headache		Furosemide (Furosemide)	SS		ORAL
300 MG (DAILY), ORAL	Metabolic Disorder Toxic Skin Eruption Vertigo		Irbesartan (Irbasartan)	SS		ORAL
300 MG (DAILY), ORAL			Tramadol Hydrochloride (Tramadol Hydrochloride)	SS		ORAL
75 MG, ORAL			Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS		ORAL
300 MG , ORAL			Acarbose (Acarbose)	SS		ORAL
			Amlodipine Besilate			

(Amlodipine Besilate)	C
Lanxoprazole (Lansoprazole)	C
Diffrarel (Betacarotene, Myrtillus)	C
Paracetamol (Paracetamol)	C
Trimetazidine Hydrochloride (Trimetazidine Hydrochloride)	C
Betacarotene (Betacarotene)	C

Date:04/16/04ISR Number: 4342567-8Report Type:Expedited (15-DaCompany Report #2004024146
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG (DAILY), ORAL		Inadequate Analgesia Myoclonic Epilepsy	Foreign Health Professional	Gabaentin (Gabapentin)	PS		ORAL
10 MG				Amitriptyline (Amitriptyline)Q	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(DAILY), ORAL

Transtec
(Buprenorphine) SS

140 MCG/H

(DAILY)

Date:04/19/04ISR Number: 4342501-0Report Type:Direct
Age:66 YR Gender:Female I/FU:I

Company Report #CTU 216899

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Seroquel 300 Mg X 2	PS		
2 TIME DAY				Neurontin 600 Mg X 3	SS		
3 TIMES DAY							

Date:04/19/04ISR Number: 4343143-3Report Type:Expedited (15-DaCompany Report #A01200401659
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hepatic Steatosis Pancreatitis White Blood Cell Count	Health Professional	Plavix- (Clopidogrel Sulfate) - Tablet- 75 Mg	PS		ORAL
75 MG QD, ORAL		Increased		Dianben (Metformin Hydrochloride) - Tablet 850 Mg	SS		ORAL
850 MG, BID ORAL				Daonil - (Glibenclamide) - Tablet - 5 Mg	SS		ORAL
15 MG QD, ORAL				Pantoprazole Gabapentin	SS SS		

Date:04/19/04ISR Number: 4343427-9Report Type:Expedited (15-DaCompany Report #2004024700
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Accident At Work	Health Professional	Neurontin (Gabapentin)	PS		
				Zoloft (Sertraline)	SS		
				Valdecoxib (Valdecoxib)	SS		
				Lansoprazole (Lansoprazole)	SS		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:04/19/04ISR Number: 4343428-0Report Type:Expedited (15-DaCompany Report #2004022937
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Pressure Inadequately Controlled	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (THREE Other TIMES A DAY),		Neuritis					
ORAL		Somnolence					
				Enalapril Maleate (Enalapril Maleate)	SS		

Freedom Of Information (FOI) Report

All Other
Therapeutic Products
(All Other
Therapeutic
Products) C

Date:04/19/04ISR Number: 4343429-2Report Type:Expedited (15-DaCompany Report #2004024140
Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aura	Consumer	Dilantin Suspension (Phenytoin Sodium)	PS		
Other		Body Height Decreased		Neurontin (Gabapentin)	SS		
		Breast Cancer		Phenobarbital (Phenobarbital)	SS		
		Drug Ineffective		Metformin Hydrochloride (Metformin Hydrochloride)	C		
		Muscle Spasms		All Other Therapeutic Products (All Other Therapeutic Products)	C		
		Somnolence		Celecoxib (Celecoxib)	C		
				Tolterodine L-Tartrate (Tolterodine L-Tartrate)	C		
				Chondroitin Sulfate (Chondroitin Sulfate)	C		
				Glucosamine (Glucosamine)	C		
				Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotinamide, Tocopherol (Tocopherol)	C		
				Calcium With Vitamin	C		

D (Calcium
Phosphate, Calcium
Sodium Lactate,
Ergocalciferol) C

Date:04/19/04ISR Number: 4343641-2Report Type:Expedited (15-DaCompany Report #2004011175
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Constipation	Consumer	Neurontin (Solution)			
Other		Dyspepsia		(Gabapentin)	PS		ORAL
SEE IMAGE,		Pharmaceutical Product					
ORAL		Complaint		All Other			
		Tooth Loss		Therapeutic Products			
				(All Other			

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Freedom Of Information (FOI) Report

Therapeutic Products)	SS
All Other Therapeutic Products (All Other Therapeutic Products)	SS
Insulin (Insulin)	SS
Magnesium Hydroxide (Magnesium Hydroxide)	C
Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C
Fentanyl (Fentanyl)	C
Diazepam (Diazepam)	C
Doxazosin Mesilate (Doxazosin Mesilate)	C
Zolpidiem Tartrate (Zolpidem Tartrate)	C
Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Metformin Hydrochloride (Metformin Hydrochloride)	C
Rosuvastatin (Rosuvastatin)	C
Losartan Potassium (Losartan Potassium)	C

Date:04/19/04ISR Number: 4343643-6Report Type:Expedited (15-DaCompany Report #2004023104
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Consumer	Neurontin (Tablets)			
3600 (TID),		Pharmaceutical Product		(Gabapentin)	PS		ORAL
ORAL		Complaint		Topiramate			

(Topiramate)	C
Pravastatin Sodium	
(Pravastatin Sodium)	C
Sertraline	
Hydrochloride	
(Sertraline	
Hydrochloride)	C
Sotalol Hydrochloride	
(Sotalol	
Hydrochloride)	C
Buspirone	
Hydrochloride	
(Buspirone	
Hydrochloride)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/19/04ISR Number: 4343816-2Report Type:Expedited (15-DaCompany Report #2004010558
Age:52 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 200 MG (BID), Other ORAL	Acute Lymphocytic Leukaemia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
	Acute Myeloid Leukaemia	Professional				
	Anaemia		Estradiol (Estradiol)	C		
	Bronchitis		Escitalopram (Escitalopram)	C		
	Drug Ineffective					
	Dyspnoea					

Date:04/19/04ISR Number: 4343818-6Report Type:Expedited (15-DaCompany Report #2004009410
Age:60 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL Other	Arrhythmia Benign Prostatic	Consumer	Neurontin (Gabapentin)	PS		ORAL
	Hyperplasia		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		
	Micturition Urgency		Buspirone Hydrochloride (Buspirone Hydrochloride)	C		
	Pollakiuria		Testosterone (Testosterone)	C		
	Urinary Incontinence					

Date:04/20/04ISR Number: 4341559-2Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040301432
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death TRANSDERMAL Two 75 ug/hr patches (with	Cardiomegaly	Health	Duragesic	PS		
	Cardiopulmonary Failure	Professional				

biocclusive

Drug Abuser

dressings)

Drug Toxicity

Hepatic Congestion
 Mucous Membrane Disorder
 Nervous System Disorder
 Oesophagitis
 Pharmaceutical Product
 Complaint
 Pulmonary Congestion
 Pulmonary Oedema
 Scoliosis

Diazepam SS
 Oxycodone SS
 Morphine SS
 Gabapentin SS
 Benadryl C

Date:04/20/04ISR Number: 4341624-XReport Type:Expedited (15-DaCompany Report #200411201EU
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hepatic Steatosis Pancreatitis		Daonil	PS	Aventis Pharmaceuticals Inc.	ORAL ORAL
				Dianben	SS		ORAL
				Pantoprazol	SS		
				Plavix			
				/Unk/ Gabapentine	SS SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/20/04ISR Number: 4342331-XReport Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 216932

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bone Density Decreased		Neurontin 100 Mg	PS		
100 MG TID				Serzone	C		

Date:04/20/04ISR Number: 4344709-7Report Type:Expedited (15-DaCompany Report #2004022255
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Sudden Death	Consumer	Neurontin (Gabapentin)	PS		

Date:04/20/04ISR Number: 4344711-5Report Type:Expedited (15-DaCompany Report #2004022254
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Sudden Death	Consumer	Neurontin (Gabapentin)	PS		

Date:04/20/04ISR Number: 4344731-0Report Type:Expedited (15-DaCompany Report #2004022259
 Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Sudden Death	Consumer	Neurontin (Gabapentin)	PS		

UNKNOWN UNKNOWN

(UNKNOWN),

UNKNOWN

Date:04/20/04ISR Number: 4344738-3Report Type:Expedited (15-DaCompany Report #2004022257

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Consumer	Neurontin (Gabapentin)	PS		
UNKNOWN	UNKNOWN						
(UNKNOWN),							
UNKNOWN							

Date:04/20/04ISR Number: 4344739-5Report Type:Expedited (15-DaCompany Report #2004022256

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Consumer	Neurontin (Gabapentin)	PS		
UNKNOWN	UNKNOWN						
(UNKNOWN),							
UNKNOWN							

... C

Date:04/20/04ISR Number: 4344915-1Report Type:Expedited (15-DaCompany Report #2004024146

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Myoclonic Epilepsy Myoclonus	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
100 MG		Pain	Professional				
(DAILY), ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

10 MG				Amitriptyline (Amitriptyline)	SS		ORAL
(DAILY), ORAL							
140 MCG/H				Transtec (Buprenorphine)	SS		
(DAILY)							

Date:04/20/04ISR Number: 4345160-6Report Type:Expedited (15-DaCompany Report #2004022286
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Sudden Death	Consumer	Neurontin (Gabapentin)	PS		

Date:04/20/04ISR Number: 4345199-0Report Type:Expedited (15-DaCompany Report #2004022252
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Sudden Death	Consumer	Neurontin (Gabapentin)	PS		

Date:04/20/04ISR Number: 4345270-3Report Type:Expedited (15-DaCompany Report #2004024145
Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Test Abnormal Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG		Headache					
(THREE TIMES		Petit Mal Epilepsy					
A DAY), ORAL		Pharmaceutical Product Complaint		Topiramate (Topiramate)	C		

Stress
Treatment Noncompliance

Amitriptyline
Hydrochloride
(Amitriptyline
Hydrochloride) C
Fluoxetine
Hydrochloride
(Fluoxetine
Hydrochloride) C
Midodrine
Hydrochloride
(Midodrine
Hydrochloride) C
Fludrocortisone
Acetate
(Fludrocortisone
Acetate) C
Midrid
(Dichloralphenazone,
Isometheptene,
Paracetamol) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/20/04ISR Number: 4345272-7Report Type:Expedited (15-DaCompany Report #2004024435
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Pancreatic Operation	Health Professional	Neurontin (Gabapentin)	PS		

Date:04/20/04ISR Number: 4345365-4Report Type:Expedited (15-DaCompany Report #2004022253
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death		Sudden Death	Consumer	Neurontin (Gabapentin)	PS		

Date:04/21/04ISR Number: 4343794-6Report Type:Direct Company Report #CTU 217042
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to 300 MG PO QD Prevent Permanent Impairment/Damage		Sedation		Gabapentin 300 Mg Po Pd Pfizer	PS	Pfizer	ORAL

Date:04/21/04ISR Number: 4344397-XReport Type:Direct Company Report #CTU 217120
Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 100 MG PO Q8 Hospitalization - H Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage		Blood Sodium Decreased Dehydration		Gabapentin Amlodipine Aspirin Pletal Surfak Pepcid	PS C C C C C		ORAL

Ferrous Sulfate	C
Oxycontin	C
Quinine	C
Senna	C
Vicodin	C
Ambien	C
Reglan	C

Date:04/21/04ISR Number: 4345385-XReport Type:Expedited (15-DaCompany Report #2004025636
 Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG		Spinal Column Stenosis Vertigo	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/21/04ISR Number: 4345617-8Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040301432
 Age:22 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accident	Health	Duragesic (Fentanyl)			
		Cardiac Failure	Professional	Patch	PS		
TRANSDERMAL	75 UG/HR, 2	Cardio-Respiratory Arrest					
IN 72 HOUR,		Cardiomegaly					
TRANSDERMAL		Drug Abuser		Diazepam (Diazepam)	SS		
		Drug Level Decreased		Oxycodone			
		Drug Tolerance		(Oxycodone)	SS		
		Drug Toxicity		Morphine (Morphine)	SS		
		Hepatic Congestion		Gabapentin			
		Medication Error		(Gabapentin)	SS		
		Nervous System Disorder					
		Obesity					
		Oesophagitis					
		Pharmaceutical Product					
		Complaint					
		Pulmonary Oedema					
		Respiratory Failure					
		Scoliosis					

Date:04/21/04ISR Number: 4345696-8Report Type:Expedited (15-DaCompany Report #2004016355
 Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Eye Disorder	Health	Neurontin			
		Visual Acuity Reduced	Professional	(Gabapentin)	PS		
900 MG (TID)		Visual Disturbance		Eflornithine			
				Hydrochloride			
				(Eflornithine			
				Hydrochloride)	C		

Date:04/21/04ISR Number: 4346150-XReport Type:Expedited (15-DaCompany Report #2004024490
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability Other		Embolia Cutis Medicamentosa Gait Disturbance Injection Site Irritation Injection Site Pain	Consumer	Vistaril (Im) (Hydroxyzine Hydrochloride)	PS		
300 MG, ORAL				Neurontin (Gabapentin)	SS		ORAL
ORAL		Medication Error Muscle Spasms Necrosis		Pethidine Hydrochloride (Pethidine Hydrochloride)	SS		ORAL
20 MG				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL
(UNKNOWN), ORAL							
2 MG				Tizanidine Hydrochloride (Tizanidine Hydrochloride)	SS		ORAL
(UNKNOWN), ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/22/04ISR Number: 4347486-9Report Type:Expedited (15-DaCompany Report #USA-2004-0014543

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aortic Atherosclerosis	Health	Oxycodone			
		Hepatic Cirrhosis	Professional	Hydrochloride			
		Hepatitis C	Other	(Similar To Nda			
		Multiple Drug Overdose		20-553) (Oxycodone			
		Pulmonary Oedema		Hydrochloride)	PS		
				Ethanol (Ethanol)	SS		
				Diazepam (Diazepam)	SS		
				Temazepam			
				(Temazepam)	SS		
				Gabapentin			
				(Gabapentin)	SS		
				Citalopram			
				(Citalopram)	SS		
				Olanzapine			
				(Olanzapine)	SS		
				Oxazepam (Oxazepam)	SS		

Date:04/22/04ISR Number: 4347585-1Report Type:Expedited (15-DaCompany Report #2004019978

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abdominal Discomfort	Consumer	Neurontin (Tablets)			
Initial or Prolonged		Asthma	Health	(Gabapentin)	PS		ORAL
ORAL		Cardiac Disorder	Professional	Acetylsalicylic Acid			
Other		Dizziness		(Acetylsalicylic			
		Drug Ineffective		Acid)	C		
		Heart Valve Replacement		Warfarin Sodium			
		Pain		(Warfarin Sodium)	C		
		Pharmaceutical Product		Metoprolol Tartrate			
		Complaint		(Metoprolol			
				Tartrate)	C		
				Furosemide			
				(Furosemide)	C		

Date:04/22/04ISR Number: 4347597-8Report Type:Expedited (15-DaCompany Report #2004010782

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Calcium Decreased Blood Creatinine	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 MG (QID), ORAL		Increased Blood Urea Increased Drug Ineffective Hearing Impaired Oedema Pain Serum Ferritin Increased Vision Blurred Xanthopsia		All Other Therapeutic Products (All Other Therapeutic Products) Hydromorphone (Hydromorphone) Potassium Chloride (Potassium Chloride) Insulin Glargine (Insulin Glargine) Insulin Lispro (Insulin Lispro) Atorvastatin (Atorvastatin) Magnesium	SS C C C C		

Freedom Of Information (FOI) Report

(Magnesium)	C
Omeprazole	
(Omeprazole)	C
Oxycodone	
Hydrochloride	
(Oxycodone	
Hydrochloride)	C
Pentoxifylline	
(Pentoxifylline)	C
Prednisone	
(Prednisone)	C
Torasemide	
(Torasemide)	C
Calcium Carbonate	
(Calcium Carbonate)	C
Paracetamol	
(Paracetamol)	C
Ascorbic Acid	
(Ascorbic Acid)	C
Tocopherol	
(Tocopherol)	C
Warfarin (Warfarin)	C
Metolazone	
(Metolazone)	C
Cetirizine	
Hydrochloride	
(Cetirizine	
Hydrochloride)	C
Allopurinol	
(Allopurinol)	C
Darbepoetin Alfa	
(Darbepoetin Alfa)	C
Diphenhydramine	
Hydrochloride	
(Diphenhydramine	
Hydrochloride)	C
Calcitrol	
(Ergocalciferol,	
Retinol, Calcium	
Carbonate)	C
Mycophenolate	
Mofetil	
(Mycophenolate	
Mofetil)	C
Clonazepam	
(Clonazepam)	C
Colchicine	
(Colchicine)	C

Losartan Potasium	
(Losartan Potassium)	C
Folic Acid (Folic	
Acid)	C
Furosemide	
(Furosemide)	C
Amitriptyline	
(Amitriptyline)	C
Loratadine	
(Loratadine)	C
Eflornithine	
Hydrochloride	
(Eflornithine	
Hydrochloride)	C

Freedom Of Information (FOI) Report

Glucose (Glucose) C
 Glucagon (Glucagon) C
 Methenamine
 Hippurate
 (Methenamine
 Hippurate) C
 Ondansetron
 Hydrochloride
 (Ondansetron
 Hydrochloride) C

Date:04/22/04ISR Number: 4347602-9Report Type:Expedited (15-DaCompany Report #2004025316
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Depression Overdose	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							
Other							

Date:04/22/04ISR Number: 4347637-6Report Type:Expedited (15-DaCompany Report #2004025134
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety Cognitive Disorder	Consumer	Neurontin (Gabapentin)	PS		
UNKNOWN,							
		Panic Disorder					
UNKNOWN							
		Weight Increased					

Date:04/22/04ISR Number: 4347906-XReport Type:Expedited (15-DaCompany Report #2004025366
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Difficulty In Walking Drug Ineffective	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1500 MG (FOUR							

TIMES A DAY),

Dyspnoea

Feeling Abnormal

ORAL

Kyphosis
Memory Impairment
Palpitations
Pharmaceutical Product
Complaint

Vicodin (Hydrocodone
Bitartrate,
Paracetamol) C

Date:04/22/04ISR Number: 4347915-0Report Type:Expedited (15-DaCompany Report #2004024707

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 4800 (TID)		Anaemia Fibromyalgia	Consumer	Neurontin (Gabapentin)	PS		
Other		Multiple Sclerosis		Levothyroxine Sodium (Levothyroxine Sodium) Esomeprazole (Esomeprazole) Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C C C		

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Freedom Of Information (FOI) Report

Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C
Lorazepam (Lorazepam)	C
Tramadol Hydrochloride (Tramadol Hydrochloride)	C
Paracetamol (Paracetamol)	C
Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride)	C
Tizanidine Hydrochloride (Tizanidine Hydrochloride)	C
Rofecoxib (Rofecoxib)	C

Date:04/22/04ISR Number: 4348005-3Report Type:Expedited (15-DaCompany Report #KII-2004-0009569
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged Other	Agitation Anoxic Encephalopathy Atrial Fibrillation Cerebellar Infarction Coma Confusional State Hallucination Hemiparesis Multiple Drug Overdose Somnolence	Study Health Professional Other	Morphine Sulfate (Similar To Nda-19-516) (Morphine Sulfate) Unknown Neurontin (Gabapentin) Tramadol			PS SS SS

Date:04/23/04ISR Number: 4348415-4Report Type:Expedited (15-DaCompany Report #2004024140
Age:83 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Body Height Decreased	Consumer	Dilantin Suspension			
		Breast Cancer		(Phenytoin Sodium)	PS		
		Drug Ineffective		Neurontin			
		Muscle Spasms		(Gabapentin)	SS		
		Somnolence		Phenobarbital			
		Stress		(Phenobarbital)	SS		
				Metformin			
				Hydrochloride	C		
				All Other			
				Therapeutic Products	C		
				Celecoxib	C		
				Tolterodine			
				L-Tartrate	C		
				Chondroitin Sulfate	C		
				Glucosamine	C		
				Multivitamins			
				(Ascorbic Acid,			

Freedom Of Information (FOI) Report

Ergocalciferol,
 Folic Acid,
 Nicotinamide, C
 Tocopherol C
 Calcium With Vitamin
 D (Calcium
 Phosphate, Calcium
 Sodium Lactate,
 Ergocalciferol) C

Date:04/23/04ISR Number: 4348781-XReport Type:Expedited (15-DaCompany Report #2004024145
 Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2400 MG (THREE TIMES A DAY), ORAL		Condition Aggravated Convulsion Headache Laboratory Test Abnormal Medication Error Stress	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Topiramate (Topiramate)	C		
				Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
				Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		
				Midodrine Hydrochloride (Midodrine Hydrochloride)	C		
				F;Udrocortisone Acetate (Fludrocortisone Acetate)	C		
				Midrid (Dichloralphenazone, Isometheptene, Paracetamol)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (TID), Other ORAL	Abdominal Pain Abnormal Faeces Arthritis Blood Glucose Decreased Body Height Decreased Condition Aggravated Fatigue Food Allergy Hypersensitivity Irritable Bowel Syndrome Pancreatic Enzymes Abnormal Reaction To Medical Agent Preservatives Vision Blurred	Consumer Health Professional	Neurontin (Gabapentin) Glucosamine Sulfate/Minerals/Mul tivitamins (Glucosamine Sulfate, Vitamins Hyoscyamine Sulfate (Hyoscyamine Sulfate) Pancrelipase	PS SS SS C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/26/04ISR Number: 4349494-0Report Type:Expedited (15-DaCompany Report #2004010456

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		No Adverse Drug Effect	Health Professional	Neurontin (Gabapentin)	PS		

Date:04/26/04ISR Number: 4349814-7Report Type:Expedited (15-DaCompany Report #2004012533

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion	Consumer	Dilantin Suspension (Phenytoin Sodium)	PS		
Other		Hot Flush		Neurontin (Gabapentin)	SS		
2400 MG TID		Muscle Twitching		Phenobarbital (Phenobarbital)	C		
				Valproate Semisodium (Valproate Semisodium)	C		
				Clonazepam (Clonazepam)	C		

Date:04/27/04ISR Number: 4349013-9Report Type:Direct Company Report #CTU 217413

Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Abnormal Behaviour		Neurontin	PS		
400 MG ONCE		Amnesia					
DAILY		Educational Problem					

Date:04/27/04ISR Number: 4350475-1Report Type:Expedited (15-DaCompany Report #2004025953

Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Dizziness	Consumer	Neurontin			
Initial or Prolonged		Dyspepsia		(Gabapentin)	PS		ORAL
3200 MG (FOUR							
Other		Gastritis Fungal					
TIMES A DAY),							
ORAL		Haemorrhage					
		Nausea					
		Tonsillectomy					

Date:04/27/04ISR Number: 4350737-8Report Type:Expedited (15-DaCompany Report #2004-DE-02030GD
Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Literature	Clonidine			
		Drug Tolerance Decreased		(Clonidine)	PS		
INTRATHECAL	150 MCG, IT	9 DAY					
		Drug Tolerance Increased		Paracetamol			
4G		Metastases To Spine		(Paracetamol)	SS		
		Motor Dysfunction		Amitriptyline			
25 MG		Nausea		(Amitriptyline)	SS		
		Refusal Of Treatment By		Gabapentin			
1800 MG		Patient		(Gabapentin)	SS		
		Somnolence		Dexamethasone			
4.5 MG		Vomiting		(Dexamethasone)	SS		
				Fentanyl (Fentanyl)	SS		
TRANSDERMAL	7.2 MG, TD			Morphine (Morphine)	SS		
INTRATHORACIC	SEE IMAGE	3 DAY					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

INTRATHECAL	60 MG, IT	27	DAY	Bupivacaine (Bupivacaine)	SS
INTRATHORACIC	20 MG, IT	3	DAY	Ketamine (Ketamine)	SS

Date:04/27/04ISR Number: 4350758-5Report Type:Expedited (15-DaCompany Report #2004025316
Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Other		Depression Intentional Misuse	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
				Carisoprodol (Carisoprodol)	C		
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		

Date:04/27/04ISR Number: 4350863-3Report Type:Expedited (15-DaCompany Report #2004025616
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cerebrovascular Accident Difficulty In Walking Fall Myocardial Infarction	Consumer	Neurontin (Gabapentin)	PS		
				Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
				Pravastatin Sodium(Pravastatin Sodium)	C		
				Labetalol (Labetalol)	C		
				Hydrochlorothiazide(Hydrochlorothiazide)	C		
				Clopidogrel Sulfate			

(Clopidogrel Sulfate) C

Date:04/27/04ISR Number: 4351011-6Report Type:Expedited (15-DaCompany Report #2004025785
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Hepatitis Cholestatic	Foreign Health Professional	Neurontin (Gabapentin) Hydromorphone Hydrochloride (Hydromorphone Hydrochloride) Tramadol/Acetaminophen (Paracetamol/Tramadol) Paracetamol (Paracetamol)	PS C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/27/04ISR Number: 4351714-3Report Type:Expedited (15-DaCompany Report #2004021921
Age:80 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Balance Disorder	Foreign	Neurontin			
Initial or Prolonged	Blood Sodium Decreased	Health	(Gabapentin)	PS		ORAL
300 MG						
	Dizziness	Professional				
(DAILY), ORAL						
	Fall	Company	Fentanyl (Fentanyl)	SS		
TRANSDERMAL	25 MCG/H					
		Representative				
(DAILY),						
TRANSDERMAL						
			Aporex			
			(Dextropropoxyphene			
			Hydrochloride,			
			Pracetamol)	C		
			Lisinopril			
			(Lisinopril)	C		
			Atenolol (Atenolol)	C		
			Omeprazole			
			(Omeprazole)	C		
			Temazepam			
			(Temazepam)	C		

Date:04/27/04ISR Number: 4351719-2Report Type:Expedited (15-DaCompany Report #2004009875
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Convulsion	Foreign	Neurontin			
	Drug Withdrawal Syndrome	Health	(Gabapentin)	PS		ORAL
ORAL						
		Professional				

Date:04/27/04ISR Number: 4351722-2Report Type:Expedited (15-DaCompany Report #2003125090
Age:60 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					

Hospitalization - Initial or Prolonged 1800 MG	Hepatotoxicity Transaminases Increased	Foreign Health	Neurontin (Gabapentin)	PS
(TID),		Professional		
		Company Representative	Antineoplastic Agents (Antineoplastic Agents)	SS

Date:04/28/04ISR Number: 4350396-4Report Type:Direct Company Report #CTU 217525
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG TWO		Drug Effect Decreased		Gabapentin (Novo)	PS	Novo	ORAL
TID PO		Pharmaceutical Product Complaint					

Date:04/28/04ISR Number: 4350821-9Report Type:Expedited (15-DaCompany Report #2004018498
 Age:44 YR Gender:Female I/FU:F

Outcome Other	PT Arthralgia Insomnia Mood Altered Recurrent Cancer
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tongue Neoplasm Malignant
Stage Unspecified

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
200 MG	(DAILY), ORAL	Health Professional	Zoloft (Sertraline)	PS		ORAL
1200 MG	(DAILY), ORAL		Neurontin (Tablets) (Gabapentin)	SS		ORAL
			Antiinflammatory/- Antirheumatic Products (Antiinflammatory-/A ntirheumatic	C		

Date:04/28/04ISR Number: 4350897-9Report Type:Expedited (15-DaCompany Report #2004026311
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	1800-2700MG (DAILY), ORAL	Abdominal Pain Upper Disease Recurrence Dyspepsia	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Food Poisoning Gastritis		All Other Non-Therapeutic Products (All Other Non-Therapeutic Products)	SS		ORAL
				Vitamins (Vitamins) Triamcinolone Acetonide Lorazepam All Other Therapeutic Products (All Other Products)	SS C C C		
				Lansoprazole Ranitidine Hydrochloride	C C C		

Date:04/28/04ISR Number: 4350901-8Report Type:Expedited (15-DaCompany Report #2004026316

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Amnesia	Consumer	Neurontin			
Initial or Prolonged	Burning Sensation		(Gabapentin)	PS		ORAL
900 MG (TID),						
Other	Condition Aggravated					
ORAL						
	Disturbance In Attention		Rofecoxib	C		
	Facial Pain					
	Headache					
	Hypoaesthesia					
	Neuropathy					
	Pruritus					
	Weight Decreased					

Date:04/28/04ISR Number: 4351000-1Report Type:Expedited (15-DaCompany Report #2004024723

Age:53 YR Gender:Male I/FU:I

Outcome	PT
Other	Amnesia
	Hepatitis C

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Freedom Of Information (FOI) Report

Speech Disorder

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1800 MG, (TID), ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
			Levetiracetam	C		
			Diazepam	C		
			Codeine Sulfate	C		

Date:04/28/04ISR Number: 4351305-4Report Type:Expedited (15-DaCompany Report #2004026318
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Colon Injury Diabetes Mellitus Difficulty In Walking Eczema Infection Loss Of Consciousness Nerve Injury Pain Paraesthesia	Consumer	Neurontin (Gabapentin) Celecoxib (Celecoxib) All Other Therapeutic Products (All Other Therapeutic Products)	PS SS C		

Date:04/28/04ISR Number: 4352420-1Report Type:Expedited (15-DaCompany Report #2004026305
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Dependence	Foreign Health	Neurontin (Gabapentin)	PS		
UNKNOWN	1600 (TID),		Professional				
UNKNOWN			Company Representative	Morphine Sulfate (Morphine Sulfate) Analesics (Analgesics) Anxiolytics	C C		

(Anxiolytics) C
Antidepressants
(Antidepressants) C

Date:04/29/04ISR Number: 4352749-7Report Type:Expedited (15-DaCompany Report #2012272

Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Consumer	Oxycontin Tablets			
Hospitalization -		Asthenia	Health	(Oxycodone			
Initial or Prolonged		Back Pain	Professional	Hydrochloride) Cr			
Other		Drug Abuser	Other	Tablet	PS		
SEE IMAGE		Erectile Dysfunction		Diazepam (Diazepam)	SS		
		Haematemesis		Oxazepam (Oxazepam)	SS		
		Headache		Temazepam			
		Iron Deficiency Anaemia		(Temazepam)	SS		
		Mallory-Weiss Syndrome		Cannabnoids(Cannabis			
		Tachycardia)	SS		
				Lorazepam	SS		
				Diphenhydramine			
				Hydrochloride			
				(Diphenhydramine			

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Hydrochloride)	SS
Gabapentin	
(Gabapentin)	SS
Claritin	
(Loratadine)	C
Zoloft (Sertraline	
Hydrochloride)	C
Altace (Ramipril)	C
Allopurinol	
(Allopurinol)	C
Depakote (Valproate	
Semisodium)	C
Soma (Carisoprodol)	C
Relafen (Nabumetone)	C

Date:04/29/04ISR Number: 4353142-3Report Type:Expedited (15-DaCompany Report #2004026162
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatitis Acute	Foreign Health	Neurontin (Gabapentin)	PS		
900 MG (TID)			Professional Company Representative	Aporex (Dextropropoxyphene Hydrochloride, Paracetamol)	C		
				Hydroxyzine Hydrochloride	C		
				Thiocolchicoside	C		

Date:04/29/04ISR Number: 4353143-5Report Type:Expedited (15-DaCompany Report #2004026081
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser Drug Dependence	Foreign Health	Neurontin (Gabapentin)	PS		
1200 (TID)		Drug Ineffective Psychopathic Personality	Professional Company Representative	Antipsychotics Anxiolytics	C C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haemorrhoids	Consumer	Neurontin			
ORAL		Neuralgia		(Gabapentin)	PS		ORAL
		Pharmaceutical Product		Baclofen (Baclofen)	C		
		Complaint		Diazepam (Diazepam)	C		
		Surgery		Amlodipine Besilate			
				(Amlodipine Besilate)	C		
				Simvastatin			
				(Simvastatin)	C		
				Oxybutynin			
				(Oxybutynin)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/29/04ISR Number: 4353207-6Report Type:Expedited (15-DaCompany Report #2004026285
Age:16 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 300 MG (DAILY		Abasia Asthenia	Consumer	Neurontin (Gabapentin)	PS		
		Balance Disorder Fall Feeling Abnormal Hypoaesthesia Nervous System Disorder Tremor					

Date:04/29/04ISR Number: 4353209-XReport Type:Expedited (15-DaCompany Report #2004005888
Age:69 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (THREE TIMES A DAY), ORAL		Abdominal Pain Abdominal Tenderness Abnormal Faeces Alopecia Anxiety Costovertebral Angle Tenderness Depression Diverticulitis Dysuria Ear Pain Eating Disorder Haematuria Insomnia Middle Ear Effusion Nausea Nitrite Urine Present Osteoarthritis Protein Urine Present Rectal Haemorrhage Sinusitis Urine Leukocyte Esterase	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Ketorolac Tromethamine (Ketorolac Tromethamine) Esomeprazole (Esomeprazole) Atorvastatin (Atorvastatin) Metronidazole (Metronidazole) Levothyroxine Sodium (Levothyroxine Sodium) Indapamide (Indapamide) Panadeine Co (Codeine Phosphate, Paracetamol)	C C C C C		

Positive

Zolpidem Tartrate
(Zolpidem Tartrate) C
Ultracet
(Paracetamol,
Tramadol
Hydrochloride) C
Alprazolam
(Alprazolam) C

Date:04/30/04ISR Number: 4351493-XReport Type:Direct
Age:90 YR Gender:Female I/FU:I

Company Report #CTU 217747

Outcome PT
Life-Threatening Acute Respiratory Failure
Hospitalization - Coma
Initial or Prolonged Dysarthria
Mucosal Inflammation
Neutrophil Count
Decreased
Pyrexia
White Blood Cell Count

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Decreased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
600 MG PO Q 8			Methadone	PS		
			Gabapentin	SS		ORAL
HRS			Lasix	C		
			Theo-Dur	C		
			Celexa	C		
			Nitropan Xl	C		
			Ultram	C		
			Aricept	C		

Date:04/30/04ISR Number: 4353384-7Report Type:Expedited (15-DaCompany Report #2004027040

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Calcium Increased	Consumer	Neurontin			
		Bone Formation Increased		(Gabapentin)	PS		ORAL
1800 MG, BID,		Delusion					
ORAL		Spinal Operation		Teriparatide			
				(Teriparatide)	SS		
DAILY				Anaesthetic			
				S(Anaesthetics)	SS		
				Morphine (Morphine)	SS		
				Enalapril			
				(Enalapril)	C		
				Nifedipine			
				(Nifedipine)	C		
				One-A-Day (Ascorbic			
				Acid,			
				Cyanocobalamin,			
				Ergocalciferol,			
				Nicotinamide,	C		
				Chlorphenamine			
				Maleate			
				(Chlorphenamine			
				Maleate)	C		

Calcium Magnesium
Zinc (Calcium,
Magnesium, Zinc) C

Date:04/30/04ISR Number: 4353471-3Report Type:Expedited (15-DaCompany Report #2004027304
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neutrophil Count Decreased	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL		White Blood Cell Count Decreased	Professional	Interferon Beta Citalopram Hydrobromide Ergocalciferol Calcium Risedronate Sodium Ranitidine Hydrochloride Paracetamol Carbamazepine	C C C C C C C C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/30/04ISR Number: 4353494-4Report Type:Expedited (15-DaCompany Report #2004026037

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Geodon (Ziprasidone)	PS		ORAL
Other		Eye Haemorrhage	Professional				
40 MG (BID),			Company Representative	Neurontin (Gabapentin)			
ORAL				(Gabapentin)	SS		ORAL
1800 MG							
(TID), ORAL				Zoloft (Sertraline)	SS		ORAL
50 MG							
(DAILY), ORAL							

Date:04/30/04ISR Number: 4353506-8Report Type:Expedited (15-DaCompany Report #2004021469

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other		Dysgeusia Ovarian Cancer	Professional				
3200 MG QID,			Pharmaceutical Product	Lisinopril (Lisinopril)	C		
ORAL			Complaint	Glibenclamide (Glibenclamide)	C		

Date:04/30/04ISR Number: 4353508-1Report Type:Expedited (15-DaCompany Report #2004027046

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Neurontin (Gabapentin)	PS		ORAL
Other		Drug Dependence Euphoric Mood	Professional				
1200 MG TID,							

Medication Error

ORAL

Clonazepam (Clonazepam)	C
Temazepam (Temazepam)	C
Pantoprazole (Pantoprazole)	C
Propranolol Hydrochloride (Propranolol Hydrochloride)	C

Date:04/30/04ISR Number: 4353606-2Report Type:Expedited (15-DaCompany Report #2004026961
Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dizziness Eyelid Oedema	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1200 MG			Professional				
DAILY, ORAL				Diazepam (Diazepam)	C		

Date:04/30/04ISR Number: 4353630-XReport Type:Expedited (15-DaCompany Report #2004021304
Age:59 YR Gender:Female I/FU:F

Outcome Other	PT Alanine Aminotransferase 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
900 (TID), ORAL		Aspartate Aminotransferase Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Blood Alkaline Phosphatase Increased	Professional				
		Cholelithiasis		Amitriptyline (Amitriptyline)	SS		
		Gamma-Glutamyltransferase Increased Hepatotoxicity		Tamoxifen (Tamoxifen)	C		
				Paracetamol (Paracetamol)	C		
				Insulin (Insulin)	C		
				Nifedipine (Nifedipine)	C		

Date:04/30/04ISR Number: 4353635-9Report Type:Expedited (15-DaCompany Report #2004021300
Age:58 YR Gender:Female I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (THREE TIMES A DAY), ORAL		Alanine Aminotransferase Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Aspartate Aminotransferase Increased	Professional				
		Blood Alkaline Phosphatase Increased		Ranitidine (Ranitidine)	SS		
		Gamma-Glutamyltransferase Increased Hepatotoxicity		Diazepam (Diazepam)	C		
				Dexamethasone (Dexamethasone)	C		

Date:04/30/04ISR Number: 4353637-2Report Type:Expedited (15-DaCompany Report #2004027303
Age:50 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization - Initial or Prolonged 200 MG (QID), Other ORAL	Neutropenic Sepsis	Foreign Health Professional	Gabapentin (Gabapentin)	PS	ORAL
			Antibiotics (Antibiotics)	C	
			Morphine Sulfate (Morphine Sulfate)	C	
			Trazodone Hydrochloride (Trazodone Hydrochloride)	C	
			Clonazepam (Clonazepam)	C	
			Pantoprazole (Pantoprazole)	C	
			All Other Therapeutic Products (All Other Therapeutic Products)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/30/04ISR Number: 4353651-7Report Type:Expedited (15-DaCompany Report #2004026614
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fall	Health	Neurontin			
6400 MG		Feeling Abnormal	Professional	(Gabapentin)	PS		ORAL
(QID), ORAL		Head Injury					
		Incoherent		Venlafaxien			
		Loss Of Consciousness		Hydrochloride			
		Panic Reaction		(Venlafaxine			
		Skin Laceration		Hydrochloride)	C		
		Urinary Retention					

Date:04/30/04ISR Number: 4353654-2Report Type:Expedited (15-DaCompany Report #2004027041
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin			
900 (TID),		Pharmaceutical Product		(Gabapentin)	PS		ORAL
ORAL		Complaint					
		Pruritus		Glimepiride			
				(Glimepiride)	C		
				Hydrocodone			
				(Hydrocodone)	C		
				Baclofen (Baclofen)	C		
				Metoprolol Succinate			
				(Metoprolol			
				Succinate)	C		
				Buspirone			
				Hydrochloride			
				(Buspirone			
				Hydrochloride)	C		
				Zaleplon (Zaleplon)	C		
				Rofecoxib			
				(Rofecoxib)	C		
				Lisinopril			
				(Lisinopril)	C		
				Rabeprazole Sodium			
				(Rabeprazole Sodium)	C		

Metformin
Hydrochloride
(Metformin
Hydrochloride) C
Pravastatin Sodium
(Pravastatin Sodium) C
Trazodone
(Trazodone) C

Date:04/30/04ISR Number: 4353679-7Report Type:Expedited (15-DaCompany Report #2002UW16093
Age:37 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Eructation
Other	Fatigue
	Headache
	Hot Flush
	Hypertension
	Medication Error
	Muscle Spasms

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nausea Oedema Overdose	Report Source				
TOOK 3-5 PER DAY		Vomiting	Consumer	Nexium	PS		
400 MG TID		Weight Increased		Neurontin	SS		
				Zocor	C		
				Lasix	C		
				Diovan	C		
				Protonix	C		
				Celexa	C		

Date:04/30/04ISR Number: 4353820-6Report Type:Expedited (15-DaCompany Report #2004026445
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agranulocytosis Leukopenia	Foreign Health	Neuronttin (Tablets) (Gabapentin)	PS		ORAL
900 MG			Professional				
(UNKNOWN), ORAL				Ponstan (Mefenamic Acid) (Mefenamic Acid)	SS		ORAL
1500 MG							
(UNKNOWN), ORAL				Cefepime Hydrochloride) (Cefepime Hydrochloride)	SS		
INTRAVENOUS	6 GRAM						
(UNKNOWN), INTRAVENOUS							

40 MG

(UNKNOWN),

ORAL

Omeprazole
(Omeprazole) SS

Oxycodone
Hydrochloride
(Oxycodone
Hydrochloride) C
Morphine C
Ketorolac
Tromethamine
(Ketorolac
Tromethamine) C
Co-Dafalgan (Codeine
Phosphate
Hemihydrate
Paracetamol) C
Droperidol
(Droperidol) C
Diclofenac Sodium
(Diclofenac Sodium) C
Amitriptyline
Hydrochloride
(Amitriptyline
Hydrochloride) C
Bromazepam
(Bromazepam) C
Clorazepate
Dipotassium

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Freedom Of Information (FOI) Report

(Clorazepate
Dipotassium) C
Vancomycin
Hydrochloride
(Vancomycin
Hydrochloride) C

Date:04/30/04ISR Number: 4353821-8Report Type:Expedited (15-DaCompany Report #2004016481
Age:59 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNKNOWN		Acute Psychosis Aggression	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
(UNKNOWN);		Confusional State	Professional				
ORAL		Drug Withdrawal Syndrome					
				All Other Therapeutic Products	C		

Date:05/03/04ISR Number: 4352542-5Report Type:Direct Company Report #CTU 217886
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG TID		Confusional State		Neurontin 600 Mg Tid	PS		
		Drug Interaction		Morphine Sr	SS		
				Clonazepam	C		
				Trazadone	C		
				Zoloft	C		
				Zyprexa	C		
				Kadian	C		
				Mobic	C		

Date:05/03/04ISR Number: 4354328-4Report Type:Expedited (15-DaCompany Report #2004026722
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Abdominal Pain Upper	Consumer	Neurontin (Tablets)		
2400 (TID),	Intervertebral Disc		(Gabapentin)	PS	ORAL
ORAL	Protrusion				
	Pharmaceutical Product		Levothyroxide Sodium		
	Complaint		(Levothyroxide		
			Sodium)	C	
			Etodolac (Etodolac)	C	
			Alendronate Sodium		
			(Alendronate Sodium)	C	
			Calcium (Calcium)	C	
			Sertraline		
			Hydrochloride		
			(Sertraline		
			Hydrochloride)	C	
			Lorazepam		
			(Lorazepam)	C	

Date:05/03/04ISR Number: 4354414-9Report Type:Expedited (15-DaCompany Report #2004027042
Age:47 YR Gender:Male I/FU:I

Outcome PT
Other Cerebral Haemorrhage
Headache

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nervous System Disorder

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
200 MG	(DAILY),	Health Professional	Neurontin (Gabapentin)	PS		

Date:05/03/04ISR Number: 4354423-XReport Type:Expedited (15-DaCompany Report #2004016094
Age:45 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	900 MG	(DAILY), ORAL	Retinal Detachment	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
					Olanzapine (Olanzapine)	C		

Date:05/03/04ISR Number: 4354435-6Report Type:Periodic Company Report #PHEH2003US11582
Age:69 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	5 MG		Angioneurotic Oedema	Health Professional Distributor	Lotrel (Amlodipine Besilate, Benazepril Hydrochloride) Capsule, 5/10 Mg	PS		
					Plavix /Unk/(Clopidogrel Sulfate)	SS		
					Neurontin /Unk/(Gabapentin)	SS		
					Trileptal			

(Oxcarbazepine,
 Oxcarbazapine)
 Unknown SS
 Glyburide
 (Glibenclamide) SS

Date:05/03/04ISR Number: 4354442-3Report Type:Expedited (15-DaCompany Report #2004003992
 Age:30 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG Other (DAILY), ORAL	Asthenia Muscle Spasms Muscle Twitching	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
	Pollakiuria Sensory Loss Urinary Incontinence	Company Representative	Nimesulide (Nimesulide) Paracetamol (Paracetamol)	C C		

Date:05/03/04ISR Number: 4354570-2Report Type:Expedited (15-DaCompany Report #2004-04-0933
 Age:45 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
Disability
Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Aphagia Aplastic Anaemia Bacterial Infection	Health Professional Company	Temodar (Temozolomide) Capsules	PS		ORAL
SEE IMAGE		Catheter Related	Representative	Neurontin Capsules	SS		ORAL
1200 MG TID		Infection					
ORAL		Conjunctival Haemorrhage		Septra Ds	SS		ORAL
ORAL		Ear Pain		Levaquin Tablets	SS		ORAL
X 7 DAYS ORAL	7 DAY	Headache		Thalidomide	SS		
1200 MG QD		Herpetic Stomatitis		Heparin Injectable	SS		
		Malaise		Famciclovir	C		
		Myalgia		Codeine	C		
		Petechiae		Phenobarbital	C		
		Pyrexia		Aspirin	C		
		Sepsis		Fragmin	C		
				Cft-11	C		
				Bcnu-70	C		
				Taxol	C		
				Tamiflu (Oseltamivir)	C		
				Ceftriaxone	C		

Date:05/04/04ISR Number: 4354946-3Report Type:Expedited (15-DaCompany Report #2003125945

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Dreams	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Anger					
300 MG (TID),		Body Height Decreased					
ORAL		Condition Aggravated		Clonidine (Clonidine)	C		
		Diarrhoea		Lorazepam			
		Drug Effect Decreased		(Lorazepam)	C		
		Eating Disorder					

Grand Mal Convulsion
 Hyperhidrosis
 Muscle Spasms
 Nausea
 Negative Thoughts
 Psychiatric Symptom
 Tremor
 Vomiting
 Weight Increased

Fosinopril Sodium
 (Fosinopril Sodium) C

Date:05/04/04ISR Number: 4355004-4Report Type:Expedited (15-DaCompany Report #2004024254
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Hyperkeratosis Scleroderma	Foreign Health Professional	Neurontin (Gabapentin) Metoprolol Succinate (Metoprolol Succinate) Calcium Acetate (Calcium Acetate) Sevelamer Hydrochloride (Sevelamer	PS C C		ORAL

Freedom Of Information (FOI) Report

Hydrochloride)	C
Alfacalcidol	
(Alfacalcidol)	C
Colecalciferol	
(Colecalciferol)	C
Madopar (Benserazide	
Hydrochloride,	
Levodopa)	C
Flunitrazepam	
(Flunitrazepam)	C
Phenprocoumon	
(Phenprocoumon)	C

Date:05/05/04ISR Number: 4355626-0Report Type:Expedited (15-DaCompany Report #2004024327
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Hypersensitivity	Foreign	Neurontin			
Initial or Prolonged	Rash	Health	(Gabapentin)	PS		
	Swelling	Professional	Carbamazepine			
	Testicular Swelling		(Carbamazepine)	SS		

Date:05/05/04ISR Number: 4355817-9Report Type:Expedited (15-DaCompany Report #2004027301
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Cognitive Disorder	Health	Neurontin			
Initial or Prolonged	Drug Interaction	Professional	(Gabapentin)	PS		
Other	Malaise		All Other			
			Therapeutic Products			
			(All Other			
			Therapeutic			
			Products)	SS		
			Amlodipine Besilate	C		
			Atorvastatin	C		
			Insulin Human			
			Injection, Isophane	C		
			Insulin Human	C		
			Warfarin	C		
			Furosemide	C		
			Potassium	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Grand Mal Convulsion	Health	Neurontin (Tablets)			
3200 MG		Laboratory Test Abnormal	Professional	(Gabapentin)	PS		ORAL
(THREE TIMES		Sedation					
A DAY), ORAL							
				Paxil (Paroxetine Hydrochloride)	C		
				Depakote (Valproate Semisodium)	C		
				Tylenol With Codeine (Codeine Phosphate, Paracetamol)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/05/04ISR Number: 4355822-2Report Type:Expedited (15-DaCompany Report #2004027664

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Operation Cardiac Pacemaker	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG DAILY		Insertion					
ORAL		Decreased Appetite Dementia Alzheimer'S Type Dizziness Insomnia		Donepezil Hydrochloride (Donepezil Hydrochloride)	SS		
OPHTHALMIC	5 MG DAILY						
OPHTHALMIC				Carbamazepine	C		

Date:05/05/04ISR Number: 4355823-4Report Type:Expedited (15-DaCompany Report #2004027834

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Cough Drug Level Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
2400 (TID)							
Other		Foreign Body Aspiration					
ORAL		Pneumonia Aspiration Urosepsis		Tolterodine L-Tartrate Lamotrigine Pantoprazole Amitriptyline Hydrochloride	C C C C		

Date:05/05/04ISR Number: 4355829-5Report Type:Expedited (15-DaCompany Report #2004027658

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Activities Of Daily	Consumer	Neurontin (Tablets)			

Other 2400 MG	Living Impaired	(Gabapentin)	PS	ORAL
(THREE TIMES A DAY), ORAL	Anxiety			
	Cognitive Disorder			
100 MG	Confusional State	Topiramate		
(DAILY), ORAL	Difficulty In Walking	(Topiramate)	SS	ORAL
	Disorientation			
	Fall	Hypnotics And		
	Insomnia	Sedatives (Hypnotics		
	Memory Impairment	And Sedatives)	SS	
	Mobility Decreased	Amlodipine Besilate		
	Sleep Walking	(Amlodipine		
	Treatment Noncompliance	Besilate)	C	
		Metformin		
		(Metformin)	C	
		Karvea Hct		
		(Hydrochlorothiazide		
		, Irbesartan)	C	
		Oxybutynin		
		(Oxybutynin)	C	
		Simvastatin		
		(Simvastatin)	C	
		Alendronate Sodium		
		(Alendronate Sodium)	C	
		Propafenone		
		(Propafenone)	C	
		Acetylsalicylic Acid		
		(Acetylsalicylic		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Acid) C

Date:05/05/04ISR Number: 4355830-1Report Type:Expedited (15-DaCompany Report #2004027685
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Anger	Consumer	Neurontin			
Other		Anxiety		(Gabapentin)	PS		ORAL
400 MG		Asthenia					
(DAILY), ORAL		Change In Sustained Attention					
		Cognitive Disorder					
		Confusional State					
		Depression					
		Drug Dependence					
		Drug Withdrawal Syndrome					
		Dysstasia					
		Fatigue					
		Feeling Abnormal					
		Hyperhidrosis					
		Hypotonia					
		Impaired Work Ability					
		Insomnia					
		Masked Facies					
		Nervous System Disorder					
		Peripheral Coldness					
		Weight Decreased					

Date:05/05/04ISR Number: 4355843-XReport Type:Expedited (15-DaCompany Report #2004011279
 Age:82 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abasia	Consumer	Neurontin			
ORAL		Amnesia	Health	(Gabapentin)	PS		ORAL
		Loss Of Consciousness	Professional	Magnesium Oxide			
		Speech Disorder		(Magnesium Oxide)	C		
				Digoxin (Digoxin)	C		
				Atorvastatin			
				(Atorvastatin)	C		

Gemfibrozil
(Gemfibrozil) C
Ramipril (Ramipril) C
Vitamins (Vitamins) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C
Insulin (Insulin) C

Date:05/06/04ISR Number: 4356283-XReport Type:Expedited (15-DaCompany Report #2002057832
Age:42 YR Gender:Male I/FU:F

Outcome	PT
Life-Threatening	Agranulocytosis
Hospitalization -	Bacterial Infection
Initial or Prolonged	Blood Lactate
Other	Dehydrogenase Increased
	Body Temperature
	Increased
	Bone Marrow Toxicity

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (TID), ORAL		Drug Toxicity Electrocardiogram St-T Segment Elevation Parvovirus Infection Pericardial Effusion Splenomegaly Staphylococcal Infection Troponin Increased Viral Myocarditis	Foreign Consumer Health Professional	Neurontin (Gabapentin) Cefotaxime Sodium (Cefotaxime Sodium) Cefalexin Monohydrate (Cefalexin Monohydrate)	PS C C		ORAL

Date:05/06/04ISR Number: 4356606-1Report Type:Expedited (15-DaCompany Report #2004028032
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 1200 MG Initial or Prolonged DAILY, ORAL Other		Alanine Aminotransferase Increased Aspartate Aminotransferase Increased Blood Alkaline Phosphatase Increased Blood Lactate Dehydrogenase Increased Circulatory Collapse Dizziness Gamma-Glutamyltransferase Increased Hepatocellular Damage Hepatotoxicity Nausea Restlessness	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Zinc	PS C		ORAL

Date:05/06/04ISR Number: 4357329-5Report Type:Expedited (15-DaCompany Report #2004027799
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Neurontin (Tablets)			
1800 (TID),		Infection	Professional	(Gabapentin)	PS		ORAL
ORAL		Pharmaceutical Product					
		Complaint		Gemfibrozil			
		Staring		(Gemfibrozil)	C		

Date:05/07/04ISR Number: 4356598-5Report Type:Expedited (15-DaCompany Report #200410158BFR
Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Eczema	Foreign	Glucor (Acarbose)	PS		ORAL
300 MG, TOTAL		Erythema	Health				
Initial or Prolonged		Fall	Professional	Furosemide	SS		ORAL
DAILY, ORAL		Pruritus	Other				
20 MG, TOTAL		Self-Medication		Neurontin			
DAILY, ORAL		Toxic Skin Eruption		(Gabapentin)	SS		ORAL
600 MG, TOTAL		Urticaria					
DAILY ,ORAL				Aprovel (Irbesartan)	SS		ORAL
300 MG TOTAL							
DAILY ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

150 MG BID	ORAL	Contramel (Tramadol Hydrochloride)	SS	ORAL
75 MG TOTAL	DAILY ORAL	Effexor (Venlafaxine Hydrochloride)	SS	ORAL
15 MG TOTAL	DAILY ORAL	Ogast (Lansoprazole)	SS	ORAL
400 MG TOTAL	DAILY ORAL	Difrarel	SS	ORAL
BID ORAL		Dafalgan (Paracetamol)	SS	ORAL
10 MG TOTAL	DAILY ORAL	Amlor (Amlodipine Besilate)	SS	ORAL
60 MG TOTAL	DAILY ORAL	Vastarel (Trimetazidine Hydrochloride)	SS	ORAL

Date:05/07/04ISR Number: 4356766-2Report Type:Expedited (15-DaCompany Report #2004028513

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dysgeusia	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1600 MG (4),		Complaint		Neurontin (Tablets)			
ORAL							

3200 MG (4),

(Gabapentin)

SS

ORAL

ORAL

Vicodin (Hydrocodone Bitartrate, Paracetamol)	C
Cetirizine Hydrochloride (Cetirizine Hydrochloride)	C
Lidocaine (Lidocaine)	C
Lactulose (Lactulose)	C
Salmeterol Xinafoate (Salmeterol Xinafoate)	C
Doxepin (Doxepin)	C
Salbutamol (Salbutamol)	C

Date:05/07/04ISR Number: 4356768-6Report Type:Expedited (15-DaCompany Report #2004028587

Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anaemia	Consumer	Neurontin (Gabapentin)	PS		ORAL
UNKNOWN		Faeces Discoloured					
Other (UNKNOWN),		Gastrointestinal					
		Haemorrhage					

ORAL

Simvastatin (Simvastatin)	C
Gliclazide	

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Freedom Of Information (FOI) Report

(Gliclazide) C
 Lansoprazole
 (Lansoprazole) C
 Latanoprost
 (Latanoprost) C
 Cosopt (Dorzolamide
 Hydrochloride,
 Timolol Maleate) C
 Amitriptyline
 Hydrochloride
 (Amitriptyline
 Hydrochloride) C

Date:05/07/04ISR Number: 4356774-1Report Type:Expedited (15-DaCompany Report #2004023410
 Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (BID), Other ORAL		Compression Fracture Convulsion Hyponatraemia Muscle Spasms Nausea Neuralgia Vomiting	Consumer Health Professional	Neurontin (Gabapentin) Rofecoxib (Rofecoxib) Ibuprofen (Ibuprofen) Lotrel (Amlodipine, Benazepril Hydrochloride) Alendronate Sodium (Alendronate Sodium) Provella-14 (Estrogens Conjugated, Medroxyprogesterone Acetate)	PS SS C C C C		ORAL

Date:05/07/04ISR Number: 4356776-5Report Type:Expedited (15-DaCompany Report #2004027642
 Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Test Abnormal	Health	Neurontin (Tablets)			

3200 MG
 (THREE TIMES
 A DAY), ORAL

Grand Mal Convulsion
 Pharmaceutical Product
 Complaint
 Sedation

Professional
 (Gabapentin)
 PS

Paxil (Paroxetine
 Hydrochloride)
 Depakote (Valproate
 Semisodium)
 Tylenol With Codeine
 (Codeine Phosphate,
 Paracetamol)

C
 C
 C

Date:05/10/04ISR Number: 4356571-7Report Type:Direct
 Age:38 YR Gender:Female I/FU:I

Company Report #CTU 218261

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Menorrhagia Menstruation Irregular		Neurontin 300 Mg Parke Davis	PS	Parke Davis	ORAL
300 MG ONCE A DAY ORAL				Rebiff	C		

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Freedom Of Information (FOI) Report

Motrin C
 Tylenol C
 Zoloft C

Date:05/11/04ISR Number: 4357749-9Report Type:Expedited (15-DaCompany Report #FRWYE593019FEB04
 Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other	37.5 MG 2X PER 1 DAY, ORAL	Bradyarrhythmia Cerebral Atrophy Eczema Electroencephalogram Abnormal	Health Professional Other	Effexor Lp (Venlafaxine Hydrochloride, Capsule, Extended Release, 0)	PS		ORAL
		Fall Headache					
	300 MG 1X PER 1 DAY, ORAL; LONG COURSE THERAPY	Metabolic Disorder Nervous System Disorder Self-Medication Toxic Skin Eruption Urticaria Vertigo		Aprovel (Irbesartan, , 0)	SS		ORAL
	2 TABLET 1X PER 1 DAY, ORAL; LONG COURSE THERAPY			Contramal (Tramadol Hydrochloride, , 0)	SS		ORAL
	100 MG 3X PER 1 DAY, ORAL;			Glucor (Acarbose, , 0)	SS		ORAL

LONG COURSE

THERAPY

Lasilix (Furosemide,
, 0) SS

ORAL

1 DOSE 1X PER

1 DAY, ORAL;

LONG COURSE

THERAPY

Neurontin
(Gabapentin, , 0) SS

ORAL

2 DOSE 1X PER

1 DAY, ORAL;

LONG COURSE

THERAPY

Amlor (Amlodipine
Besilate) C
Ogast (Lansoprazole) C
Difrarel
(Betacarotene/Myrtil
lus) C
Dafalgan
(Paracetamol) C
Vastarel
(Trimetazidine
Hydrochloride) C

Date:05/11/04ISR Number: 4358175-9Report Type:Expedited (15-DaCompany Report #2004011471

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

Outcome PT
Other Abnormal Behaviour
Brain Neoplasm Malignant

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Condition Aggravated Pharmaceutical Product Complaint	Report Source	Product	Role	Manufacturer	Route
		Simple Partial Seizures	Consumer	Neurontin (Tablets) (Gabapentin)	PS		
				Valproate Semisodium (Valproate Semisodium)	C		
				Desloratadine(Deslor atadine)	C		

Date:05/11/04ISR Number: 4358177-2Report Type:Expedited (15-DaCompany Report #2004029598
Age:65 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Condition Aggravated Oedema Peripheral	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG , TID), ORAL		Poor Peripheral Circulation	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Skin Discolouration		Paroxetine Hydrochloried (Paroxetine Hydrochloride)	C		
				Drug Used In Diabetes 9drug Used In Diabetes)	C		

Date:05/11/04ISR Number: 4358179-6Report Type:Expedited (15-DaCompany Report #2004015518
Age: Gender:Male I/FU:F

Outcome Dose Death Other	Duration	PT Bladder Disorder Chronic Obstructive Pulmonary Disease	Report Source	Product	Role	Manufacturer	Route
600 MG(DAILY),			Consumer Health	Neurontin (Gabapentin)	PS		ORAL
			Professional				

ORAL
 Condition Aggravated
 Disease Recurrence
 Dyspepsia
 Emphysema
 Gastric Disorder
 Lung Neoplasm Malignant
 Pain
 Pruritus

Diuretics
 (Diuretics) C
 All Otehr
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C

Date:05/11/04ISR Number: 4358204-2Report Type:Expedited (15-DaCompany Report #2004028897
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Effect Decreased Pharmaceutical Product	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG TID,		Complaint					

ORAL

Ibuprofen
 (Ibuprofen) C
 Budesonide
 (Budesonide) C
 Triamcinolone
 Acetonide
 (Triamcinolone
 Acetonide) C

Freedom Of Information (FOI) Report

Salmeterol Xinafoate
 (Salmeterol
 Xinafoate) C
 Narine Repetabs
 (Loratadine,
 Pseudoephedrine
 Sulfate) C
 Dyazide
 (Hydrochlorothiazide
 , Triamterene) C
 Cimetidine
 (Cimetidine) C
 Fluocinonide
 (Fluocinonide) C

Date:05/11/04ISR Number: 4358220-0Report Type:Expedited (15-DaCompany Report #2004029359
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Coordination Abnormal Drug Withdrawal Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Dyskinesia		Alprazolam (Alprazolam)	SS		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		

Date:05/11/04ISR Number: 4358267-4Report Type:Expedited (15-DaCompany Report #2004028419
 Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (TID), ORAL		Arthralgia Back Pain Neck Pain Pain In Extremity	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Propranolol Hydrochloride	C		
				Fluoxetine Hydrochloride	C		

Date:05/11/04ISR Number: 4358333-3Report Type:Expedited (15-DaCompany Report #2004028507

Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Neurontin (Gabapentin)	PS		
1800 MG			Company Representative	Methadone (Methadone)	SS		
40 MG (10 MG,							
4 IN 1 D)							

Date:05/11/04ISR Number: 4358335-7Report Type:Expedited (15-DaCompany Report #2003121603

Age: Gender:Male I/FU:F

Outcome	PT
Other	Abasia Aphasia Arthralgia Bone Pain

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FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	Confusional State Convulsion Decreased Activity	Report Source	Product	Role	Manufacturer	Route
1600 MG	(QID), ORAL	Depression Diarrhoea	Consumer	Dilantin Suspension (Phenytoin Sodium)	PS		ORAL
1200 MG	(QID), ORAL	Drug Ineffective Drug Toxicity Dysphemia		Neurontin (Gabapentin)	SS		ORAL
		Epilepsy Grand Mal Convulsion Humerus Fracture Loss Of Employment Memory Impairment Myalgia Personality Disorder Petit Mal Epilepsy Rotator Cuff Syndrome Tremor Weight Increased		Antiepileptics (Antiepileptics) Escitalopram (Escitalopram) Cannabis (Cannabis)	SS C C		

Date:05/11/04ISR Number: 4358457-0Report Type:Expedited (15-DaCompany Report #2004019964
Age:80 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	3600 MG	Cholestasis Hepatocellular Damage	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other (DAILY), ORAL		Nausea Vomiting	Professional				

Date:05/11/04ISR Number: 4358700-8Report Type:Expedited (15-DaCompany Report #2004028910
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Suicide Attempt	Health	Neurontin			

Initial or Prolonged
900 MG TID
Other

Professional

(Gabapentin)

PS

Quetiapine Fumarate
(Quetiapine
Fumarate)
Trazodone
(Trazodone)
Venlafaxine
Hydrochloride
(Venlafaxine
Hydrochloride)
Paracetamol
(Paracetamol)

C
C
C
C

Date:05/12/04ISR Number: 4358830-0Report Type:Expedited (15-DaCompany Report #2004019973
Age:30 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 800 MG Other (DAILY), ORAL	Convulsion Coordination Abnormal Fall Head Injury Hypertension	Foreign Consumer	Gabapentin (Tablets) (Gabapentin) Phenytoin (Phenytoin) Clonazepam (Clonazepam)	PS C C		ORAL

Freedom Of Information (FOI) Report

Ginkgo Biloba
(Ginkgo Biloba) C

Date:05/13/04ISR Number: 4360035-4Report Type:Expedited (15-DaCompany Report #2004029925

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1400 MG Other (EVERY DAY), ORAL		Abnormal Behaviour Aggression Anxiety Convulsion Depersonalisation Dissociation Head Banging Head Injury Loss Of Consciousness Medication Error Overdose Personality Change Psychomotor Hyperactivity Sexual Assault Victim	Consumer	Neurontin (Gabapentin) Gabapentin (Gabapentin) Carisoprodol (Carisoprodol) Sertraline Hydrochloride (Sertraline Hydrochloride) Obetrol Amfetamine Aspartate, Amfetamine Sulfate, Dexamfetamine Saccharate,	PS SS SS C C		ORAL

Date:05/13/04ISR Number: 4360037-8Report Type:Expedited (15-DaCompany Report #2004029813

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 4500 MG (300 MG, TID), ORAL		Dizziness Drug Effect Decreased Fall Gait Disturbance Gout Insomnia	Consumer	Neurontin (Gabapentin) Torasemide (Torasemide)	PS C		ORAL

Neuropathic Pain

Clonidine	
(Clonidine)	C
Lansoprazole	
(Lansoprazole)	C
Paroxetine	
Hydrochloride	
(Paroxetine	
Hydrochloride)	C
Allopurinol	
(Allopurinol)	C
Ezetimibe	
(Ezetimibe)	C
Candesartan	
Cilexetil	
(Candesartan	
Cilexetil)	C

Date:05/13/04ISR Number: 4360217-1Report Type:Expedited (15-DaCompany Report #2004016191

Age: 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

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, TID), ORAL		Abdominal Pain Agitation Anxiety Crying Diabetes Mellitus Economic Problem Fatigue Hepatic Cirrhosis Hepatic Steatosis Hepatomegaly Inappropriate Affect Intestinal Obstruction Irritability Liver Induration Tremor	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Diltiazem Hydrochloride (Diltiazem Hydrochloride)	SS		
				Benazepil Hydrochloride (Benazepil Hydrochloride)	C		
				Hydrochlorothiazide (Hydrochlorothiazide)	C		

Date:05/13/04ISR Number: 4360512-6Report Type:Expedited (15-DaCompany Report #2004-DE-02030GD
Age:77 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death INTRATHECAL Other 4 GM, NR 25MG 1800 MG 4.5 MG TRANSDERMAL INTRATHECAL INTRATHECAL		150 MCG, IT	Drug Ineffective Metastases To Spine Refusal Of Treatment By Patient	Foreign Literature Health Professional Other	Clonidine Paracetamol Amitriptyline Gabapentin Dexamethasone Fentanyl Morphine Bupivacaine	PS SS SS SS SS SS SS SS		
		7.2 MG						
		10MG; 20MG	3 DAY					
		60MG	27 DAY					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Consumer	Oxycodone			
		Cardiomegaly	Health	Hydrochloride			
		Chronic Obstructive	Professional	(Similar To Nda			
		Pulmonary Disease	Other	20-553)	PS		
		Coma		Hydrocodone			
		Drug Toxicity		Bitartrate (Similar			
		Fall		To Ind 59,175)	SS		
				Fluoxetine			
				(Fluoxetine)	SS		
				Caffeine (Caffeine)	SS		
				Alprazolam			
				(Alprazolam)	SS		
				Citalopram			
				(Citalopram)	SS		
				Metoclopramide			
				(Metoclopramide)	SS		
				Nicotine (Nicotine)	SS		
				Gabapentin			
				(Gabapentin)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/14/04ISR Number: 4359824-1Report Type:Expedited (15-DaCompany Report #2004030444

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Dilantin Suspension			
400 MG (200		Drug Level Below	Professional	(Phenytoin Sodium)	PS		
MG, BID),		Therapeutic					
UNKNOWN							
				Neurontin			
3600 MG (900				(Gabapentin)	SS		
MG, QID),							
UNKNOWN							
				Benzatropine			
				Mesilate	C		
				Lorazepam	C		
				Thioridazine	C		
				Diazepam	C		

Date:05/14/04ISR Number: 4359825-3Report Type:Expedited (15-DaCompany Report #2004029727

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abdominal Distension	Health	Neurontin			
Initial or Prolonged		Abdominal Pain	Professional	(Gabapentin)	PS		
Disability		Abdominal Pain Upper		Insulin (Insulin)	C		
Other		Activities Of Daily		Insulin Glargine			
		Living Impaired		(Insulin Glargine)	C		
		Acute Stress Disorder					
		Ascites					
		Crying					
		Diabetes Mellitus					
		Inadequate Control					
		Dyspnoea					
		Dysstasia					
		Emotional Disorder					
		Fatigue					
		Hepatic Cirrhosis					

Hepatic Steatosis
Herpes Zoster
Homicidal Ideation
Impaired Work Ability
Intestinal Obstruction
Movement Disorder
Nervousness
Postoperative Adhesion
Simple Partial Seizures
Visual Disturbance

Date:05/14/04ISR Number: 4359827-7Report Type:Expedited (15-DaCompany Report #2004029944
Age: Gender:Female I/FU:I

Outcome PT
Other Activities Of Daily
Living Impaired
Catatonia
Dry Mouth
Dysphonia
Eye Rolling
Impaired Driving Ability
Memory Impairment

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Somnolence

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
(THREE TIMES A DAY), ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
			Flexeril (Cyclobenzaprine Hydrochloride)	C		

Date:05/14/04ISR Number: 4360417-0Report Type:Expedited (15-DaCompany Report #2004030577
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 4800 MG (1600 MG, THREE TIMES A DAY), ORAL		Blindness Transient Somnolence Tinnitus Visual Disturbance	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Tolterodine L-Tartrate (Tolterodine L-Tartrate)	C		
				Bactrim (Sulfamethoxazole, Trimethoprim)	C		

Date:05/14/04ISR Number: 4360418-2Report Type:Expedited (15-DaCompany Report #2004029933
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other (100 MG,		Asthenia Difficulty In Walking	Consumer	Neurontin (Gabapentin)	PS		ORAL

DAILY), ORAL	Dysphagia			
20 MG (20 MG,	Feeling Abnormal	Bextra (Valdecoxib)	SS	ORAL
ONCE DAILY),	Gait Disturbance			
ORAL	Mobility Decreased			
	Muscle Disorder	Metoprolol Tartrate		
	Nervous System Disorder	(Metoprolol		
	Rash	Tartrate)	C	
	Somnolence	Isosorbide		
		Mononitrate		
		(Isosorbide		
		Mononitrate)	C	
		Fenofibrate		
		(Fenofibrate)	C	
		Ezetimibe		
		(Ezetimibe)	C	
		Pravastatin Sodium		
		(Pravastatin Sodium)	C	

Date:05/14/04ISR Number: 4360438-8Report Type:Expedited (15-DaCompany Report #2004025366
Age: Gender:Female I/FU:F

Outcome PT
Other Amnesia
Difficulty In Walking
Drug Ineffective

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dyspnoea Feeling Abnormal Neck Pain	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1500 MG (FOUR TIMES A DAY); ORAL		Neuralgia Palpitations Posture Abnormal	Professional	Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		

Date:05/14/04ISR Number: 4360440-6Report Type:Expedited (15-DaCompany Report #2004-DE-02030GD
Age:77 YR Gender:Female I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other INTRATHECAL	150 MCG , IT 9 DAY	Drug Ineffective Refusal Of Treatment By Patient	Foreign Literature Health Professional Other	Clonidine (Clonidine) Paracetamol (Paracetamol) Amitriptyline (Amitriptyline)	PS SS SS		
25 MG				Gabapentin (Gabapentin)	SS		
1800 MG				Dexamethasone (Dexamethasone)	SS		
4.5 MG				Fentanyl (Fentanyl)	SS		
TRANSDERMAL	7/2 MG, TD			Morphine (Morphine)	SS		
INTRATHORACIC	10 MG, IT			Bupivacaine (Bupivacaine)	SS		
INTRATRACHEAL	60 MG, IT 27 DAY			Ketamine (Ketamine)	SS		
INTRATHECAL	20 MG, IT 3 DAY						

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 600MG, TID, Initial or Prolonged BY MOUTH	Confusional State Dizziness Lethargy		Gabapentin Dipyridamole Calcium Carbonate Aspirin (Enteric Coated) Docusate Sodium (Colace) Isosorbide Dinitrate Nortriptyline Hcl Vitamin B Complex Flunisolide (Aerobid) Atenolol Insulin , Human Regular (Novolin-R) Diltiazem	PS C C C C C C C C C C C C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/17/04ISR Number: 4361378-0Report Type:Expedited (15-DaCompany Report #2004024327

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Hypersensitivity	Foreign	Neurontin			
Initial or Prolonged	Rash	Health	(Gabapentin)	PS		
	Swelling	Professional	Carbamazepine			
	Testicular Swelling		(Carbamazepine)	SS		

Date:05/17/04ISR Number: 4361397-4Report Type:Expedited (15-DaCompany Report #2004029398

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Aplastic Anaemia	Health	Neurontin			
Initial or Prolonged	Bacterial Infection	Professional	(Gabapentin)	PS		ORAL
3600 MG						
Other	Bone Marrow Depression					
(TID), ORAL						
	Bronchitis		Temozolomide			
	Catheter Related		(Temozolomide)	SS		ORAL
80 MG (QD),						
	Infection					
ORAL						
	Catheter Site Pain		Bactrim			
	Conjunctival Haemorrhage		(Sulfamethoxazole,			
	Cough		Trimethoprim)	SS		ORAL
ORAL						
	Ear Pain		Levofloxacin			
	Fluid Intake Reduced		(Levofloxacin)	SS		ORAL
ORAL						
	Headache		Thalidomide			
	Herpetic Stomatitis		(Thalidomide)	SS		
1200 MG (QD)						
	Malaise		Heparin (Heparin)	SS		
INTRAVENOUS	INTRAVENOUS					
	Myalgia		Famciclovir			
	Oral Intake Reduced		(Famciclovir)	C		
	Oral Mucosal Petechiae		Codeine (Codeine)	C		
	Pain		Phenobarbital			
	Petechiae		(Phenobarbital)	C		
	Sepsis		Acetylsalicylic			
	Upper Respiratory Tract		Acid			
	Infection		(Acetylsalicylic			

Acid)	C
Heparin-Fraction, Sodium Salt	
(Heparin-Fraction, Sodium Salt)	C
Carmustine	
(Carmustine)	C
Paclitaxel	
(Paclitaxel)	C
Irinotecan	
(Irinotecan)	C
Ceftriaxone	
(Ceftriaxone)	C

Date:05/18/04ISR Number: 4362409-4Report Type:Expedited (15-DaCompany Report #2004030834
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
ORAL				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Tocopherol (Tocopherol)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Biotin (Biotin) C
 Esomeprazole
 (Esomeprazole) C

Date:05/18/04ISR Number: 4362696-2Report Type:Expedited (15-DaCompany Report #2004028507
 Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800 MG (3 IN 1 D)	Death	Health Professional Company	Neurontin (Gabapentin)	PS		
	40 MG (10 MG, 4 IN 1 D)		Representative	Methadone (Methadone)	SS		

Date:05/18/04ISR Number: 4362751-7Report Type:Expedited (15-DaCompany Report #2004030542
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly Other TRANSPLACENTAL	600 MG (300 MG, TWICE A DAY), PLACENTAL	Developmental Coordination Disorder Maternal Drugs Affecting Foetus Myoclonic Epilepsy	Consumer	Neurontin (Gabapentin)	PS		
		Nuclear Magnetic Resonance Imaging Brain Abnormal Pregnancy		Montelukast Sodium (Montelukast Sodium)	C		

Date:05/18/04ISR Number: 4362965-6Report Type:Expedited (15-DaCompany Report #2004030937
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased Carpal Tunnel Syndrome Headache	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Cephalosporins And Related Substances (Cephalosporins And Related Substances) Nimesulide (Nimesulide) Antihypertensives (Antihypertensives)	PS C C C		

Date:05/18/04ISR Number: 4362968-1Report Type:Expedited (15-DaCompany Report #2004031169
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged DAILY		Insomnia Muscle Spasms Neuropathic Pain	Foreign Consumer	Neurontin (Gabapentin)	PS		
INTERVAL				Insulin (Insulin) Fluoxetine (Fluoxetine) Levothyroxine (Levothyroxine)	C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/19/04ISR Number: 4363110-3Report Type:Expedited (15-DaCompany Report #2004031170
 Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG , 3 IN 1 D), ORAL		Hepatic Function Abnormal Pruritus Generalised Rash Maculo-Papular	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG (200 MG, 3 IN 1 D), ORAL				Carbamazepine (Carbamazepine)	SS		ORAL
				Bendroflumethiazide (Bendroflumethiazide) Amitriptyline (Amitriptyline) Ramipril (Ramipril) Tramadol (Tramadol)	C C C C		

Date:05/19/04ISR Number: 4364212-8Report Type:Expedited (15-DaCompany Report #2004032207
 Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Completed Suicide Injury Self Injurious Behaviour	Consumer	Neurontin (Gabapentin)	PS		

Date:05/19/04ISR Number: 4364217-7Report Type:Expedited (15-DaCompany Report #2004030444
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Affective Disorder	Health	Dilantin Suspension			

Life-Threatening 400 MG (200 Other MG BID)	Anticonvulsant Drug Level Decreased Convulsion	Professional	(Phenytoin Sodium)	PS
3600 MG (900 MG, QID)			Neurontin (Gabapentin)	SS
			Benzatropine Mesilate	C
			Lorazepam	C
			Thioridazine	C
			Diazepam	C

Date:05/19/04ISR Number: 4364483-8Report Type:Expedited (15-DaCompany Report #04P-163-0260284-00
Age:55 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Abdominal Pain Upper
Initial or Prolonged	Cataract
Required	Cellulitis
Intervention to	Condition Aggravated
Prevent Permanent	Decreased Appetite
Impairment/Damage	Depressed Mood
	Depression
	Diarrhoea
	Disturbance In Attention
	Drug Abuser
	Feeling Of Despair

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration		Report Source	Product	Role	Manufacturer	Route
		Gait Disturbance Hallucination, Auditory Hallucination, Visual					
		Influenza Like Illness Insomnia Myalgia Nausea Neuropathic Pain Nightmare Obsessive Thoughts	Consumer Health Professional Other	Vicodin Es (Hydrocodone/Acetaminophen) (Hydrocodone/Acetaminophen) Peginterferon Alfa-2b	PS SS		
SUBCUTANEOUS	150 MCG, 1 IN	Pancytopenia					
1 WK,		Pollakiuria					
SUBCUTANEOUS;		Sedation					
SEE IMAGE		Suicidal Ideation		Ribavirin	SS		ORAL
1200 MG, 1 IN		Toothache					
1 D, ORAL;		Vomiting					
SEE IMAGE		Weight Decreased		Gabapentin	SS		
300 MG, 3 IN							
1 D; SEE							
IMAGE				Rofecoxib	C		
				Lexapro	C		
				Omeprazole	C		
				Ondansetron			
				Hydrochloride	C		
				Prochlorperazine			
				Edisylate	C		
				Buspirone			
				Hydrochloride	C		
				Multivitamins	C		
				Nystatin	C		
				Lidocaine	C		
				Filgrastim	C		
				Erythropoietin	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Neurontin Cap 300 Mg	PS		ORAL
ONE PO TID		Pharmaceutical Product Complaint					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abnormal Behaviour	Consumer	Neurontin			
Disability		Anhedonia		(Gabapentin)	PS		
Other		Emotional Disorder Mental Disorder Pain Pharmaceutical Product Complaint Polytraumatism					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/20/04ISR Number: 4365319-1Report Type:Expedited (15-DaCompany Report #2004031206
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 800 MG (400 MG, 2 IN 1 D), ORAL		Ageusia Anosmia Dizziness Laryngitis Nasopharyngitis	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Metformin (Metformin)	C		
				Glimepiride (Glimepiride)	C		
				Celecoxib (Celecoxib)	C		
				Pravastatin Sodium (Pravastatin Sodium)	C		
				Ibuprofen (Ibuprofen)	C		
				Omeprazole (Omeprazole)	C		

Date:05/20/04ISR Number: 4365320-8Report Type:Expedited (15-DaCompany Report #2004026316
Age:63 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, 3 IN 1 D), ORAL		Amnesia Condition Aggravated Disturbance In Attention Drug Ineffective Facial Pain Fear Headache Hypoaesthesia Neuropathy Peripheral Pruritus Vision Blurred Weight Decreased	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
			Professional				
				Rofecoxib (Rofecoxib)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Atrial Fibrillation	Consumer	Neurontin (Tablets)			
Other		Cardiac Operation		(Gabapentin)	PS		ORAL
2400 MG (800		Cardiac Pacemaker					
MG, 3 IN 1		Insertion					
D), ORAL		Decreased Appetite		Donepezil			
		Dementia Alzheimer'S Type		Hydrochloride			
		Diarrhoea		(Donepezil			
		Hypoglycaemia		Hydrochloride)	SS		ORAL
10 MG (10 MG,		Insomnia					
1 IN 1 D),		Myocardial Infarction					
ORAL		Tricuspid Valve		Carbamazepine			
		Incompetence		(Carbamazepine)	C		
				Warfarin Sodium			
				(Warfarin Sodium)	C		
				Amiodarone			
				Hydrochloride			
				(Amiodarone			
				Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Digoxin (Digoxin)	C
Furosemide (Furosemide)	C
Isosorbide Mononitrate (Isosorbide Mononitrate)	C
Potassium Chloride (Potassium Chloride)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C
Centrum (Minerals Nos, Vitamins Nos)	C
Atorvastatin (Atorvastatin)	C
All Other Therapeutic Products (All Other Therapeutic Products)	C
Magnesium (Magnesium)	C
Zinc (Zinc)	C
Metoprolol (Metoprolol)	C
Enalapril Maleate (Enalapril Maleate)	C

Date:05/20/04ISR Number: 4365721-8Report Type:Expedited (15-DaCompany Report #2004032003
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident	Foreign	Neurontin			
Other		Femur Fracture	Consumer	(Gabapentin)	PS		ORAL
150 MG		Medication Error					
(DAILY), ORAL							

Date:05/21/04ISR Number: 4364598-4Report Type:Expedited (15-DaCompany Report #2004025614
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blindness Nausea	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
3200 MG (800, QID), ORAL		Visual Acuity Reduced	Professional				
50 MG (DAILY), ORAL		Vomiting		Zolmitriptan (Zolmitriptan)	SS		ORAL
				Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
				Atenolol (Atenolol)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Simvastatin (Simvastatin)	C		
				Celecoxib (Celecoxib)	C		
				Metoclopramide			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Metoclopramide) C
 Levothyroxine Sodium
 (Levothyroxine Sodium) C

Date:05/21/04ISR Number: 4364611-4Report Type:Expedited (15-DaCompany Report #2004033113
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaphylactic Reaction	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:05/21/04ISR Number: 4364612-6Report Type:Expedited (15-DaCompany Report #2004019177
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Aspiration Pneumonia	Consumer	Neurontin (Gabapentin)	PS		
Other		Relapsing Polychondritis					

Date:05/21/04ISR Number: 4364946-5Report Type:Expedited (15-DaCompany Report #2004031827
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Aggression Agitation	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG							
Other (DAILY) ORAL		Anger					
		Fatigue Oral Intake Reduced Stress Suicide Attempt Treatment Noncompliance Weight Decreased		Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		

Date:05/21/04ISR Number: 4364975-1Report Type:Expedited (15-DaCompany Report #2004031739
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cystic Fibrosis Lung	Foreign	Gabapentin			
Other		Neoplasm Malignant	Health	(Gabapentin)	PS		ORAL
1800 MG (300		Renal Failure Acute	Professional				
MG) ORAL				Gliclazide			
				(Gliclazide)	C		

Date:05/21/04ISR Number: 4364992-1Report Type:Expedited (15-DaCompany Report #2004001236
Age:80 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Asthenia
Initial or Prolonged	Bacterial Infection
Other	Cardiomegaly
	Chlamydia Serology
	Positive
	Diastolic Dysfunction
	Headache

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Freedom Of Information (FOI) Report

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Iatrogenic Injury Inflammation Interstitial Lung Disease			Report Source				
Pulmonary Embolism Pulmonary Fibrosis			Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Viral Infection			Professional				
				Dihydroergotoxine (Dihydroergotoxine)	SS		ORAL
				Alendroante Sodium (Alendronate Sodium)	SS		ORAL

Date:05/21/04ISR Number: 4365023-XReport Type:Expedited (15-DaCompany Report #2004014131
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other MG, 3 IN 1			Professional				
				Amitriptyline (Amitriptyline)	C		
				Medroxyprogesterone Acetate (Medroxyprogesterone Acetate)	C		
				Fusidic Cid (Fusidic Acid)	C		

Date:05/21/04ISR Number: 4365025-3Report Type:Expedited (15-DaCompany Report #2004031968
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Foreign	Neurontin			

Initial or Prolonged 300 MG, (300 MG, 1 IN 1 D)	Hypersensitivity Productive Cough Rash Skin Discolouration Upper Respiratory Tract Infection Vomiting	Consumer	(Gabapentin)	PS
			Amitriptyline Hydrochloride (Amitriptyline Hydrochloride) Bi Predonium (Indapamide, Perindopril Erbumine) Estrogens Conjugated (Estrogens Conjugated) Diazepam (Diazepam)	C C C C

Date:05/24/04ISR Number: 4366185-0Report Type:Expedited (15-DaCompany Report #2004016481
Age:59 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Acute Psychosis Aggression Confusional State Disorientation Drug Withdrawal Syndrome Restlessness	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/24/04ISR Number: 4366187-4Report Type:Expedited (15-DaCompany Report #2004032454

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Toxic Skin Eruption	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Antiepileptics (Antiepileptics)	PS C		

Date:05/24/04ISR Number: 4366643-9Report Type:Expedited (15-DaCompany Report #2004014815

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG (DAILY), ORAL	Dissociation Loss Of Consciousness Medication Error	Foreign Health Professional	Neurontin (Gabapentin) All Other Therapeutic Products (All Other Therapeutic Products) Valproate Sodium	PS SS C		ORAL

Date:05/24/04ISR Number: 4366959-6Report Type:Expedited (15-DaCompany Report #2004019177

Age:42 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other	Pneumonia Aspiration Relapsing Polychondritis	Consumer	Neurontin (Gabapentin)	PS		

Date:05/24/04ISR Number: 4366971-7Report Type:Expedited (15-DaCompany Report #2004030444

Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anticonvulsant Drug Level	Health	Dilantin Suspension			
Life-Threatening		Decreased	Professional	(Phenytoin Sodium)	PS		
Other		Convulsion					
	400 MG (200 MG, BID)			Neurontin (Gabapentin)	SS		
	3600 MG, (900 MG.QID)			Benzatropine Mesilate (Benzatropine Mesilate)	C		
				Lorazepam (Lorazepam)	C		
				Thioridazine (Thioridazne)	C		
				Diazepam (Diazepam)	C		

Date:05/25/04ISR Number: 4365375-0Report Type:Expedited (15-DaCompany Report #2004032264
Age:60 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Other	Ageusia Anosmia	Foreign Health

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
ORAL		Gabapentin (Gabapentin)	PS		ORAL

Date:05/25/04ISR Number: 4368942-3Report Type:Expedited (15-DaCompany Report #2004032194
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Depression Multiple Sclerosis	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL		Pharmaceutical Product Complaint					

Interferon Beta (Interferon Beta)	SS
Tamsulosin Hydrochloride (Tamsulosin Hydrochloride)	C
Oxybutynin Hydrochloride (Oxybutynin Hydrochloride)	C

Date:05/25/04ISR Number: 4368971-XReport Type:Expedited (15-DaCompany Report #2004032769
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Delirium Eating Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
1600 MG (400 MG, QID INTERVAL:		Pain In Extremity Staphylococcal Infection					

Weight Decreased

EVERY DAY),

ORAL

Metformin	
Hydrochloride	C
Insulin	C
Ramipril	C
Potassium Chloride	C
Multivitamins	
(Ascorbic Acid,	
Ergocalciferol,	
Folic Acid,	
Nicotinamide,	C
Calcium	C
Furosemide	C

Date:05/25/04ISR Number: 4368980-0Report Type:Expedited (15-DaCompany Report #2004032770

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Anxiety	Consumer	Neurontin			
Initial or Prolonged	Drug Ineffective		(Gabapentin)	PS		ORAL
ORAL						
Other	Drug Withdrawal Syndrome		Ibuprofen	C		
	Mental Disorder					
	Tremor					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/26/04ISR Number: 4365631-6Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 219580

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hypoxia Respiratory Disorder		Gabapentin 300-600 Mg Park-Davis -Pfizer	PS	Park-Davis-Pfizer	ORAL
600-900 MG							
TID-QID ORAL				Oxycontin	C		
				Decadron	C		
				Vioxx	C		

Date:05/26/04ISR Number: 4366166-7Report Type:Direct
 Age:69 YR Gender:Female I/FU:I

Company Report #CTU 219493

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Sedation		Gabapentin Simvastatin	PS C		

Date:05/26/04ISR Number: 4367202-4Report Type:Expedited (15-DaCompany Report #2012272
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Consumer	Oxycontin Tablets			
Hospitalization - Initial or Prolonged		Asthenia Back Pain	Health Professional	(Oxycodone Hydrochloride) Cr			
Other		Drug Abuser	Other	Tablet	PS		
SEE IMAGE Required		Drug Level Below		Diazepam (Diazepam)	SS		
Intervention to Prevent Permanent Impairment/Damage		Therapeutic Drug Toxicity		Oxazepam (Oxazepam) Temazepam	SS		
		Erectile Dysfunction		(Temazepam)	SS		
		Haematuria		Lorazepam			
		Headache		(Lorazepam)	SS		
		Iron Deficiency Anaemia		Cannabnoids			
		Mallory-Weiss Syndrome		(Cannabis)	SS		
		Neck Pain		Diphenhydramine			
		Respiratory Arrest		Hydrochloride			

Tachycardia
Upper Gastrointestinal
Haemorrhage

(Diphenhydramine
Hydrochloride) SS
Gabapentin
(Gabapentin) SS
Claritin
(Loratadine) C
Zoloft (Sertraline
Hydrochloride) C
Altace (Ramipril) C
Allopurinol
(Allopurinol) C
Depakote (Valproate
Semisodium) C
Soma (Carisoprodol) C
Relafen (Nabumetone) C

Date:05/26/04ISR Number: 4367562-4Report Type:Expedited (15-DaCompany Report #2004032696
Age:48 YR Gender:Female I/FU:I

Outcome PT
Disability Asthenia
Other Coordination 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
900 MG (300 MG, 3 IN 1 D), ORAL		Dizziness Impaired Driving Ability	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:05/26/04ISR Number: 4367565-XReport Type:Expedited (15-DaCompany Report #2004033139
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Bedridden Full Blood Count Decreased Hypotension Mobility Decreased Pneumonitis	Consumer	Neurontin (Gabapentin) Rofecoxib Simvastatin	PS C C		

Date:05/26/04ISR Number: 4367567-3Report Type:Expedited (15-DaCompany Report #2004032420
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, TID), ORAL		Bone Pain Nerve Graft Pain Post Procedural Pain Suicidal Ideation	Consumer	Neurontin (Gabapentin) Diazepam Lidocaine Hydrochloride Ibuprofen	PS C C C		ORAL

Date:05/26/04ISR Number: 4367572-7Report Type:Expedited (15-DaCompany Report #2004033210
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Benign Prostatic Hyperplasia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Hypotonic Urinary Bladder Urinary Retention Urinary Tract Infection Bacterial		Lisinopril Hydrochlorothiazide	C C		

Date:05/26/04ISR Number: 4367574-0Report Type:Expedited (15-DaCompany Report #2004033211
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 3 IN D		Aggression Anorexia	Consumer	Neurontin (Gabapentin)	PS		
Other		Diet Refusal Drug Intolerance Psychotic Disorder Self-Medication Weight Decreased		Trazodone	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/26/04ISR Number: 4372430-8Report Type:Expedited (15-DaCompany Report #GB-JNJFOC-20040503085

Age:80 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged TRANSDERMAL	Duration Balance Disorder Blood Sodium Decreased 25 UG/HR, Dizziness	Foreign Health Professional	Durogesic (Fentanyl) Patch	PS		
TRANSDERMAL 300 MG, 1 IN 1 DAY, ORAL	Drug Interaction Fall		Neurontin (Gabapentin)	Unknown SS		ORAL
			Co-Proxamol (Aporex) Lisinopril (Lisinopril) Atenolol (Atenolol) Unknown Temazepam (Temazepam) Omeprazole (Omeprazole)	C C C C C		

Date:05/26/04ISR Number: 4372740-4Report Type:Expedited (15-DaCompany Report #2004032843

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL	Duration Anorexia Blindness Oral Intake Reduced Retinopathy	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:05/26/04ISR Number: 4372741-6Report Type:Expedited (15-DaCompany Report #2004032760

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization -	Duration Loss Of Consciousness	Consumer	Neurontin			

Initial or Prolonged Weight Increased (Gabapentin) PS
 3500 MG (1200
 Other
 MG, 3 IN 1 D)
 Celecoxib
 (Celecoxib) C

Date:05/26/04ISR Number: 4372743-XReport Type:Expedited (15-DaCompany Report #2004032455
 Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG (600 Other MG, TID), ORAL		Arthralgia Aspiration Cognitive Disorder Extradural Abscess Fall Influenza Mobility Decreased Muscle Abscess Myalgia Rotator Cuff Syndrome Sepsis Speech Disorder Staphylococcal Infection	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/26/04 ISR Number: 4372745-3 Report Type:Expedited (15-DaCompany Report #2004022432
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Anaemia Astigmatism	Consumer Health	Neurontin (Gabapentin)			ORAL
7200 MG (600 MG, 3 IN 1 D), ORAL		Constipation Dysarthria Endometriosis Fatigue Libido Decreased Muscle Twitching Myalgia Myositis Sleep Disorder	Professional	Propacet (Dextropropoxyphene Napsilate, Paracetamol) Hs (Estrogens Esterified, Methyltestosterone) Paroxetine Hydrochloride (Paroxetine Hydrochloride) Nortriptyline (Nortriptyline) Buspirone Hydrochloride (Buspirone Hydrochloride) Clonazepam (Clonazepam) Dextropropoxyphene Hydrochloride (Dextropropoxyphene Hydrochloride) Laxatives (Laxatives) Calcium Carbonate (Calcium Carbonate) Becosym Forte (Nicotinamide, Pyridoxine Hydrochloride, Riboflavin, Thiamine Multivitamins (Ascorbic Acid/Ergocalciferol/			C C C C C C

Date:05/26/04ISR Number: 4372762-3Report Type:Expedited (15-DaCompany Report #2004032979
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Beta Haemolytic Streptococcal Infection	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL		Drug Ineffective Headache Pain In Extremity Pharmaceutical Product Complaint		Estrogens Conjugated (Estrogens Conjugated) Levothyroxine Sodium (Levothyroxine Sodium) All Other	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Therapeutic Products
(All Other
Therapeutic
Products) C

Date:05/26/04ISR Number: 4373097-5Report Type:Expedited (15-DaCompany Report #2004032454
Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 800 MG Other		Dysphagia Inflammation	Foreign Health	Neurontin (Gabapentin)	PS		
		Limb Injury Platelet Count Decreased Renal Disorder Toxic Skin Eruption White Blood Cell Count Decreased	Professional Company Representative	Antiepileptics (Antiepileptics) Phenobarbital (Phenobarbital) Verapamil Hydrochloride (Verapamil Hydrochloride) Antihypertensives (Antihypertensives) Prednisone (Prednisone)	C C C C		

Date:05/27/04ISR Number: 4369076-4Report Type:Expedited (15-DaCompany Report #2004032849
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Loss Of Consciousness Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		

Date:05/27/04ISR Number: 4369117-4Report Type:Expedited (15-DaCompany Report #2004033431
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Adrenal Neoplasm	Consumer	Neurontin			

ORAL

Cardiac Failure

(Gabapentin)

PS

ORAL

Congestive
Dizziness
Dyspnoea
Emotional Distress
Feeling Abnormal
Feeling Drunk
Oedema Peripheral
Weight Increased

Date:05/27/04ISR Number: 4369120-4Report Type:Expedited (15-DaCompany Report #2004032845
Age:28 YR Gender:Female I/FU:I

Outcome PT
Other Cataract
Diabetes Mellitus
Gastrooesophageal Reflux
Disease
Oral Intake Reduced
Retinal Detachment
Vomiting

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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
10 MG (10 MG, DAILY)		Consumer	Neurontin (Gabapentin)	PS		
			Zyrtec (Tablets) (Cetirizine)	SS		

Date:05/27/04ISR Number: 4369123-XReport Type:Expedited (15-DaCompany Report #2004032848
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Upper Asthenia Depression Diarrhoea Feeling Abnormal Gastric Disorder Gastrointestinal Pain Intestinal Spasm Muscle Spasms Nausea Oral Intake Reduced Oropharyngeal Spasm Suicidal Ideation Vomiting	Consumer	Neurontin (Gabapentin)	PS		
				Simvastatin (Simvastatin)	C		

Date:05/27/04ISR Number: 4369125-3Report Type:Expedited (15-DaCompany Report #2004033208
Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Infection Neutropenia White Blood Cell Count	Health Professional	Neurontin (Gabapentin)	PS		ORAL
100 MG (100 MG, 1 IN 1							

D), ORAL

Abnormal

White Blood Cell Count
Decreased

Fluoxetine
(Fluoxetine) C
Panadeine Co
(Codeine Phosphate,
Paracetamol) C

Date:05/27/04ISR Number: 4369129-0Report Type:Expedited (15-DaCompany Report #2004009823

Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Arthritis Drug Intolerance	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1		Exostosis Knee Arthroplasty	Professional				
D), ORAL		Knee Operation Memory Impairment Post Procedural Complication Somnolence		Insulin (Human Injection, Isophane (Insulin Human Injection, Isophane) Nateglinide (Nateglinide) Insulin (Insulin) Doxepin	C C C		

Freedom Of Information (FOI) Report

Hydrochloride
 (Doxepin Hydrochloride) C
 Atenolol (Atenolol) C
 Simvastatin
 (Simvastatin) C
 Alprazolam
 (Alprazolam) C
 Clorazepate
 Dipotassium
 (Clorazepate
 Dipotassium) C

Date:05/27/04ISR Number: 4369132-0Report Type:Expedited (15-DaCompany Report #2004013496
 Age:10 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Abnormal Behaviour - Aggression Anger Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:05/27/04ISR Number: 4369133-2Report Type:Expedited (15-DaCompany Report #2004032749
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (300 MG, 2 IN 1 D), ORAL		Balance Disorder Coordination Abnormal Dysarthria Fall Head Injury Loss Of Consciousness	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Clonazepam (Clonazepam0 Raloxifene Hydrochloride (Raloxifene Hydrochloride) Esomeprazole (Esomeprazole)	C C C		

Fexofenadine
 Hydrochloride
 (Fexofenadine
 Hydrochloride) C
 Rofecoxib
 (Rofecoxib) C

Date:05/27/04ISR Number: 4369137-XReport Type:Expedited (15-DaCompany Report #2004032764

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (100 MG, TID INTERVAL: EVERY DAY), ORAL ORAL 10 MEQ (10		Aortic Valve Incompetence Cardiac Murmur Drug Effect Decreased Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Calcium (Calcium)	SS		ORAL
				Potassium Chloride (Potassium Chloride)	SS		

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Freedom Of Information (FOI) Report

MEQ,K QD

INTERVAL:

EVERY DAY)

Magnesium	
(Magnesium)	SS
Selenium (Selenium)	SS
Alendronate Sodium	
(Alendronate Sodium)	C
Digoxin (Digoxin)	C
Levothyroxine Sodium	
(Levothyroxine Sodium)	C
Warfarin Sodium	
(Warfarin Sodium)	C
Fluvastatin	
(Fluvastatin)	C
Famotidine	
(Famotidine)	C
Paracetamol	
(Paracetamol)	C
Diphenhydramine Hydrochloride	
(Diphenhydramine Hydrochloride)	C
Estrogens Conjugated	
(Estrogens Conjugated)	C
Furosemide	
(Furosemide)	C

Date:05/27/04ISR Number: 4369139-3Report Type:Expedited (15-DaCompany Report #2003039126
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anovulatory Cycle	Consumer	Neurontin			
Other		Condition Aggravated		(Gabapentin)	PS		ORAL
2400 MG (600		Drug Effect Decreased					
MG, 4 IN 1		Hypoaesthesia					
D), ORAL		Infertility Female		Esomeprazole			

Libido Decreased
Weight Increased

(Esomeprazole) C
Panadeine Co
(Codeine Phosphate,
Paracetamol) C
Mometasone Furoate
(Mometasone Furoate) C
Cyclobenzaprine
Hydrochloride
(Cyclobenzaprine
Hydrochloride) C

Date:05/27/04ISR Number: 4369140-XReport Type:Expedited (15-DaCompany Report #2004032851
Age:77 YR Gender:Female I/FU:I

Outcome PT
Death Abnormal Behaviour
Hospitalization - Agitation
Initial or Prolonged Dehydration
Depressed Level Of
Consciousness
Moaning

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Dose	Duration	Oral Intake Reduced Sedation Treatment Noncompliance	Report Source	Product	Role	Manufacturer	Route
3900 MG (300 MG, 3 IN 1 D), ORAL		Urinary Tract Infection	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Warfarin Sodium (Warfarin Sodium)	C		
				Spiroinolactone (Spiroinolactone)	C		
				Lisinopril (Lisinopril)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Metoprolol Tartrate (Metopropol Tartrate)	C		
				Colchicine (Colchicine)	C		

Date:05/27/04ISR Number: 4369854-1Report Type:Expedited (15-DaCompany Report #2004032842

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Depressed Level Of Consciousness	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL
		Feeling Abnormal Feeling Drunk Gait Disturbance Impaired Driving Ability Road Traffic Accident					

Date:05/27/04ISR Number: 4369855-3Report Type:Expedited (15-DaCompany Report #2004033685

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Foreign	Neurontin			
Other		Intentional Misuse	Health	(Gabapentin)	PS		ORAL
ORAL			Professional				

Date:05/27/04ISR Number: 4369857-7Report Type:Expedited (15-DaCompany Report #2004000436

Age:98 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abdominal Pain	Foreign	Neurontin			
Hospitalization -		Blood Glucose Increased	Health	(Gabapentin)	PS		ORAL
400 MG (100		Colitis Ischaemic	Professional				
Initial or Prolonged		Gangrene	Company				
MG, 4 IN 1		Glucose Urine Present	Representative	Metoclopramide			
Other		Glycosylated Haemoglobin		Hydrochloride	C		
D), ORAL		Increased		Lormetazepam			
		Hyponatraemia		(Lormetazepam)	C		
		Pain In Extremity		Latanoprost			
		Peripheral Ischaemia		(Latanoprost)	C		
		White Blood Cell Count		Gliclazide			
		Increased		(Gliclazide)	C		

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Freedom Of Information (FOI) Report

Lansoprazole
 (Lansoprazole) C
 Paramol-118
 (Dihydrocodeine
 Bitartrate,
 Paracetamol) C
 Brimonidine Tartrate

 (Brimonidine
 Tartrate) C
 Peptac C
 Dorzolamide
 Hydrochloride C
 Amlodipine Besilate

 (Amlodipine
 Besilate) C
 Prochlorperazine
 (Prochlorperazine) C
 Furosemide
 (Furosemide) C

Date:05/28/04ISR Number: 4368734-5Report Type:Expedited (15-DaCompany Report #2004034120
 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		Complex Partial Seizures Tremor	Foreign Health Professional Other	Neurontin (Gabapentin)	PS		ORAL

Date:05/28/04ISR Number: 4368747-3Report Type:Expedited (15-DaCompany Report #2004032476
 Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 900 MG (300		Blood Glucose Fluctuation Convulsion	Foreign Health	Neurontin (Gabapentin)	PS		ORAL

Initial or Prolonged Laboratory Test Abnormal Professional

MG, 3 IN 1

Other

D), ORAL

Amitriptyline
(Amitriptyline)

C

Date:05/28/04ISR Number: 4368761-8Report Type:Expedited (15-DaCompany Report #2004033914

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2100 MG (D), Other ORAL		Anaemia Epilepsy Hepatic Enzyme Increased Oedema Peripheral Pleural Effusion Pulmonary Oedema Renal Impairment Tachycardia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

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Date:05/28/04ISR Number: 4368762-XReport Type:Expedited (15-DaCompany Report #2004033716

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (DAILY), ORAL		Hepatic Function Abnormal	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
15 MG (DAILY), ORAL				Prochlorperazine (Prochlorperazine)	SS		ORAL
				Warfarin Sodium Clathrate (Warfarin Sodium Clarthrate)	C		
				Lansoprazole (Lansoprazole)	C		
				Dexamethasone (Dexamethasone)	C		
				Metformin Hydrochloride (Metformin Hydrochloride)	C		
				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		

Date:05/28/04ISR Number: 4368797-7Report Type:Expedited (15-DaCompany Report #2004033874

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other OPHTHALMIC 4 IN 1 D), ORAL	3600 (600 MG,	Leg Amputation Staphylococcal Infection Tooth Discolouration Tooth Disorder Toothache	Consumer	Neurontin (Tablets) (Gabapentin)	PS		
				Insulin Lispro (Insulin Lispro)	C		

Humalog Mix 25	
(Insulin Lispro,	
Insulin Lispro	
Protamine	
Suspension)	C
Nateglinide	
(Nateglinide)	C
All Other	
Therapeutic Product	
(All Other	
Therapeutic	
Products)	C
Carvedilol	
(Carvedilol)	C
Ezetimibe	
(Ezetimibe)	C
Ramipril (Ramipril)	C
Tolterodine	
L-Tartrate	
(Tolterodine	
L-Tartrate)	C
Venlafaxine	
Hydrochloride	
(Venlafaxine	
Hydrochloride)	C

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Furosemide (Furosemide)	C
Atorvastatin (Atorvastatin)	C
Loperamide Hydrochloride (Loperamide Hydrochloride)	C
Rosiglitazone Maleate (Rosiglitazone Maleate)	C
Fexofenadine Hydrochloride (Fexofenadine Hydrochloride)	C
Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotinamide, Valaciclovir Hydrochloride (Valaciclovir Hydrochloride)	C
Isosorbide (Isosorbide)	C
Montelukast Sodium (Montelukast Sodium)	C
Esomeprazole (Esomeprazole)	C
Oxycocet (Oxycodone Hydrochloride, Paracetamol)	C

Date:05/28/04ISR Number: 4368798-9Report Type:Expedited (15-DaCompany Report #2004033869
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Brain Damage Confusional State	Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG (D),							

ORAL

Diverticular Perforation

Memory Impairment
Post Procedural
Complication

All Other
Therapeutic Products
(All Other
Therapeutic
Products) C
Estradiol C
(Estradiol)
Nadolol (Ndolol) C

Date:05/28/04ISR Number: 4368802-8Report Type:Expedited (15-DaCompany Report #2004033212
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Gangrene	Consumer	Neurontin (Gabapentin)	PS		
Other		Leg Amputation					
100 MG,				Simvastatin			

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(Simvastatin) C
 Valsartan
 (Valsartan) C
 Paracetamol
 (Paracetamol) C
 Ramipril (Ramipril) C
 Allergy Medication
 (Allergy Medication) C

Date:05/28/04ISR Number: 4369733-XReport Type:Expedited (15-DaCompany Report #2004009355
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG Other (TID), ORAL	Abdominal Pain Back Disorder Blood Cholesterol	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
20 MG (DAILY), ORAL	Increased Chest Pain Drug Ineffective		Lipitor (Atorvastatin)	SS		ORAL
1000 MG (BID), ORAL	Drug Intolerance Erectile Dysfunction Facial Pain Feeling Cold Gastrooesophageal Reflux Disease General Physical Health Deterioration		Lithium (Lithium) (Lithium) Naproxen (Naproxen) Rofecoxib (Rofecoxib) Amoxicillin (Amoxicillin)	SS SS SS SS		ORAL
	Gingival Disorder Glossodynia Medication Error Mobility Decreased Neck Pain Oesophageal Spasm Pain In Jaw Sleep Disorder Somnolence Tooth Abscess Tremor		All Other Therapeutic Products (All Other Therapeutic Products) Levothyroxine Sodium (Levothyroxine Sodium) Vitamins (Vitamins) Diltiazem Hydrochloride (Diltiazem)	C C C		

Hydrochloride) C

Date:05/28/04ISR Number: 4369734-1Report Type:Expedited (15-DaCompany Report #2004033139
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Bedridden	Consumer	Neurontin			
Initial or Prolonged	Blood Pressure Decreased		(Gabapentin)	PS		
800 MG						
Other	Full Blood Count		Rofecoxib			
	Decreased		(Rofecoxib)	C		
	Mobility Decreased		Simvastatin			
	Pneumonitis		(Simvastatin)	C		

Date:06/01/04ISR Number: 4367895-1Report Type:Direct Company Report #CTU 219829
Age:71 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Dizziness		Gabapentin	PS		
Initial or Prolonged	Hypovolaemia		Oxycontin	SS		
	Ischaemia		Placebo-Per Study			

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Protocol Titration	
Sche	SS
Placebo-Per Study	
Protocol Titration	
Sche	SS
Famvir 500 Mg Tid	SS
Acetaminophen 500 Mg	
Q6 Hr	SS
Hydrochlorothiazide	C
Celexa	C
Miralax	C
Mobic	C
Multivit	C
Calcium And Vit C	C
Phenergan	C

Date:06/01/04ISR Number: 4371466-0Report Type:Expedited (15-DaCompany Report #2004023332

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intervertebral Disc Protrusion Panic Attack	Consumer	Neurontin (Gabapentin)	PS		

Date:06/01/04ISR Number: 4371488-XReport Type:Expedited (15-DaCompany Report #2004022537

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Amnesia Anorexia	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other		Aphasia					
3600 MG (7 TIMES DAILY)		Balance Disorder					
ORAL		Breast Mass Breast Pain		Dilantin Suspension (Phenytoin Sodium)	SS		ORAL
500 MG ORAL		Convulsion Delusional Perception Drug Level Fluctuating Drug Toxicity		Protein Supplements Phenobarbital	C C		

Dysphagia
Dysuria
Eating Disorder
Fall
Gynaecomastia
Headache
Hypoaesthesia
Impatience
Insomnia
Liver Function Test
Abnormal
Mood Swings
Nausea
Personality Change
Pruritus Generalised
Thermal Burn
Thirst Decreased
Tooth Injury
Tooth Loss
Vomiting
Weight Decreased
Weight Increased

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Freedom Of Information (FOI) Report

Date:06/01/04ISR Number: 4371602-6Report Type:Expedited (15-DaCompany Report #2004033942
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 50 MG, ORAL	3 YR	Abnormal Dreams	Consumer	Zoloft (Sertraline)	PS		ORAL
		Accident		Neurontin			
		Condition Aggravated		(Gabapentin)	SS		
		Depression		Esomeprazole			
		Drug Effect Decreased		(Esomeprazole)	SS		
		Gastrooesophageal Reflux		Hydromorphone			
		Disease		Hydrochloride			
		Headache		(Hydromorphone			
		Hypoaesthesia		Hydrochloride)	C		
		Nerve Injury		Morphine	C		
		Pain					
		Suicidal Ideation					
		Treatment Noncompliance					

Date:06/01/04ISR Number: 4371633-6Report Type:Expedited (15-DaCompany Report #2004030542
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (300 MG, 2 IN 1 D), ORAL		Congenital Anomaly	Consumer	Neurontin			
		Developmental	Health	(Gabapentin)	PS		ORAL
		Coordination Disorder					
		Drug Exposure During	Professional				
		Pregnancy					
		Myoclonic Epilepsy		Montelukast Sodium			
		Nuclear Magnetic		(Montelukast Sodium)	C		
		Resonance Imaging Brain					
		Abnormal					

Date:06/01/04ISR Number: 4371635-XReport Type:Expedited (15-DaCompany Report #2004034123
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Distension	Consumer	Neurontin			

300 MG (300	Abdominal Pain Upper	(Gabapentin)	PS	ORAL
MG, DAILY	Drug Ineffective			
INTERVAL:	Exostosis			
EVERY DAY),	Hysterectomy			
ORAL	Inflammation			
	Insomnia			
	Neck Pain			
	Nerve Compression			
	Ovarian Cyst			
	Swelling			

Date:06/02/04ISR Number: 4371489-1Report Type:Expedited (15-DaCompany Report #2004028507
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dyspnoea	Health	Neurontin			
Other		Feeling Abnormal	Professional	(Gabapentin)	PS		
1800 MG (600		Respiratory Arrest	Company				
MG, 3 IN 1			Representative				
D), UNKNOWN				Methadone			
				(Methadone)	SS		
40 MG (10 MG,							
4 IN 1 D),							
UNKNOWN							
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Freedom Of Information (FOI) Report

Date:06/02/04ISR Number: 4371491-XReport Type:Expedited (15-DaCompany Report #2004034119
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Oxcarbazepine (Oxcarbazepine)	C		
Other		Device Failure Post Procedural	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
2400 MG (600		Complication					
MG, 4 IN 1		Weight Increased					
D), ORAL				Furosemide (Furosemide)	C		
				Metformin Hydrochloride (Metformin Hydrochloride)	C		
				Pioglitazone (Pioglitazone)	C		
				Glimepiride (Pioglitazone)	C		
				Glimepiride (Glimepride)	C		
				Rofecoxib (Rofecoxib)	C		
				Cetirizine Hydrochloride (Cetirizine Hydrochloride)	C		
				Ramipril (Ramipril)	C		
				Zolpidem Tartrate (Zolpidem Tartrate)	C		
				Amitriptyline (Amitriptyline)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Ultracet (Paracetamol,			

Tramadol
Hydrochloride) C

Date:06/02/04ISR Number: 4371492-1Report Type:Expedited (15-DaCompany Report #2004019974
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Neurontin			
1800 MG		Anaemia Vitamin B12	Health	(Gabapentin)	PS		ORAL
(TID), ORAL		Deficiency	Professional				
		Cognitive Disorder		Panadeine Co			
		Herpes Zoster		(Codeine Phosphate, Paracetamol)	C		
				Benadryl			
				(Diphenhydramine Hydrochloride)	C		
				Albuterol			
				(Salbutamol)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/02/04ISR Number: 4371493-3Report Type:Expedited (15-DaCompany Report #2004034128
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Condition Aggravated Coordination Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
3600 MG (900 MG, 4 IN 1 D), ORAL		Fibromyalgia Muscle Spasms					
		Psychotic Disorder Raynaud'S Phenomenon Treatment Noncompliance		Moduretic "Msd" (Amiloride Hydrochloride, Hydrochlorothiazide) Vitamins (Vitamins) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Alprazolam (Alprazolam) Fentanyl (Fentanyl)	C C C C		

Date:06/02/04ISR Number: 4371496-9Report Type:Expedited (15-DaCompany Report #2004007885
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Upper Arthralgia Condition Aggravated Convulsion Difficulty In Walking	Consumer Health Professional	Dilantin Suspension (Phenytoin Sodium) (Phenytoin Sodium) Neurontin (Gabapentin)	PS SS		ORAL
1200 MG (TID), ORAL		Feeling Abnormal					
75 MG (DAILY), ORAL		Hypoaesthesia Insomnia Pain Pain In Extremity		Phenobarbital (Phenobarbital) Topiramate	SS		ORAL

100 MG	Partial Seizures	(Topiramate)	SS	
(DAILY)	Periarthritis			
	Rash	Oxcarbazepine		
	Vaginal Haemorrhage	(Oxcarbazepine)	SS	
		Levetiracetam		
		(Levetiracetam)	SS	ORAL
500 MG (250				
MG, BID),				
ORAL				
		Fluoxetine		
		(Fluoxetine)	C	

Date:06/02/04ISR Number: 4371556-2Report Type:Expedited (15-DaCompany Report #2004020149
Age:59 YR Gender:Unknown I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Blood Disorder	Health	Neurontin			
Initial or Prolonged	Gout	Professional	(Gabapentin)	PS		ORAL
2700 MG						
(BID), ORAL	Laboratory Test Abnormal					
	Pain		All Other			
			Therapeutic Products			
			(All Other			
			Therapeutic			
			Products)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/02/04ISR Number: 4371573-2Report Type:Expedited (15-DaCompany Report #2004021501
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Balance Disorder Condition Aggravated	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
300 MG (DAILY), ORAL		Depression	Professional				
20 MG (DAILY)		Drug Interaction		Escitalopram	SS		
		Fall Feeling Drunk Hallucination, Visual Migraine Nightmare Suicide Attempt		Allegra-D (Fexofenadine, Pseudoephedrine Hydrochloride) Propacet (Dextropropoxyphene Napsilate, Paracetamol) Pantoprazole (Pantoprazole) Docusate (Docusate)	C C C C		

Date:06/02/04ISR Number: 4372071-2Report Type:Expedited (15-DaCompany Report #200410158BFR
Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG TOTAL Initial or Prolonged DAILY ORAL		Eczema	Foreign Health	Glucor (Acarbose)	PS		ORAL
20 MG, TOTAL DAILY, ORAL		Self-Medication	Professional	Furosemide	SS		ORAL
600 MG, TOTAL DAILY , ORAL		Toxic Skin Eruption	Other	Neurontin (Gabapentin)	SS		ORAL
300 MG, TOTAL				Aprovel (Irbesartan)	SS		ORAL

DAILY, ORAL		Contramal (Tramadol Hydrochloride)	SS	ORAL
150 MG, BID,				
ORAL		Effexor (Venlafaxine Hydrochloride)	SS	ORAL
75 MG, TOTAL				
DAILY, ORAL		Ogast (Lansoprazole)	SS	ORAL
15 MG, TOTAL				
DAILY, ORAL		Difrarel	SS	ORAL
300 MG, TOTAL				
DAILY, ORAL		Dafalgan (Paracetamol)	SS	ORAL
BID, ORAL		Amlor (Amlodipine Besilate)	SS	ORAL
10 MG, TOTAL				
DAILY, ORAL		Vastarel (Trimetazidine Hydrochloride)	SS	ORAL
60 MG, TOTAL				
DAILY, ORAL				

Date:06/02/04ISR Number: 4372280-2Report Type:Expedited (15-DaCompany Report #2004034256
Age:43 YR Gender:Male I/FU:I

Outcome	PT	Report Source
Other	Aplastic Anaemia	Foreign Health Professional

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Freedom Of Information (FOI) Report

Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
1800 MG ORAL			Neurontin (Gabapentin)	PS		ORAL

Date:06/02/04ISR Number: 4372443-6Report Type:Expedited (15-DaCompany Report #2004030937
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Pressure Increased Carpal Tunnel Syndrome Headache	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Cephalosporins And Related Substances (Cephalosporins And Related Substances) Ketorolac (Ketorolac) Nimesulide Nimesulide) Antihypertensives (Antihypertensives) All Other Therapeutic Products (All Other Therapeutic Products)	PS C C C C C		

Date:06/03/04ISR Number: 4373304-9Report Type:Expedited (15-DaCompany Report #2004024700
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Accident At Work	Health Professional	Neurontin (Gabapentin) Zoloft (Sertraline) Valdecoxib (Valdecoxib) Lansoprazole (Lansoprazole)	PS SS SS SS		

All Other
Therapeutic Products
(All Other
Therapeutic
Products) C

Date:06/03/04ISR Number: 4373311-6Report Type:Expedited (15-DaCompany Report #2004034121
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cellulitis	Consumer	Neurontin	PS		
Other		Cystitis		(Gabapentin)			
(300 MG,)		Depression					
		Diabetes Mellitus					
		Dizziness					
		Drug Ineffective					
		Feeling Abnormal					
		Lung Neoplasm Malignant					
		Vomiting					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/04/04ISR Number: 4374569-XReport Type:Expedited (15-DaCompany Report #2004005958

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2400 MG (QID), ORAL		Abnormal Behaviour Bronchitis Feeling Abnormal	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1000 MG (BID), ORAL		Pharmaceutical Product Counterfeit		Naproxen (Naprosyn)	SS		ORAL
		Sinusitis Tonic Clonic Movements		Clonazepam (Clonazepam)	C		
				Lamotrigine (Lamotrigine)	C		
				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Calcium (Calcium) Fish Oil (Fish Oil)	C C C		

Date:06/04/04ISR Number: 4374590-1Report Type:Expedited (15-DaCompany Report #2004034717

Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 Other MG, 2 IN 1 D), ORAL		Bone Infection Leukopenia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Bactrim (Sulfamethoxazole, Trimethoprim)	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin (Tablets)			
		Pharmaceutical Product		(Gabapentin)	PS		ORAL
3600 MG							
(TID), ORAL		Complaint					
				Topiramate			
				(Topiramate)	C		
				Pravastatin Sodium			
				(Pravastatin Sodium)	C		
				Sertraline			
				Hydrochloride			
				(Sertraline			
				Hydrochloride)	C		
				Sotalol			
				Hydrochloride			
				(Sotalol			
				Hydrochloride)	C		
				Buspirone			
				Hydrochloride			
				(Buspirone			
				Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/04/04ISR Number: 4374854-1Report Type:Expedited (15-DaCompany Report #2004034724

Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Difficulty In Walking	Consumer	Neurontin			
Other		Dysphagia		(Gabapentin)	PS		ORAL
300 MG, ORAL		Hypoaesthesia		All Other			
		Pain		Therapeutic Products			
		Pharmaceutical Product		(All Other			
		Complaint		Therapeutic			
		Throat Irritation		Products)	C		

Date:06/04/04ISR Number: 4374894-2Report Type:Expedited (15-DaCompany Report #2004034979

Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Cytolytic Hepatitis	Foreign	Neurontin			
Initial or Prolonged		Hepatitis Cholestatic	Health	(Gabapentin)	PS		ORAL
(TID), ORAL			Professional	Tramadol			
				Hydrochloride			
				(Tramadol			
				Hydrochloride)	SS		ORAL
(BID), ORAL							

Date:06/04/04ISR Number: 4374904-2Report Type:Expedited (15-DaCompany Report #2004034981

Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation	Foreign	Gabapentin			
ORAL		Condition Aggravated	Health	(Gabapentin)	PS		ORAL
		Depression	Professional	Venlafaxine			
				Hydrochloride			
				(Venlafaxine			
				Hydrochloride)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation	Consumer	Neurontin			
		Cerebrovascular Accident		(Gabapentin)	PS		ORAL
400 MG, ORAL		Confusional State		Clopidogrel Sulfate			
		Convulsion		(Clopidogrel			
		Fall		Sulfate)	C		
		Headache		Vicodin			
		Loss Of Consciousness		(Paracetamol,			
				Hydrocodone			
				Bitartrate)	C		
				Diazepam (Diazepam)	C		
				Fexofenadine			
				Hydrochloride			
				(Fexofenadine			
				Hydrochloride)	C		
				Budesonide			
				(Budesonide)	C		
				Furosemide			
				(Furosemide)	C		
				Potassium			
				(Potassium)	C		

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Freedom Of Information (FOI) Report

Date:06/07/04ISR Number: 4375634-3Report Type:Expedited (15-DaCompany Report #8086

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Guillain-Barre Syndrome	Study	Methotrexate	PS		
SUBCUTANEOUS	25 MG WEEKLY	Intestinal Perforation	Health				
SC			Professional	Remicade	SS		
INTRAVENOUS	3 MG/KG FREQ						
IV				Prednisone	SS		
20 MG BID				Neurontin	SS		
300 MG				Tricor	SS		
54 MG BID				Folic Acid	SS		
1 MG DAILY				Bumex	SS		
1 MG DAILY				Atenolol	SS		
25 MG DAILY				Darvocet-N	SS		

Date:06/07/04ISR Number: 4375693-8Report Type:Expedited (15-DaCompany Report #2004034700

Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Abdominal Pain Cerebral Toxoplasmosis Convulsion Drug Interaction Drug Intolerance Dysaesthesia Dyspnoea Folate Deficiency Hepatic Function Abnormal Hepatocellular Damage	Foreign Health Professional	Viracept (Tablet) (Nelfinavir Mesilate) Neurontin (Gabapentin) (Gabapentin) Doxorubicin Hydrochloride (Doxorubicin Hydrochloride)	PS SS SS		
INTRAVENOUS	35 MG (35 MG	Hyperaesthesia					

MG)

INTRAVENOUS	Hyperlactacidaemia			
	Kaposi'S Sarcoma	Tenofovir Disoproxil		
	Liver Disorder	Fumarate (Tenofovir		
245 MG, ORAL	Neuralgia	Disoproxil Fumarate)	SS	ORAL
	Oedema	Kaletra (Lopinavir,		
(2 IN D),	Pancreatic Disorder	Ritonavir)	SS	ORAL
ORAL	Pancytopenia			
	Pulmonary Hypertension	Domperidone		
ORAL	Rash	(Domperidone)	SS	ORAL
	Renal Failure Chronic	Zidovudine		
		(Zidovudine)	C	
		Zalcitabine		
		(Zalcitabine)	C	
		Stavudine		
		(Stavudine)	C	
		Lamivudine		
		(Lamivudine)	C	
		Indinavir		
		(Indinavir)	C	
		Ritonavir		
		(Ritonavir)	C	
		Efavirenz		
		(Efavirenz)	C	
		Didanosine		
		(Didanosine)	C	
		Bosentan (Bosentan)	C	
		C	
		Pyrimethamine		
		(Pyrimethamine)	C	
		Sulfadiazine		

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(Sulfadiazine)	C
Calcium Foliniate	
(Calcium Folate)	C
Pentamidine	
(Pentamidine)	C
Clindamycin	
(Clindamycin)	C
Vitamins (Vitamins)	C
Levocarnitine	
(Levocarnitine)	C
Furosemide	
(Furosemide)	C
Potassium Chloride	
(Potassium Chloride)	C
Bleomycin	
(Bleomycin)	C
Abacavir Sulfate	
(Abacavir Sulfate)	C
Ondansetron	
Hydrochloride	
(Ondansetron	
Hydrochloride)	C

Date:06/07/04ISR Number: 4376065-2Report Type:Expedited (15-DaCompany Report #2004036067
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Thrombocytopenia	Foreign Health Professional Company Representative	Gabapentin (Gabapentin)	PS		ORAL

Date:06/07/04ISR Number: 4376075-5Report Type:Expedited (15-DaCompany Report #2004028026
 Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG		Balance Disorder Dysarthria	Foreign Health	Neurontin (Gabapentin)	PS		ORAL

(DAILY), ORAL

Motor Dysfunction

Professional

Sedation
Somnolence
Vertigo

Company
Representative

Bisoprolol
(Bisoprolol) C
Candesartan
(Candesartan) C
Atorvastatin
(Atorvastatin) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C
Sertraline
(Sertraline) C

Date:06/08/04ISR Number: 4373005-7Report Type:Expedited (15-DaCompany Report #US-MERCK-0307USA01625
Age:57 YR Gender:Male I/FU:F

Outcome

Death

Hospitalization -
Initial or Prolonged

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Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hyperhidrosis		Vioxx	PS	Merck & Co., Inc	ORAL
53 DAY		Hyponatraemia		Tracleer	SS		ORAL
		Oesophageal Disorder		Tracleer	SS		ORAL
UNKNOWN		Pneumonia Aspiration		Cardizem	SS		
UNKNOWN		Respiratory Arrest		Neurontin	SS		
UNKNOWN		Sepsis		Flolan	SS		
INTRAVENOUS		Thrombocytopenia		Elavil	C		ORAL
UNKNOWN				Coumadin	C		
UNKNOWN				Aldactone	C		
UNKNOWN				Oxycontin	C		
UNKNOWN				Celexa	C		
UNKNOWN				Allegra	C		
UNKNOWN				Prevacid	C		
UNKNOWN				Lasix (Furosemide)	C		

Date:06/08/04ISR Number: 4373255-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0263843A
 Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300MG Twice		Abdominal Pain Upper		Combivir	PS	Glaxosmithkline	ORAL
Initial or Prolonged per day 3 WK		Bilirubin Conjugated					
400MG Twice		Increased		Kaletra	SS		ORAL
per day 3 WK		Blood Alkaline					
500MG Three		Phosphatase Increased		Neurontin	SS		ORAL

times per day	3	WK	Blood Amylase Increased				
			Blood Bilirubin Increased	Norvir	SS		ORAL
			Hepatitis Cholestatic	Malocide	C	Glaxosmithkline	ORAL
75MG Per day			Jaundice	Adiazine	C		ORAL
500MG Six			Lipase Increased				
times per day			Neutropenia	Osfolate	C	Glaxosmithkline	ORAL
			Pancreatitis	Urbanyl	C		ORAL
2	DAY		Rash Maculo-Papular				
			Transaminases Increased				

Date:06/08/04ISR Number: 4376398-XReport Type:Expedited (15-DaCompany Report #USA-2004-0015309
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dizziness Hypovolaemia	Study Health Professional	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet			ORAL
ORAL					PS		
500 MG, TID				Placebo (Placebo) Famvir (Famciclovir)	SS SS		
				Gabapentin/Placebo (Gabapentin) Phenergan "Natrappharm" (Promethazine Hydrochloride)	SS SS		
50 MG, Q6H							
PRN				Miacalcin (Calcitonin, Salmon) Mobic (Meloxicam) Acetaminophen Tablet Hydrochlorothiazide	C C C		

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(Hydrochlorothiazide
)
Miracox (Macrogol) C
Multivit (Vitamins
Nos) Tablet C
Calcium Tablet C
Celexa (Citalopram
Hydrobromide) C
Vitamin C (Ascorbic
Acid) Tablet C

Date:06/08/04ISR Number: 4376716-2Report Type:Expedited (15-DaCompany Report #2004035634

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour Anxiety	Health Professional	Geodon (Ziprasidone)	PS		ORAL
40 MG (20 MG, 2 IN 1 D), ORAL		Delusion Dyskinesia					
ORAL		Insomnia Knee Arthroplasty		Neurontin (Gabapentin)	SS		ORAL
		Loss Of Employment Movement Disorder Obsessive Thoughts Obsessive-Compulsive Disorder Somnolence Thinking Abnormal Toe Deformity Tremor		Clonazepam (Clonazepam) Dyazide (Hydrochlorothizide, Triameterene) Lorazepam (Lorazepam)	C C C		

Date:06/09/04ISR Number: 4377441-4Report Type:Expedited (15-DaCompany Report #230008M04CAN

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neutrophil Count Decreased	Foreign Health	Rebif (Interferon Beta)	PS		
44 MCG, 3 IN							

1 WEEKS,	White Blood Cell Count	Professional				
	Decreased			Carbamazepine	SS	
				Gabapentin	SS	
Date:06/09/04ISR Number: 4378032-1Report Type:Expedited (15-DaCompany Report #2004035776						
Age:22 YR	Gender:Male	I/FU:I				
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Renal Tubular Disorder	Foreign	Neurontin			
Initial or Prolonged		Health	(Gabapentin)	PS		
1800 MG (600						
		Professional				
MG, 3 IN 1 D)						
Date:06/09/04ISR Number: 4378445-8Report Type:Expedited (15-DaCompany Report #2004023104						
Age:45 YR	Gender:Female	I/FU:F				
Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Convulsion	Consumer	Neurontin (Tablets)			
	Pharmaceutical Product	Health	(Gabapentin)	PS		ORAL
3600 MG						
(TID), ORAL	Complaint	Professional				
			Topiramate			
			(Topiramate)	C		
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Pravastatin Sodium
 (Ravastatin Sodium) C
 Sertraline
 Hydrochloride) C
 Sotalol
 Hydrochloride
 (Sotalol
 Hydrochloride) C
 Buspirone
 Hydrochloride
 (Buspirone
 Hydrochloride) C

Date:06/09/04ISR Number: 4378472-0Report Type:Expedited (15-DaCompany Report #2004029359
 Age:67 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL Other	Coordination Abnormal Dyskinesia Medication Error Self-Medication	Consumer Health Professional	Neurontin (Gabapentin) Alprazolam (Alprazolam) Levothyroxine Sodium (Levothyroxine Sodium)	PS SS C		ORAL

Date:06/09/04ISR Number: 4378508-7Report Type:Expedited (15-DaCompany Report #2004029398
 Age:45 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 3600 MG (3 IN Other 1 D), ORAL	Aplastic Anaemia Bacterial Sepsis Bone Marrow Depression Bronchitis Conjunctival Haemorrhage Cough Ear Pain	Health Professional	Neurontin (Gabapentin) Temozolomide (Temozolomide) Bactrim	PS SS		ORAL ORAL

	Headache	(Sulfamethoxazole,		
	Herpetic Stomatitis	Trimethoprim)	SS	ORAL
(BID), ORAL				
	Implant Site Infection	Levofloxacin		
	Implant Site Reaction	(Levofloxacin)	SS	ORAL
500 MG (500				
MG, DAILY),	Malaise			
ORAL	Myalgia			
	Petechiae	Thalidomide		
	Pyrexia	(Thalidomide)	SS	
1200 MG (1200				
MG, DAILY	Sepsis			
	Tenderness	Heparin (Heparin)	SS	
INTRAVENOUS	INTRAVENOUS			
	Upper Respiratory Tract	Famciclovir		
	Infection	(Famciclovir)	C	
		Codeine (Codeine)	C	
		Phenobarbital		
		(Phenobarbital)	C	
		Acetylsalicylic Acid		
		(Acetylsalicylic		
		Acid)	C	
		Heparin-Fraction,		
		Sodium Salt		
		(Heparin-Fraction,		
		Sodium Salt)	C	
		Carmustine		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Carmustine) C
 Paclitaxel
 (Paclitaxel) C
 Irinotecan
 (Irinotecan) C
 Ceftriaxone
 (Ceftriaxone) C

Date:06/09/04ISR Number: 4378795-5Report Type:Expedited (15-DaCompany Report #2004035959
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eyelid Function Disorder	Consumer	Neurontin			
		Feeling Abnormal		(Gabapentin)	PS		ORAL
ORAL		Incoherent		Antidepressants			
		Treatment Noncompliance		(Antidepressants)	C		
				Cholesterol			
				(Cholesterol)	C		
				All Other			
				Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	C		

Date:06/10/04ISR Number: 4378904-8Report Type:Expedited (15-DaCompany Report #2004036612
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cytolytic Hepatitis	Foreign	Neurontin			
Initial or Prolonged		Impaired Healing	Health	(Gabapentin)	PS		
			Professional				
			Company				
			Representative				

Date:06/10/04ISR Number: 4378910-3Report Type:Expedited (15-DaCompany Report #2004036550
 Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged (600 MG,,),	Coma Condition Aggravated	Foreign Consumer	Neurontin (Tablets) (Gabapentin)	PS	ORAL
ORAL	Dizziness				
	Facial Neuralgia Nausea		Carbamazepine (Carbamazepine)	C	

Date:06/10/04ISR Number: 4379085-7Report Type:Expedited (15-DaCompany Report #2004024707
Age:35 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 4800 MG (1600 Other MG, 3 IN 1 D)	Anaemia Fibromyalgia Multiple Sclerosis Pain	Consumer Health Professional	Neurontin (Gabapentin)	PS		
			Levothyroxine Sodium (Levothyroxine Sodium)	C		
			Esomeprazole (Esomeprazole) Oxycodone Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

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(Oxycodone Hydrochloride)	C
Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C
Lorazepam (Lorazepam)	C
Tramadol Hydrochloride (Tramadol Hydrochloride)	C
Paracetamol (Paracetamol)	C
Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride)	C
Tizanidine Hydrochloride (Tizanidine Hydrochloride)	C
Rofecoxib (Rofecoxib)	C

Date:06/10/04ISR Number: 4379086-9Report Type:Expedited (15-DaCompany Report #2004036483

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arachnoiditis	Health	Neurontin			
Other		Chemical Poisoning Condition Aggravated	Professional	(Gabapentin)	PS		

Date:06/10/04ISR Number: 4379088-2Report Type:Expedited (15-DaCompany Report #2004036509

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gastric Bypass	Health	Neurontin			
Other			Professional	(Gabapentin)	PS		

1800 MG (600

MG, 3 IN 1 D)

Vicodin
(Hydrochloride
Bitartrate,
Paracetamol)

C

Date:06/10/04ISR Number: 4379104-8Report Type:Expedited (15-DaCompany Report #2004021849

Age: Gender:Female I/FU:F

Outcome PT
Other Balance Disorder
Body Height Decreased
Breast Cancer
Drug Effect Decreased
Drug Ineffective
Dysgeusia
Fatigue
Fear Of Falling
Feeling Abnormal
Irritable Bowel Syndrome

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Mental Disorder Obsessive Thoughts Stress					
45 MG (3 IN 1 D), ORAL		Tarsal Tunnel Syndrome Treatment Noncompliance	Consumer Health Professional	Nardil (Phenelzine Sulfate)	PS		ORAL
600 MG (2 IN 1 D), ORAL				Neurontin (Gabapentin)	SS		ORAL
				Alprazolam (Alprazolam) Triobe (Cyanocobalamin, Folic Acid, Pyridoxine) Vitamins (Vitamins)	C C C		

Date:06/10/04ISR Number: 4379304-7Report Type:Expedited (15-DaCompany Report #04-00262
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Abdominal Wall Abscess Blood Albumin Decreased Blood Magnesium Decreased Blood Potassium Decreased Cognitive Disorder	Literature Health Professional Other	Kadian (Morphine Sulfate Sustained Release) Capsules, 20mg (Alpharm Pharmaceuticals)	PS	Alpharma Pharmaceuticals Inc	ORAL
30MG, AS NEEDED, ORAL		Confusional State		Alprazolam Tablets Usp, 0.5mg (Purepac)	SS	Purepac	
0.5MG, AS NEEDED ,		Delirium Delusion Depressed Level Of Consciousness		Trazodone Hydrochloride Tablets,			

100MG, NIGHTLY TRANSDERMAL EVERY THREE DAYS, TRANSDERMAL 80MG, DAILY 300MG, THREE TIMES DAILY	Disorientation Distractibility Disturbance In Attention 125MCG/HR, Joint Range Of Motion Decreased Mental Impairment Movement Disorder Myoclonus Nausea Pain Pyrexia Sedation Speech Disorder Therapeutic Response Unexpected Thinking Abnormal	100mg (Purepac) Fentanyl (Janssen) Fluoxetine (Siegfried) Gabapentin (Pfizer)	SS SS SS SS	Purepac Janssen Siegfried Pfizer
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Date:06/10/04ISR Number: 4379335-7Report Type:Expedited (15-DaCompany Report #KII-2004-0011034
Age:39 YR Gender:Male I/FU:I

Outcome Hospitalization - Initial or Prolonged Other	PT Aggression Blood Pressure Diastolic Decreased Blood Pressure 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
		Heart Rate Increased Intentional Misuse Metabolic Acidosis					
ORAL		Respiratory Acidosis Respiratory Rate Increased Urine Cannabinoids	Study Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		ORAL
ORAL		Increased Vomiting		Other Hypnotics And Sedatives ()	SS		ORAL
ORAL				Neurontin (Gabapentin)	SS		ORAL
ORAL				Ssri()	SS		ORAL
ORAL				Baclofen (Baclofen)	SS		ORAL
ORAL				Trazodone (Trazodone)	SS		ORAL
				Tetrahydrocannabinol (Tetrahydrocannabino l)	SS		

Date:06/10/04ISR Number: 4379341-2Report Type:Expedited (15-DaCompany Report #2004027642
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 3200 MG (TID), ORAL		Grand Mal Convulsion Laboratory Test Abnormal Sedation	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Paxil (Paroxetine Hydrochloride)	C		
				Depakote (Valproate Semisodium)	C		
				Tylenol With Codeine (Codeine Phosphate, Paracetamol)	C		

Date:06/10/04ISR Number: 4379342-4Report Type:Expedited (15-DaCompany Report #2004023523
Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Consumer	Neurontin			
		Nuclear Magnetic	Health	(Gabapentin)	PS		ORAL
900 MG (TID),		Resonance Imaging Brain	Professional				
ORAL		Abnormal		Levothyroxine			
		Sudden Hearing Loss		Sodium			
		Tinnitus		(Levothyroxine	C		
				Sodium)			
				Lidocaine			
				Hydrochloride			
				(Lidocaine			
				Hydrochloride)	C		

Date:06/10/04ISR Number: 4379344-8Report Type:Expedited (15-DaCompany Report #2004021959
Age:55 YR Gender:Female I/FU:F

Outcome	PT
Other	Anger
	Headache
	Postoperative Adhesion
	Skin Discolouration

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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
ORAL		Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Colchicine (Colchicine)	C		
			Estrogens Conjugated (Estrogens Conjugated)	C		
			Vicoprofen (Hydrocodone Bitartrate, Ibuprofen)	C		
			All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:06/10/04ISR Number: 4379346-1Report Type:Expedited (15-DaCompany Report #2004036533
Age:9 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 5 MG (5 MG, DAILY), ORAL		Aggression Anxiety	Consumer	Zyrtec (Tablets) (Cetirizine)	PS		ORAL
		Migraine					
		Obsessive-Compulsive Disorder		Neurontin (Gabapentin)	SS		ORAL
		Personality Disorder Photophobia Sinus Congestion Sinus Disorder Sinus Headache Swelling Swelling Face					

Date:06/14/04ISR Number: 4379218-2Report Type:Expedited (15-DaCompany Report #2004028507
Age:32 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Health Professional	Neurontin (Gabapentin)	PS		
1800 MG (3 IN 1 D)			Company Representative	Methadone (Methadone)	SS		
40 MG (10 MG, 4 IN 1 D)							

Date:06/14/04ISR Number: 4379244-3Report Type:Expedited (15-DaCompany Report #2004025316
Age:41 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Other	Acute Respiratory Failure Agitation Alopecia Amenorrhoea Anxiety Arthralgia

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Freedom Of Information (FOI) Report

Asthenia
Back Injury
Blood Bicarbonate
Increased
Blood Bilirubin Increased
Blood Chloride Decreased
Blood Creatine
Phosphokinase Increased
Blood Glucose Increased
Blood Potassium Decreased
Blood Pressure Systolic
Increased
Blood Urea Decreased
Bowel Sounds Abnormal
Bursitis
Calcinosis
Circulatory Collapse
Coma
Complex Regional Pain
Syndrome
Confusional State
Crying
Decreased Appetite
Depression
Disorientation
Dysphagia
Electrocardiogram Qt
Corrected Interval
Prolonged
Fibromyalgia
Goitre
Haematocrit Decreased
Hypertension
Hypokalaemia
Hypothyroidism
Intentional Misuse
Irritability
Loss Of Consciousness
Lymphocyte Count
Decreased
Memory Impairment
Multiple Allergies
Mydriasis
Neuropathic Pain
Neutrophil Count
Increased
Ovarian Enlargement

Pco2 Decreased
Pharyngitis
Pharyngolaryngeal Pain
Respiratory Depression
Stress
Suicidal Ideation
Tachycardia
Tenosynovitis
Thyroid Neoplasm
Tri-Iodothyronine
Increased
Urine Abnormality
Vocal Cord Polyp
White Blood Cell Count

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Decreased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
			Tricyclic Antidepressants (Tricyclic Antidepressants)	SS		
			Benzodiazepine Derivatives (Benzodiazepine Derivatives)	SS		
			All Other Therapeutic Products (All Other Therapeutic Products)	SS		
			Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
			Carisoprodol (Carisoprodol)	C		
			Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		

Date:06/14/04ISR Number: 4379285-6Report Type:Expedited (15-DaCompany Report #2004030542

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly Other TRANSPLACENTAL MG, TWICE A DAY), PLACENTAL	600 MG	(300 Drug Exposure During Pregnancy Myoclonic Epilepsy Nuclear Magnetic	Consumer	Neurontin (Gabapentin)	PS		
				Montelukast Sodium			

Resonance Imaging Brain
Abnormal

(Montelukast Sodium) C

Date:06/14/04ISR Number: 4379528-9Report Type:Expedited (15-DaCompany Report #2004037137
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Asthenia Bronchial Obstruction	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Feeling Abnormal Lung Disorder Neuropathic Pain Pneumonia Pyrexia					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/14/04ISR Number: 4379529-0Report Type:Expedited (15-DaCompany Report #2004009410
 Age:60 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL Other	Arrhythmia Atrial Fibrillation	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
	Benign Prostatic Hyperplasia Micturition Urgency Pollakiuria Urinary Incontinence Urinary Retention	Professional	Venlafaxine Hydrochloride (Venlafaxine Hydrochloride) Buspirone Hydrochloride (Buspirone Hydrochloride) Testosterone (Testosterone) Olanzapine (Olanzapine)	C C C C		

Date:06/14/04ISR Number: 4379823-3Report Type:Expedited (15-DaCompany Report #2004032843
 Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability Other ORAL	Anorexia Blindness	Consumer	Neurontin (Gabapentin)	PS		ORAL
	Condition Aggravated Diabetes Mellitus Exercise Lack Of Fatigue Retinopathy Somnolence					

Date:06/14/04ISR Number: 4379824-5Report Type:Expedited (15-DaCompany Report #2004036523
 Age:62 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Disability	Condition Aggravated	Consumer	Neurontin (Tablets)			

Other
3200 MG (800
MG, QID),
ORAL

Fibromyalgia

(Gabapentin)

PS

ORAL

Amitriptyline
(Amitriptyline) C
Pioglitazone
(Pioglitazone) C
Glibomet
(Glibenclamide,
Metformin
Hydrochloride) C
Simvastatin
(Simvastatin) C
Thyroid (Thyroid) C

Date:06/14/04ISR Number: 4379825-7Report Type:Expedited (15-DaCompany Report #2004037133
Age:34 YR Gender:Male I/FU:I

Outcome PT
Disability Amnesia
Other Drug Ineffective
Gastrointestinal Pain

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Irritable Bowel Syndrome

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
3200 MG (1600 MG, BID) ORAL		Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Abacavir Sulfate (Abacavir Sulfate)	C		
			Atazanavir Sulfate (Atazanavir Sulfate)	C		
			Stavudine (Stavudine)	C		
			Ritonavir (Ritonavir)	C		
			Desloratadine (Desloratadine)	C		
			Triamcinolone Acetonide (Triamcinolone Acetonide)	C		
			Aciclovir (Aciclovir)	C		
			Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotinamide,	C		

Date:06/14/04ISR Number: 4379826-9Report Type:Expedited (15-DaCompany Report #2004033212

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Gangrene	Consumer	Neurontin (Gabapentin)	PS		
Other		Leg Amputation	Health Professional	Simvastatin (Simvastatin)	C		
				Valsartan (Valsartan)	C		
				Paracetamol (Paracetamol)	C		
				Ramipril (Ramipril)	C		

Date:06/14/04ISR Number: 4380249-7Report Type:Expedited (15-DaCompany Report #033-0945-990076
Age:48 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (400 MG, 3 IN 1 D) ORAL	Sudden Hearing Loss	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/14/04ISR Number: 4380369-7Report Type:Expedited (15-DaCompany Report #2004030937

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased	Foreign	Neurontin			
		Carpal Tunnel Syndrome	Health	(Gabapentin)	PS		
		Headache	Professional	Cephalosporins And			
			Company	Related Substances			
			Representative	Cephalosporins And			
				Related Substances)	C		
				Ketorolac			
				(Ketorolac)	C		
				Nimesulide			
				(Nimesulide)	C		
				Antihypertensives			
				(Antihypertensives)	C		
				All Other			
				Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	C		

Date:06/14/04ISR Number: 4380370-3Report Type:Expedited (15-DaCompany Report #2004031169

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -							
Initial or Prolonged		Condition Aggravated	Foreign	Neurontin			
INTERVAL		Insomnia	Consumer	(Gabapentin)	PS		
		Muscle Spasms		Insulin (Insulin)	C		
		Neuropathy		Fluoxetine			
		Pain In Extremity		(Fluoxetine)	C		
				Levothyroxine			
				(Levothyroxine)	C		

Date:06/14/04ISR Number: 4380431-9Report Type:Expedited (15-DaCompany Report #2004030834

Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anticonvulsant Drug Level	Consumer	Neurontin (Tablets)			

ORAL	Increased	(Gabapentin)	PS	ORAL
	Convulsion	Levothyroxine Sodium (Levothyroxine Sodium)	C	
		Tocopherol (Tocopherol)	C	
		Biotin (Biotin)	C	
		Esomeprazole (Esomeprazole)	C	

Date:06/15/04ISR Number: 4382100-8Report Type:Expedited (15-DaCompany Report #2004032194
Age:61 YR Gender:Male I/FU:F

Outcome	PT
Other	Depression Feeling Abnormal Multiple Sclerosis Pain In Extremity Pharmaceutical Product Complaint

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Treatment Noncompliance

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, 3 IN 1 D), ORAL		Consumer Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
INTRAMUSCULAR	30 MCG (30 MCG, 1 IN 1 WK),		Interferon Beta (Interferon Beta)	SS		
INTRAMUSCULAR			Tamsulosin Hydrochloride (Tamsulosin Hydrochloride)	C		
			Oxybutynin Hydrochloride (Oxybutynin Hydrochloride)	C		

Date:06/16/04ISR Number: 4380912-8Report Type:Expedited (15-DaCompany Report #2004032696
 Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 900 MG (300 MG , 3 IN 1 D) ORAL		Asthenia Coordination Abnormal	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Dizziness Impaired Driving Ability					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Alcohol Use Cholestasis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Cytolytic Hepatitis	Professional	Carbamazepine (Carbamazepine)	SS		ORAL
ORAL				Aporex (Dextropropoxyphene Hydrochloride, Paracetamol)	SS		ORAL
				Pristinamycin (Pristinamycin)	C		
				Omeprazole(Omeprazol e)	C		
				Furosemide(Furosemid e)	C		
				Buflomedil (Buflomedil)	C		
				Lisinopril(Lisinopri l)	C		
				Insulin (Insulin)	C		
				Zopiclone (Zopiclone)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/16/04ISR Number: 4381197-9Report Type:Expedited (15-DaCompany Report #2012272

Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Accident	Consumer	Oxycontin Tablets			
Hospitalization -	Accidental Overdose	Health	(Oxycodone			
Initial or Prolonged	Asthenia	Professional	Hydrochloride)Cr			
Other	Back Pain	Other	Tablet	PS		
SEE IMAGE						
Required	Drug Abuser		Diazepam (Diazepam)	SS		
Intervention to	Erectile Dysfunction		Oxazepam (Oxazepam)	SS		
Prevent Permanent	Haematemesis		Temazepam			
Impairment/Damage	Haemorrhage		(Temazepam)	SS		
	Headache		Lorazepam			
	Iron Deficiency Anaemia		(Lorazepam)	SS		
	Mallory-Weiss Syndrome		Cannabnoids			
	Respiratory Arrest		(Cannabis)	SS		
	Tachycardia		Diphenhydramine			
	Tooth Disorder		Hydrochloride			
	Tooth Extraction		(Diphenhydramine			
			Hydrochloride)	SS		
			Gabapentin			
			(Gabapentin)	SS		
			Claritin			
			(Loratadine)	C		
			Zoloft (Sertraline			
			Hydrochloride)	C		
			Altace (Ramipril)	C		
			Allopurinol			
			(Allopurinol)	C		
			Depakote (Valproate			
			Semisodium)	C		
			Soma (Carisoprodol)	C		
			Relafen (Nabumetone)	C		

Date:06/16/04ISR Number: 4381456-XReport Type:Expedited (15-DaCompany Report #001-0945-M0101475

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Completed Suicide	Health	Neurontin			
Life-Threatening	Overdose	Professional	(Gabapentin)	PS		
	Suicidal Ideation		Hydroxyzine			
	Victim Of Homicide		Hydrochloride			

(Tablet)(Hydroxyzine
) (Hydroxyzine
Hydrochloride) SS
Hydrocodone
(Hydrocodone) C
Fentanyl (Fentanyl) C

Date:06/16/04ISR Number: 4381464-9Report Type:Expedited (15-DaCompany Report #2004038007

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Hospitalization - Initial or Prolonged Other						
	Brain Neoplasm Malignant	Consumer	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/16/04ISR Number: 4381509-6Report Type:Expedited (15-DaCompany Report #2004037837

Age:73 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG (600 Other MG, 3 IN 1 D), ORAL	Ankle Fracture Blood Cholesterol Increased Foot Fracture	Consumer	Neurontin (Gabapentin)	PS		ORAL
10 MG (10 MG, 1 IN 1 D), ORAL	Herpes Zoster Myalgia Osteoporosis		Lipitor (Atorvastatin)	SS		ORAL
			Rosiglitazone Maleate (Rosiglitazone Maleate)	C		
			Insulin Human (Insulin Human)	C		
			Insulin Glargine (Insulin Glargine)	C		
			Furosemide (Furosemide)	C		
			Amlodipine Besilate (Amlodipien Besilate)	C		
			Folic Acid (Folic Acid)	C		
			Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
			Carisoprodol (Carisoprodol)	C		
			Norgesic Forte (Acetylsalicylic Acid, Caffeine, Orphenadrine Citrate)	C		
			Tocopherol (Tocopherol)	C		

Date:06/16/04ISR Number: 4381537-0Report Type:Expedited (15-DaCompany Report #2004032764

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aortic Valve Incompetence Cardiac Murmur	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG, TID		Condition Aggravated	Professional				
INTERVAL: EVERY DAY)		Drug Effect Decreased					
ORAL		Drug Interaction					
ORAL		Grand Mal Convulsion		Calcium (Calcium)	SS		ORAL
10 MEQ (10 MEQ, QD				Potassium Chloride (Potassium Chloride)	SS		
INTERVAL: EVERY DAY)							
				Magnesium (Magnesium)	SS		
				Selenium (Selenium)	SS		

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Freedom Of Information (FOI) Report

Alendronate Sodium
 (Alendronate Sodium) C
 Digoxin (Digoxin) C
 Levothyroxine Sodium
 (Levothyroxine
 Sodium) C
 Warfarin Sodium
 (Warfarin Sodium) C
 Fluvastatin
 (Fluvastatin) C
 Famotidine
 (Famotidine) C
 Paracetamol
 (Paracetamol) C
 Diphenhydramine
 Hydrochloride
 (Diphenhydramine
 Hydrochloride) C
 Estrogens Conjugated
 (Estrogens
 Conjugated) C
 Furosemide
 (Furosemide) C

Date:06/16/04ISR Number: 4381538-2Report Type:Expedited (15-DaCompany Report #2004037863
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Back Pain Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG (400 MG, 3 IN 1 D) ORAL		Depression Intervertebral Disc Protrusion Medical Device Discomfort Muscle Spasms Nausea Oral Discomfort Weight Decreased		Alprazolam (Alprazolam) Vicodin (Hydrocodone Bitartrate, Paracetamol) Estrogens Conjugated (Estrogens Conjugated) Dicycloverine	C C C		

Hydrochloride
(Dicycloverine
Hydrochloride)

C

Date:06/17/04ISR Number: 4381007-XReport Type:Direct
Age: Gender: I/FU:I

Company Report #USP 56603

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Neurontin	PS	Roche *Pfizer Per Red Book	

LIQUID

Date:06/17/04ISR Number: 4381406-6Report Type:Direct
Age:46 YR Gender:Female I/FU:I

Company Report #CTU 220916

Outcome	PT
Other	Balance Disorder Feeling Abnormal

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Headache Pharmaceutical Product Complaint	Report Source	Product	Role	Manufacturer	Route
1 1/2 TAB				Neurontin Nt 16 600 Mg Tabs	PS		ORAL
BSPO				Premarin	C		
				Synthroid	C		
				Lotisone	C		
				Keppra	C		
				Seroquel	C		
				Zoloft	C		
				Ditropan	C		
				Neurontin	C		

Date:06/17/04ISR Number: 4382515-8Report Type:Expedited (15-DaCompany Report #2004038392
Age:83 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Fatigue	Foreign	Neurontin			
Hospitalization - 1800 MG		Heart Rate Increased	Health	(Gabapentin)	PS		
Initial or Prolonged		Jaundice	Professional Company Representative	Oxcarbazepine (Oxcarbazepine)	C		

Date:06/17/04ISR Number: 4382752-2Report Type:Expedited (15-DaCompany Report #2004037988
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Amnesia	Health	Neurontin			
		Convulsion	Professional	(Gabapentin)	PS		
		Ischaemia					

Date:06/17/04ISR Number: 4382753-4Report Type:Expedited (15-DaCompany Report #2004038283
Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Ineffective Feeling Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG, 3 IN 1 D) ORAL							

Hyoscyamine Sulfate (Hyoscymaine Sulfate)	C
Librax (Chlordiazepoxide Hydrochloride, Clidinium Bromide)	C
Primidone (Primidone)	C

Date:06/17/04ISR Number: 4382755-8Report Type:Expedited (15-DaCompany Report #2004033431
Age: Gender:Female I/FU:F

Outcome	PT
Other	Adrenal Neoplasm Cardiac Failure

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Doxepin (Doxepin) C
Salbutamol
(Salbutamol) C

Date:06/17/04ISR Number: 4382926-0Report Type:Expedited (15-DaCompany Report #001-0945-M0200490
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abdominal Pain	Consumer	Neurontin			
Other		Cholecystectomy		(Gabapentin)	PS		
		Completed Suicide		Zoloft (Sertraline)	SS		
75 MG		Drug Effect Decreased					
(DAILY),		Eating Disorder		Oxycodone			
		Gastrointestinal Disorder		Hydrocdhloride			
		Gun Shot Wound		(Oxycodone			
		Idiosyncratic Drug		Hydrochloride)	SS		
		Reaction					
		Pain					
		Self-Medication					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/17/04ISR Number: 4383157-0Report Type:Expedited (15-DaCompany Report #2004037986
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Amnesia Concussion	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D) ORAL		Condition Aggravated Fall Neck Mass Pain Urinary Incontinence		Atenolol (Atenolol) Esomeprazole (Esomeprazole) Vicodin (Hydrocodone Bitartrate, Paracetamol) Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C C C		

Date:06/18/04ISR Number: 4383100-4Report Type:Expedited (15-DaCompany Report #2004038419
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Brain Damage Feeling Abnormal Injury Loss Of Consciousness Memory Impairment Pain Pain In Extremity Sexual Assault Victim Syncope Vision Blurred	Consumer	Neurontin (Gabapentin) Geodon (Ziprasidone Hydrochloride) (Ziprasidone)	PS SS		

Date:06/18/04ISR Number: 4383147-8Report Type:Expedited (15-DaCompany Report #2004027799
Age:66 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Infection	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 (TID),		Pharmaceutical Product					
ORAL		Complaint		Gemfibrozil	C		

Date:06/18/04ISR Number: 4383148-XReport Type:Expedited (15-DaCompany Report #2004038421
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence	Consumer	Neurontin (Gabapentin)	PS		

Date:06/18/04ISR Number: 4383360-XReport Type:Expedited (15-DaCompany Report #2004217204TH
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 200 MG, BID,		Dizziness Melaena	Foreign Health	Celebrex (Celecoxib) Capsule	PS		ORAL
ORAL	7 DAY	Oedema Peripheral Shock	Professional Other	Neurontin			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

300 MG, BID,
 ORAL
 (Gabapentin) SS ORAL
 Neurobion C
 Brewer'S Yeast C
 (Yeast Dried) C
 Valium C

Date:06/18/04ISR Number: 4383380-5Report Type:Expedited (15-DaCompany Report #2004003994
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Developmental	Foreign	Neurontin			
Other		Coordination Disorder	Health	(Gabapentin)	PS		
2400 MG		Developmental Delay	Professional				
(TID),		Drug Exposure During					
PLACENTAL		Pregnancy					
		Mental Impairment					
		Speech Disorder					
		Developmental					

Date:06/18/04ISR Number: 4384184-XReport Type:Expedited (15-DaCompany Report #2004038659
 Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Condition Aggravated	Consumer	Neurontin			
		Rash		(Gabapentin)	PS		ORAL
ORAL		Scar		Omeprazole			
		Sensation Of Foreign Body		(Omeprazole)	C		
		Skin Infection		Paracetamol			
		Staphylococcal Infection		(Paracetamol)	C		
				Naproxen Sodium			
				(Naproxen Sodium)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other 900 MG (300 MG, 3 IN 1 D), ORAL		Back Pain Condition Aggravated Drug Ineffective Intervertebral Disc Degeneration	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Metformin Hydrochloride (Metformin Hydrochloride) Glipizide (Glipizide) Rosiglitazone Maleate (Rosiglitazone Maleate) Gemfibrozil (Germfibrozil)	C C C C		

Freedom Of Information (FOI) Report

Date:06/21/04ISR Number: 4391143-XReport Type:Periodic
Age:53 YR Gender:Female I/FU:I

Company Report #USA-2003-0011604

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553) (Oxycodone Hydrochloride) Hydrocodone Bitartrate (Similar To Ind 59,175) (Hydrocodone Bitartrate) Fentanyl (Fentanyl) Acetaminophen (Paracetamol) Caffeine (Caffeine) Meprobamate (Meprobamate) Promethazine (Promethazine) Trazodone (Trazodone) Alprazolam (Alprazolam) Citalopram (Citalopram) Desipramine (Desipramine) Imipramine (Imipramine) Lamotrigine (Lamotrigine) Nicotine (Nicotine) Carisoprodol (Carisoprodol) Dextromethorphan (Dextromethorphan) Gabapentin (Gabapentin)	PS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS		

Date:06/21/04ISR Number: 4391247-1Report Type:Periodic
Age:25 YR Gender:Female I/FU:F

Company Report #USA-2002-0000707

Outcome
Hospitalization -
Initial or Prolonged
Other

PT
Abdominal Pain
Agitation
Anxiety
Asthenia
Buttock Pain
Constipation
Cough
Decreased Appetite
Depression
Diarrhoea
Discomfort
Drug Abuser
Drug Dependence
Drug Exposure During
Pregnancy

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Ineffective Drug Withdrawal Syndrome Foot Deformity	Report Source	Product	Role	Manufacturer	Route
10 MG, SEE TEXT, ORAL		Fracture Nonunion Generalised Anxiety Disorder	Consumer Health Professional	Oxycontin Tablets 10 Mg (Oxycodone Hydrochloride)	PS		ORAL
		Hypertension	Other				
5 MG, ORAL		Hypoaesthesia Insomnia Joint Contracture		Oxyir Capsules 5 Mg (Oxycodone Hydrochloride)	SS		ORAL
INT (CONT)		Limb Deformity Motor Dysfunction Muscle Atrophy Muscle Spasms		Morphine Sulfate (Similar To Nda 19-516) (Morphine Sulfate)	SS		
300 MG, ORAL		Myalgia Nausea		Neurontin (Gabapentin)	SS		ORAL
1 TABLET, BID, ORAL		Neuropathic Pain Oedema Peripheral Pain Pain In Extremity		Lortab (Paracetamol, Hydrocodone Bitartrate)	SS		ORAL
		Peroneal Nerve Palsy Radiculopathy Rash Macular Rhinorrhoea Road Traffic Accident Scar Sciatica Sensory Loss Stress Tenderness Urinary Hesitation Vomiting Weight Decreased Weight Increased		Vicoprofen Hydrocodone W/Acetaminophen Relafen Doxepin Prochlorperazine Vioxx Ultram Ambien Penicillin V-K Dexamethasone Metronidazole Meprozine Promethazine Clonazepam Celebrex Depo Provera Thiamine Phenergan "Specia"	C C C C C C C C C C C C C C C C C C C C		

Zantac	C
Sudafed	C
Dolobid	C
Glucola (Does Not Code)	C
Tylox	C
Theragran	C
Desyrel	C
Dolophine	C
Hydrochloride	C

Date:06/21/04ISR Number: 4395357-4Report Type:Periodic Company Report #KII-2002-0007213
 Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dependence Medication Error	Health Professional Company	Oxycontin Tablets (Oxycodone Hydrochloride)	PS		
UNKNOWN	UNK UNK, UNK,		Representative				
UNKNOWN				Elavil (Amitriptyline)			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

UNKNOWN	25 MG, UNK,		Hydrochloride)	SS
UNKNOWN			Percocet (Paracetamol, Oxycodone Hydrochloride)	SS
UNKNOWN	325 MG, SEE			
TEXT,				
UNKNOWN			Neurontin (Gabapentin)	SS
UNKNOWN	300 MG, UNK,			
UNKNOWN				

Date:06/21/04ISR Number: 4395952-2Report Type:Periodic Company Report #USA-2003-0010893
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Accidental Overdose	Health	Oxycodone			
Other		Drug Abuser	Professional	Hydrochloride			
		Multiple Drug Overdose	Other	(Similar To Nda			
		Multiple Drug Overdose		20-553) (Oxycodone			
		Accidental		Hydrochloride)	PS		
		Overdose		Paroxetine			
				(Paroxetine)	SS		
				Gabpentin			
				(Gabapentin)	SS		
				Carisoprodol			
				(Carisoprodol)	SS		
				Meprobamate			
				(Meprobamate)	SS		

Date:06/21/04ISR Number: 4396063-2Report Type:Periodic Company Report #USA-2003-0011073
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Accidental Overdose	Health	Oxycontin	PS	ORAL
ORAL	Multiple Drug Overdose	Professional Other	Hydrocodone Bitartrate Alpazolam (Alpazolam) Diphenhydramine (Diphenhydramine) Acetaminophen (Paracetamol) Gemfibrozil (Gemfibrozil) Nadolol (Nadolol) Quetiapine (Quetiapine) Fentanyl (Fentanyl) Gabapentin (Gabapentin)	SS SS SS SS SS SS SS SS	

Date:06/21/04ISR Number: 4396097-8Report Type:Periodic Company Report #USA-2003-0011031
Age:38 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 (Similar To Nda			

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Freedom Of Information (FOI) Report

20-553) (Oxycodone Hydrochloride)	PS
Choline Magnesium Trisalicylate (Choline Magnesium Trisalicylate)	SS
Zonisamide (Zonisamide)	SS
Temazepam (Temazepam)	SS
Gabapentin (Gabapentin)	SS
Neurontin (Gabapentin)	SS
Celexa (Citalopram Hydrobromide)	SS
Remeron (Mirtazapine)	SS
Citalopram (Citalopram)	SS
Oxazepam (Oxazepam)	SS
Danazol	C

Date:06/22/04ISR Number: 4385916-7Report Type:Expedited (15-DaCompany Report #2004039225
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Confusional State	Consumer	Neurontin (Gapentin)	PS		
Other		Difficulty In Walking Disorientation Pain Psychotic Disorder					

Date:06/22/04ISR Number: 4385921-0Report Type:Expedited (15-DaCompany Report #2004039112
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (200 Other MG, 3 IN 1D)		Carpal Tunnel Syndrome Dizziness Drug Level Fluctuating	Health Professional	Neurontin (Gabapentin)	PS		

Eye Injury	Celecoxib	
Fall	(Celecoxib)	C
Fatigue	Warfarin Sodiun	
Feeling Abnormal	(Warfarin Sodium)	C
Gait Disturbance	Rabeprazole	
Head Injury	Sodium(Rabeprazole	
Mental Disorder	Sodium)	C
Somnolence	Ultracet	
	(Paracetamol,	
	Tramadol	
	Hydrochloride)	C
	Zolpidem Tartrate	
	(Zolpidem Tartrate)	C
	Estrogens Conjugated	
	(Estrogens	
	Conjugated_	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/22/04ISR Number: 4385923-4Report Type:Expedited (15-DaCompany Report #2004038647
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Difficulty In Walking	Consumer	Neurontin (Gabapentin)	PS		ORAL
3200 MG, ORAL		Electroencephalogram					
		Abnormal		Citalopram			
		Eye Movement Disorder		Hydrobromide			
		Impaired Driving Ability		(Citalopram			
		Speech Disorder		Hydrobromide)	C		
		Vision Blurred		Alprazolam			
				(Alprazolam)	C		
				All Other			
				Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	C		

Date:06/22/04ISR Number: 4386147-7Report Type:Expedited (15-DaCompany Report #2004039495
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Abasia	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Drug Ineffective					
ORAL		Pain		All Other			
				Therapeutic Products	C		

Date:06/22/04ISR Number: 4386148-9Report Type:Expedited (15-DaCompany Report #2004039114
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorexia	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300		Body Height Decreased					
MG, 2 IN 1		Impaired Driving Ability					
D), ORAL		Neoplasm Malignant					

Spinal Disorder
Weight Increased

Quinine Sulfate C
Nitrofurantoin C
Propranolol
Hydrochloride C
Fluoxetine C
Zolpidem Tartrate C
Morphine C
Vicodin (Hydrocodone
Bitartrate,
Paracetamol) C

Date:06/22/04ISR Number: 4386149-0Report Type:Expedited (15-DaCompany Report #2004038650

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Neurontin			
Other		Convulsion	Professional Company Representative	(Gabapentin) Valproate Semisodium	PS C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/22/04ISR Number: 4386232-XReport Type:Expedited (15-DaCompany Report #2004039144

Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Hypoaesthesia Paralysis	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL		Swelling	Professional				
				All Other Therapeutic Prodcuts (All Other Therapeutic Products)	C		

Date:06/22/04ISR Number: 4386233-1Report Type:Expedited (15-DaCompany Report #2004034949

Age:19 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Alkaline Phosphatase Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL		Liver Function Test Abnormal	Professional				
				Paracetamol (Paracetamol)	SS		ORAL
4 GRAM (1 GRAM, 4 IN 1 D), ORAL				Paroxetine (Paroxetine)	C		
				Trazodone (Trazodone)	C		
				Amitriptyline (Amitriptyline)	C		
				Trimethoprim (Trimethoprim)	C		

Date:06/23/04ISR Number: 4386351-8Report Type:Expedited (15-DaCompany Report #2004037988
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Health	Neurontin			
		Condition Aggravated	Professional	(Gabapentin)	PS		
800 MG		Convulsion					
		Ischaemia					

Date:06/23/04ISR Number: 4386988-6Report Type:Expedited (15-DaCompany Report #2004040068
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Consumer	Neurontin			
		Cardiac Disorder		(Gabapentin)	PS		
		Chest Pain					
		Circulatory Collapse					
		Dizziness					
		Headache					
		Loss Of Consciousness					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/24/04ISR Number: 4384808-7Report Type:Expedited (15-DaCompany Report #SIE-FLU-002

Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Abscess Abdominal Pain Asterixis Blood Albumin Decreased Blood Potassium Decreased Cognitive Disorder	Literature Health Professional Other	Kadian (Morphine Sulfate Sustained Release) Capsules, 20mg (Alpharma Pharmaceuticals Inc)	PS	Alpharma Pharmaceuticals Inc	ORAL
30 MG, AS NEEDED, ORAL		Confusional State					
0.5 MG, AS NEEDED		Consciousness Fluctuating Constipation Dehydration		Alprazolam Tablets Usp, 0.5 Mg (Purepac)	SS	Purepac	
100 MG, NIGHTLY		Delirium					
TRANSDERMAL EVERY THREE DAYS, TRANSDERMAL	125 MCG/HR,	Delusion Disorientation Disturbance In Attention Infection		Trazodone Hydrochloride Tablets, 100 Mg (Purepac)	SS	Purepac	
80 MG, DAILY 300 MG, THREE TIMES DAILY		Lack Of Spontaneous Speech Mental Impairment Metabolic Disorder Myoclonus		Fentanyl (Janssen)	SS	Janssen	
		Nausea Perseveration		Fluoxetine (Siegfried)	SS	Siegfried	
		Pyrexia		Gabapentin (Pfizer)	SS	Pfizer	
		Sedation					
		Small Intestinal Obstruction Thinking Abnormal					

Date:06/24/04ISR Number: 4388508-9Report Type:Expedited (15-DaCompany Report #2004039325
Age:52 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Confusional State	Foreign	Neurontin			
Initial or Prolonged	Myoclonus	Health	(Gabapentin)	PS		ORAL
ORAL	Pericardial Effusion	Professional				

Date:06/24/04ISR Number: 4388592-2Report Type:Expedited (15-DaCompany Report #2004021934
Age:84 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Abnormal Behaviour	Foreign	Neurontin			
Hospitalization -	Anorexia	Health	(Gabapentin)	PS		
1800 MG (3 IN						
Initial or Prolonged	Anxiety	Professional				
1 D)						
	Cerebellar Syndrome	Company	Raloxifene			
	Depressed Level Of	Representative	Hydrochloride			
	Consciousness		(Raloxifene			
	Drug Intolerance		Hydrochloride)	SS		ORAL
ORAL						
	Drug Toxicity		Indapamide			
	Insomnia		(Indapamide)	C		
	Pain In Extremity		Trimetazidine			
	Restless Legs Syndrome		Hydrochloride			
	Suicidal Ideation		(Trimetazidine			
			Hydrochloride)	C		
			Ibuprofen			
			(Ibuprofen)	C		
			Paracetamol			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Paracetamol) C
 Aldactazine
 (Altizide,
 Spironolactone) C
 Rabeprazole Sodium
 (Rabeprazole Sodium) C

Date:06/24/04ISR Number: 4388682-4Report Type:Expedited (15-DaCompany Report #2004039248
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Renal Impairment	Consumer	Neurontin (Gabapentin)	PS		

Date:06/24/04ISR Number: 4389091-4Report Type:Expedited (15-DaCompany Report #2004038659
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Localised Infection Pharmaceutical Product	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Complaint Pharyngeal Neoplasm Benign Scar Skin Infection Staphylococcal Infection		Omeprazole (Omeprazole) Paracetamol (Paracetamol) Naproxen Sodium (Naproxen Sodium)	C C C		

Date:06/25/04ISR Number: 4387442-8Report Type:Expedited (15-DaCompany Report #2004038392
 Age:83 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Fatigue	Foreign	Neurontin			
Hospitalization - 1800 MG		Heart Rate Increased	Health	(Gabapentin)	PS		
Initial or Prolonged		Heart Rate Irregular Jaundice	Professional Company Representative	Oxcarbazepine (Oxcarbazepine)	C		

Outcome	PT
Life-Threatening	Alanine Aminotransferase
Hospitalization -	Increased
Initial or Prolonged	Alpha 1 Globulin
Other	Increased
	Aspartate
	Aminotransferase
	Increased
	Blood Alkaline
	Phosphatase Increased
	Blood Lactate
	Dehydrogenase Increased
	Circulatory Collapse
	Dizziness
	Gamma-Glutamyltransferase
	Increased
	Hepatocellular Damage
	Nausea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Restlessness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1200 MG DAILY		Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Company Representative	Zinc	C		

Date:06/25/04ISR Number: 4387672-5Report Type:Expedited (15-DaCompany Report #2004040168
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (300 MG, 2 IN 1 D), ORAL		Convulsion Dizziness Dyspnoea Hypertension	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
				All Other Therapeutic Products	C		

Date:06/25/04ISR Number: 4387677-4Report Type:Expedited (15-DaCompany Report #2004031169
 Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (300 MG, 1 IN 1 D), ORAL		Condition Aggravated Diabetes Mellitus Insomnia Muscle Spasms Neuropathic Pain Pain In Extremity	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
				Insulin (Insulin)	C		
				Fluoxetine (Fluoxetine)	C		
				Levothyroxine			

Date:06/25/04ISR Number: 4388196-1Report Type:Expedited (15-DaCompany Report #2004038419
Age:48 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Abasia	Consumer	Neurontin			
Initial or Prolonged	Bipolar Disorder		(Gabapentin)	PS		
Other	Brain Damage		Geodon (Ziprasidone			
	Dizziness		Hydrochloride)			
	Feeling Abnormal		(Ziprasidone)	SS		
	Homeless					
	Pain					
	Pain In Extremity					
	Syncope					
	Victim Of Crime					
	Vision Blurred					

Date:06/25/04ISR Number: 4388200-0Report Type:Expedited (15-DaCompany Report #2004038007
Age: Gender:Male I/FU:F

Outcome
Death
Hospitalization -
Initial or Prolonged

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Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Abasia	Consumer	Neurontin			
		Apallic Syndrome	Health	(Gabapentin)	PS		
		Aphagia	Professional				
		Aphasia					
		Gait Disturbance					
		General Physical Health					
		Deterioration					
		Impaired Driving Ability					
		Mental Impairment					
		Metastases To Central					
		Nervous System					
		Paralysis					

Date:06/25/04ISR Number: 4388207-3Report Type:Expedited (15-DaCompany Report #2004024140

Age:83 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aura	Consumer	Dilantin Suspension			
Other		Body Height Decreased	Health	(Phenytoin Sodium)	PS		
		Breast Cancer	Professional	Neurontin			
		Drug Ineffective		(Gabapentin)	SS		
2400 MG (800		Dyspnoea					
MG, 3 IN 1 D)		Fatigue		Phenobarbital	SS		
		Muscle Spasms		Metformin			
		Post Procedural		Hydrochloride			
		Complication		(Metformin			
		Radiation Fibrosis - Lung		Hydrochloride)	C		
		Somnolence		Pioglitazone			
				(Pioglitazone)	C		
				Celecoxib			
				(Celecoxib)	C		
				Tolterodine			
				L-Tartrate			
				(Tolterodine			
				L-Tartrate)	C		
				Chondroitin Sulfate			
				(Chondroitin			
				Sulfate)	C		

Glucosamine	
(Glucosamine)	C
Multivitamins	
(Ascorbic	
Acid/Ergocalciferol/	
Folic	
Acid/Nicotinamide/Pa	C
Tocopherol	
(Tocopherol)	C
Calcium With Vitamin	
D (Calcium	
Phosphate, Calcium	
Sodium Lacatte,	
Ergocalciferol)	C
Tamoxifen	
(Tamoxifen)	C

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Date:06/25/04 ISR Number: 4388213-9 Report Type:Expedited (15-DaCompany Report #2004022432
 Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Amnesia Anaemia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
7200 MG (600 MG, 3 IN 1 D), ORAL		Astigmatism Constipation Depression Dysarthria Endometriosis Fatigue Fibromyalgia Insomnia Libido Decreased Muscle Spasms Muscle Twitching Myalgia Myositis Visual Disturbance	Professional	Propacet (Dextropropoxyphene Napsilate, Paracetamol) Estratest Hs (Estrogens Esterified, Methyltestosterone) Paroxetine Hydrochloride (Paroxetine Hydrochloride) Nortriptyline (Nortriptyline) Buspirone Hydrochloride (Buspirone Hydrochloride) Clonazepam (Clonazepam) Dextropropoxyphene Hydrochloride (Dextropropoxyphene Hydrochloride) Laxatives (Laxatives) Calcium Carbonate (Calcium Carbonate) Becosym Forte (Nicotinamide, Pyridoxine Hydrochloride, Riboflavin, Thiamine Multivitamins (Ascorbic	C C C C C C		

Acid/Ergocalciferol/
 Folic
 Acid/Nicotinamide/Pa C
 Citalopram
 Hydrochloride
 (Citalopram
 Hydrochloride) C
 All Other
 Therapeutic Products C

Date:06/25/04ISR Number: 4388218-8Report Type:Expedited (15-DaCompany Report #2004030834

Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
ORAL		Drug Level Increased		Levothyroxine Sodium (Levothyroxine			

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Sodium)	C
Tocopherol	
(Tocopherol)	C
Biotin (Biotin)	C
Esomeprazole	
(Esomeprazole) C	C

Date:06/25/04ISR Number: 4388220-6Report Type:Expedited (15-DaCompany Report #2004033139

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Asthenia Bedridden	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600						
Other MG, 3 IN 1	Blood Test Abnormal	Professional				
D), ORAL	Hypotension					
	Interstitial Lung Disease		Rofecoxib			
	Mobility Decreased		(Rofecoxib)	SS		
	Paralysis		Simvastatin			
	Pneumonitis		(Simvastatin)	C		
	Respiratory Disorder		Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)			
			Pantoprazole	C		
			(Pantoprazole)	C		
			Montelukast Sodium			
			(Montelukast Sodium)	C		
			Salbutamol			
			(Salbutamol)	C		
			Fluoxetine			
			Hydrochloride			
			(Fluoxetine Hydrochloride)	C		

Date:06/25/04ISR Number: 4388259-0Report Type:Expedited (15-DaCompany Report #2004040065

Age:60 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

Other	Oedema Peripheral	Consumer	Neurontin		
	Pain In Extremity		(Gabapentin)	PS	ORAL
ORAL	Poor Peripheral		Lisinopril	C	
	Circulation		Escitalopram	C	
	Post Procedural		Azelastine	C	
	Complication		Risedronate Sodium	C	
	Spinal Fracture		Trazodone	C	
	Swelling				
	Weight Increased				

Date:06/28/04ISR Number: 4385608-4Report Type:Expedited (15-DaCompany Report #200411919EU
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated		Lantus	PS	Aventis	
		Diabetes Mellitus				Pharmaceuticals Inc.	
SUBCUTANEOUS	Dose unit:						
		Drug Interaction					
International							
units per							
millilitre				Neurontin	SS		ORAL

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SUBCUTANEOUS	Dose unit:	Insulatard	SS	
International				
units per				
millilitre				
SUBCUTANEOUS	Dose unit:	Novorapid	SS	
International				
units per				
millilitre				
		Pantoprazol	C	ORAL
		Acetylsalicylic Acid	C	ORAL

Date:06/28/04ISR Number: 4385615-1Report Type:Expedited (15-DaCompany Report #PHEH2004US06560
 Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening Hospitalization - 6 mg, BID	Abdominal Pain Agitation	Health Professional	Zelnorm	PS	Novartis Sector: Pharma	ORAL
Initial or Prolonged 75 mg, QD	Autonomic Nervous System		Effexor-Xr	SS		
Other 75 mg/qam,	Imbalance		Effexor-Xr	SS		
150 mg/qhs	Blood Creatine					
5 mg, QHS	Phosphokinase Increased Coma		Mirapex "Boehringer Ingelheim"	SS		
25 mg, BID	Electrocardiogram St		Quetiapine	SS		
UNK, TID	Segment Abnormal Electromechanical		Mirtazapine Sinemet	SS SS		
	Dissociation		Duragesic	SS		
	Heart Injury		Neurontin	SS		
	Hypertension		Pantoprazole	C		
	Hypotension		Advair	C		

Hypoxia	Lipitor	C
Intubation	Furosemide	C
Leukocytosis	Lantus	C
Lung Infiltration	Plavix	C
Mental Status Changes	Imdur	C
Metabolic Acidosis	Allegra	C
Myocardial Infarction	Atenolol	C
Myoclonus		
Opisthotonus		
Pneumonia		
Pneumonia Aspiration		
Pulmonary Oedema		
Renal Failure		
Renal Tubular Necrosis		
Respiratory Failure		
Restless Legs Syndrome		
Rhabdomyolysis		
Serotonin Syndrome		
Troponin Increased		

Date:06/28/04ISR Number: 4389067-7Report Type:Expedited (15-DaCompany Report #04P-056-0263854-00
Age:37 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Abdominal Pain Upper
Required	Abdominal Tenderness
Intervention to	Alanine Aminotransferase
Prevent Permanent	Increased
Impairment/Damage	Aspartate

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Dose	Duration	Adverse Event	Report Source	Product	Role	Manufacturer	Route
		Aminotransferase Increased					
		Asthenia					
		Blood Alkaline					
3 CAPSULE, 2 IN 1 D, PER ORAL		Phosphatase Increased Drug Intolerance Dysaesthesia Folate Deficiency	Foreign Other	Kaletra Soft Gelatin Capsules (Kaletra) (Lopinavir/Ritonavir)	PS		ORAL
		Gamma-Glutamyltransferase Increased					
		Hepatocellular Damage		Ritonavir	SS		
		Hyperaesthesia		Didanosine	SS		
		Hyperlactacidaemia		Stavudine	SS		
		Neuralgia		Indinavir	SS		
		Neutropenia		Nelfinavir Mesilate	SS		
		Oedema		Doxorubicin			
INTRAVENOUS D, INTRAVENOUS	35 MG, 1 IN 1	Pain		Hydrochloride	SS		
		Pancreatic Disorder					
		Pancytopenia					
		Pulmonary Hypertension		Zidovudine			
		Rash		W/Lamivudine	SS		
		Renal Failure		Gabapentin	SS		
245 MG, 1 IN 1 D, PER ORAL		Renal Failure Chronic		Viread	SS		ORAL
		Transaminases Increased					
				Abacavir Sulfate	SS		ORAL
300 MG, 2 IN 1 D, PER ORAL				Ondansetron Hydrochloride	SS		ORAL
PER ORAL				Motilium	SS		ORAL
PER ORAL				Zalcitabine	C		
				Lamivudine	C		
				Efavirenz	C		
				Idoxuridine	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG (600 Other MG, 3 IN 1 D)	Bronchostenosis Gastrooesophageal Reflux Disease Odynophagia Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		ORAL
,ORAL			Olanzapine (Olanzapine)	SS		ORAL
ORAL			Aripiprazole (Aripiprazole)	SS		ORAL
ORAL			Valproate Semisodium Paroxetine Hydrochloride Clonazepam Diphenhydramine Hydrochloride	C C C C		

Outcome	PT
Other	Gait Disturbance Mental Impairment Miosis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pupillary Reflex Impaired Tremor	Health Professional	Neurontin (Gabapentin)	PS		
DAILY							

Date:06/28/04ISR Number: 4389830-0Report Type:Expedited (15-DaCompany Report #2004040186
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Arthralgia Dysgraphia	Consumer	Neurontin (Gabapentin)	PS		ORAL
	600 MG (300 MG, 2 IN 1 D), ORAL		Gait Disturbance Hypoaesthesia					
	30 MG (30 MG) ORAL		Pain Pain In Extremity		Procardia Xl (Nifedipine)	SS		ORAL
	200 MG (200 MG, 1 IN 1 D)		Spinal Cord Disorder		Celebrex (Celecoxib) (Celecoxib)	SS		
					All Other Therapeutic Products	C		

Date:06/28/04ISR Number: 4389833-0Report Type:Expedited (15-DaCompany Report #2004040264
Age:36 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Drug Dependence Drug Withdrawal Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
	3200 MG, ORAL		Feeling Abnormal		Antihypertensives All Other	C		

Date:06/28/04ISR Number: 4389835-0Report Type:Expedited (15-DaCompany Report #2004027664

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Atrial Fibrillation	Consumer	Neurontin (Tablets)			
Other		Cardiac Operation		(Gabapentin)	PS		ORAL
2400 MG (800		Cardiac Pacemaker					
MG, 3 IN 1		Insertion					
D), ORAL		Decreased Appetite		Donepezil			
		Dementia Alzheimer'S Type		Hydrochloride			
		Diarrhoea		(Donepezil			
		Dizziness		Hydrochloride)	SS		ORAL
10 M (10 MG,		Hypoglycaemia					
1 IN 1 D),		Insomnia					
ORAL		Myocardial Infarction		Carbamazepine			
		Tricuspid Valve		(Carbamazepine)	C		
		Incompetence		Warfarin Sodium			
				(Warfarin Sodium)	C		
				Amiodarone			
				Hydrochloride			
				(Amiodarone			
				Hydrochloride)	C		
				Digoxin (Digoxin)	C		
				Furosemide			
				(Furosemide)	C		

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Isosorbide
 Mononitrate
 (Isosorbide
 Mononitrate) C
 Potassium Chloride
 (Potassium Chloride) C
 Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C
 Centrum (Minerals
 Nos, Vitamins Nos) C
 Atorvastatin
 (Atorvastatin) C
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C
 Magnesium
 (Magnesium) C
 Zinc (Zinc) C
 Metoprolol
 (Metoprolol) C
 Enalapril Maleate
 (Enalapril Maleate) C

Date:06/28/04ISR Number: 4389998-8Report Type:Expedited (15-DaCompany Report #2004033874
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Dental Caries	Consumer	Neurontin (Tablets)			
Other		Tooth Discolouration	Health	(Gabapentin)	PS		ORAL
1800 MG (600		Tooth Disorder	Professional				
MG, 3 IN 1		Tooth Fracture					
D), ORAL		Toothache		Insulin Lispro (Insulin Lispro) Humalog Mix 25 (Insulin Lispro, Insulin Lispro Protamine Suspension)	C		

Nateglinide	
(Nateglinide)	C
All Other	
Therapeutic Products	
(All Other	
Therapeutic	
Products)	C
Carvedilol	
(Carvedilol)	C
Ezetimibe	
(Ezetimibe)	C
Ramipril (Ramipril)	C
Tolterodine	
L-Tartrate	
(Tolterodine	
L-Tartrate)	C
Venlafaxine	
Hydrochloride	
(Venlafaxine	

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Hydrochloride)	C
Furosemide	
(Furosemide)	C
Atorvastatin	
(Atorvastatin)	C
Loperamide	
Hydrochloride	
(Loperamide	
Hydrochloride)	C
Rosiglitazone	
Maleate	
(Rosiglitazone	
Maleate)	C
Fexofenadine	
Hydrochloride	
(Fexofenadine	
Hydrochloride)	C
Multivitamins	
(Ascorbic Acid,	
Ergocalciferol,	
Folic Acid,	
Nicotinamide,	C
Valaciclovir	
Hydrochloride	
(Valaciclovir	
Hydrochloride)	C
Isosorbide	
Mononitrate	
(Isosorbide	
Mononitrate)	C
Montelukast Sodium	
(Montelukast Sodium)	C
Esomeprazole	
(Esomeprazole)	C
Oxycocet (Oxycodone	
Hydrochloride,	
Paracetamol)	C
Lactulose	
(Lactulose)	C
Nystatin (Nystatin)	C
Tramadol	
Hydrochloride	
(Tramadol	
Hydrochloride)	C
Mirtazapine	
(Mirtazapine)	C
Metolazone	
(Metolazone)	C

Olanzapine	
(Olanzapine)	C
Amitriptyline	
(Amitriptyline)	C
Bupropion	
Hydrochloride	
(Bupropion	
Hydrochloride)	C
Dimeticone,	
Activated	
(Simeticone)	C

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Freedom Of Information (FOI) Report

Date:06/28/04ISR Number: 4390321-3Report Type:Expedited (15-DaCompany Report #2004039627
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 700 MG (700 MG , 1 IN 1 D), ORAL		Facial Spasm Pharmaceutical Product Complaint Tic Vitreous Floaters	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL

Date:06/28/04ISR Number: 4390323-7Report Type:Expedited (15-DaCompany Report #2004033685
Age:43 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other ORAL		Completed Suicide Intentional Misuse	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:06/28/04ISR Number: 4390332-8Report Type:Expedited (15-DaCompany Report #2004040267
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Other 25 MG (2 IN 1 D)		Abdominal Pain Upper Abnormal Behaviour Condition Aggravated Drug Ineffective Drug Interaction Drug Withdrawal Syndrome Feeling Abnormal Hallucination Medication Error Mental Disorder Pyrexia	Consumer	Neurontin (Gabapentin) Diazepam (Diazepam) Baclofen (Baclofen) All Other Therapeutic Products (All Other Therapeutic Products) Oxybutynin	PS SS SS SS		ORAL

Hydrochloride (Oxybutynin Hydrochloride)	C
Warfarin Sodium (Warfarin Sodium)	C
Potassium (Potassium)	C
Fluticasone Propionate (Fluticasone Propionate)	C
Combivent (Ipratropium Bromide, Salbutamol Sulfate)	C

Date:06/28/04ISR Number: 4390342-0Report Type:Expedited (15-DaCompany Report #2004028513

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dysgeusia Pharmaceutical Product	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1600 (4 IN 1 D), ORAL		Complaint		Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		

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Cetirizine
 Hydrochloride
 (Cetirizine
 Hydrochloride) C
 Lidocaine
 (Lidocaine) C
 Lactulose
 (Lactulose) C
 Salmeterol Xinafoate
 (Salmeterol
 Xinafoate) C
 Doxepin (Doxepin) C
 Salbutamol
 (Salbutamol) C

Date:06/30/04ISR Number: 4388113-4Report Type:Direct
 Age:76 YR Gender:Female I/FU:I

Company Report #CTU 221827

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Discomfort		Neurontin 800 Mg			
Other		Dizziness		Pzifer	PS	Pzifer	
4/DAILY		Pain					
		Pharmaceutical Product					
		Complaint					
		Tinnitus					

Date:06/30/04ISR Number: 4389923-XReport Type:Expedited (15-DaCompany Report #2004041539
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Impaired Insulin	Foreign	Neurontin (Tablets)			
Initial or Prolonged		Secretion	Consumer	(Gabapentin)	PS		ORAL
600 MG (300		Pancreas Transplant					
MG, 2 IN 1							
D), ORAL							

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged (600 MG, UNKNOWN), ORAL	Coma Condition Aggravated Dizziness Nausea Trigeminal Nerve Disorder	Foreign Consumer	Neurontin (Tablets) (Gabapentin) Carbamazepine (Carbamazepine)	PS C		ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Difficulty In Walking Fall	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:06/30/04ISR Number: 4390005-1Report Type:Expedited (15-DaCompany Report #2004040828

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

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG (400 MG, 1 IN 1 D), ORAL		Condition Aggravated Diabetes Mellitus	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Drug Interaction	Professional				
		Trigeminal Neuralgia					
SUBCUTANEOUS (UNKNOWN), SUBCUT	100 IE/ML			Insulin Glargine (Insulin Glargine)	SS		
SUBCUTANEOUS (UNKNOWN), SUBCUT	100 IE/ML			Insulin Injection, Isophane (Insulin Injection, Isophane)	SS		
SUBCUTANEOUS (UNKNOWN), SUBCUT	100 IE/ML			Insulin Aspart (Insulin Aspart)	SS		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Pantoprazole (Pantoprazole)	C		

Date:06/30/04ISR Number: 4390536-4Report Type:Expedited (15-DaCompany Report #2004041566

Age:71 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other	Back Pain	Consumer	Neurontin (Tablets)		
1600 MG (800	Drug Effect Decreased		(Gabapentin)	PS	ORAL
MG, 2 IN 1D),	Pharmaceutical Product				
ORAL	Complaint				
	Wrist Surgery		Atorvastatin		

(Atorvastatin)	C
Metoprolol	
(Metoprolol)	C
Esomeprazole	
(Esomeprazole)	C
Tocopherol	
(Tocopherol)	C

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		Drug Hypersensitivity	Consumer	Neurontin (Tablets)			
2400 MG (800		Muscle Spasms		(Gabapentin)	PS		ORAL
MG, 3 IN 1		Nervous System Disorder					
D), ORAL		Pharmaceutical Product					
		Complaint		Bupropion			
		Tremor		Hydrochloride			
		Weight Increased		(Bupropion			
				Hydrochloride)	SS		
				Venlafaxine			
				Hydrochloride			
				(Venlafaxine			
				Hydrochloride)	C		

Freedom Of Information (FOI) Report

Levothyroxine Sodium
 (Levothyroxine Sodium) C
 Baclofen (Baclofen) C

Date:06/30/04ISR Number: 4390684-9Report Type:Expedited (15-DaCompany Report #2004041530
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG (300 Other MG, 3 IN 1 D)	Abdominal Pain Abdominal Wall Abscess	Literature Health	Gabapentin (Gabapentin)	PS		
30 MG (30 MG, PRN) ORAL		Asterixis Blood Albumin Decreased Blood Potassium Decreased	Professional	Morphine Sulphate (Morphne Sulphate)	SS		ORAL
0.5 MG (PRN)		Cognitive Disorder Confusional State Constipation		Alprazolam (Alprazolam)	SS		
100 MG (NIGHTLY)		Decreased Activity Dehydration		Trazodone (Trazodone)	SS		
TRANSDERMAL (EVERY 3 DAYS),	125 MCG/H	Delusion Disorientation Disturbance In Attention		Fentanyl (Fentanyl)	SS		
TRANSDERMAL 80 MG (DAILY)		Mental Impairment Myoclonus Nausea Pain Sedation Thinking Abnormal		Fluoxetine (Fluoxetine)	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cellulitis Depression	Consumer	Neurontin (Gabapentin)	PS		
300 MG		Dizziness Feeling Abnormal Muscle Spasms Pain Vomiting					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Cholesterol Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 1 IN 1 D), ORAL		Flatulence Myocardial Infarction					
ORAL		Paraesthesia Tinnitus		Lipitor (Atorvastatin)	SS		ORAL
				Lisinopril (Lisinopril)	SS		
				Rosuvastatin (Rosuvastatin)	SS		
				Simvastatin (Simvastatin)	SS		
10 MG (10 MG, 1 IN 1 D)							

Freedom Of Information (FOI) Report

Pyridoxine Hydrochloride (Pyridoxine Hydrochloride)	SS
Ezetimibe (Ezetimibe)	C
All Other Therapeutic Products (All Other Therapeutic Products)	C
Doxepin (Doxepin)	C
Estrogens (Estrogens)	C
Fish Oil (Fish Oil)	C
Vitamins (Vitamins)	C
Minerals Nos (Minerals Nos)	C
Vicon Forte (Folic Acid, Minerals Nos, Vitamins Nos)	C
Retinol (Retinol)	C
Tocopherol (Tocopherol)	C
Potassium (Potassium)	C
Calcium Citrate (Calcium Citrate)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C

Date:06/30/04ISR Number: 4390921-0Report Type:Expedited (15-DaCompany Report #2004028587
Age:80 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - ORAL	Anaemia	Consumer	Neurontin	PS		ORAL
Initial or Prolonged Other	Faeces Discoloured Gastrointestinal Haemorrhage	Health Professional	Simvastatin (Simvastatin) Gliclazide (Gliclazide) Lansoprazole (Lansoprazole)	C C C		

Latanoprost
(Latanoprost) C
Cosopt (Dorzolamide
Hydrochloride,
Timolol Maleate) C
Amitriptyline
Hydrochloride
(Amitriptyline
Hydrochloride) C

Date:06/30/04ISR Number: 4390923-4Report Type:Expedited (15-DaCompany Report #2004034717
Age:83 YR Gender:Female I/FU:F

Outcome
Hospitalization -
Initial or Prolonged

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Freedom Of Information (FOI) Report

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route		
600 MG (300 MG, 2 IN 1 D), ORAL		Bone Infection	Consumer	Neurontin (Gabapentin)	PS		ORAL		
		Leukopenia							
		Neutropenia							
		Osteoarthritis							
		Osteoporosis						Bactrim (Sulfamethaxozole, Trimethoprim)	SS
		Procedural Complication						Levothyroxine (Levothyroxine)	C
								Ketorolac (Ketorolac)	C
								Dalteparin (Dalteparin)	C
								Vicodin (Hydrocodone Bitartrate, Paracetamol)	C
								Lomotil (Atropine Sulfate, Diphenoxylate Hydrochloride)	C

Date:06/30/04ISR Number: 4411148-XReport Type:Periodic
Age:48 YR Gender:Female I/FU:F

Company Report #03P-163-0243402-00

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthralgia	Consumer	Humira 40 Mg/ 0.8 ML Pre-Filled Syringe (Humira) (Adalimumab)	PS		
		Hypotrichosis					
		Photopsia					
		Visual Disturbance					
SUBCUTANEOUS	40 MG, 1 IN 2						
Wk,							
SUBCUTANEOUS							
PER ORAL				Gabapentin	SS		ORAL

Prednisone	C
Esomeprazole	C
Oxycodone	
Hydrochloride	C
Clonazepam	C

Date:06/30/04ISR Number: 4415366-6Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2003031978

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Abnormal Dreams	Consumer	Zoloft (Sertraline)	PS		
Initial or Prolonged	Anxiety		Neurontin			
Other	Blood Pressure Decreased		(Gabapentin)	SS		
	Bradycardia		Alprazolam			
	Diabetes Mellitus		(Alprazolam)	SS		
	Drug Interaction					
	Heart Rate Increased					
	Insomnia					
	Neuropathy					
	Nightmare					
	Panic Attack					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/01/04ISR Number: 4388415-1Report Type:Expedited (15-DaCompany Report #FR-ROCHE-371766
Age:76 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 9 DAY Initial or Prolonged	Toxic Skin Eruption		Rivotril	PS	Roche	ORAL
15 DAY			Tinzaparine Sodique	SS		ORAL
			Neurontin	SS		ORAL
2 DAY			Neurontin	SS		ORAL
			Enoxaparine	SS		ORAL
			Tardyferon B9	C		
			Naftidrofuryl	C		ORAL
			Depakine Chrono	C		ORAL

Date:07/01/04ISR Number: 4391721-8Report Type:Expedited (15-DaCompany Report #2004042649
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2700 MG (900 Other MG, 3 IN 1 D), ORAL	Accidental Overdose Balance Disorder Memory Impairment Movement Disorder Vision Blurred	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL			Carbamazepine (Carbamazepine)	SS		ORAL
			Potassium Chloride (Potassium Chloride)	C		
			Simvastatin (Simvastatin)	C		
			Furosemide (Furosemide)	C		
			Levothyroxine Sodium (Levothyroxine Sodium)	C		
			Baclofen (Baclofen)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour	Health	Neurontin			
Hospitalization - 900 MG (300 Initial or Prolonged MG, 3 IN 1 D), ORAL		Agitation	Professional	(Gabapentin)	PS		ORAL
		Delirium					
		Depressed Level Of Consciousness		Warfarin Sodium (Warfarin Sodium)	C		
		Fluid Intake Reduced		Spirolactone (Spirolactone)	C		
		Hypersomnia		Lisinopril (Lisinopril)	C		
		Moaning		Levothyroxine Sodium (Levothyroxine Sodium)	C		
		Oral Intake Reduced		Metoprolol Tartrate (Metoprolol Tartrate)	C		
		Sedation		Colchicine (Colchicine)	C		
		Urinary Tract Infection		Magnesium Gluconate (Magnesium Gluconate)	C		
				Calcium (Calcium)	C		

Freedom Of Information (FOI) Report

Pantoprazole
 (Pantoprazole) C
 Torasemide
 (Torasemide) C
 Insulin (Insulin) C

Date:07/02/04ISR Number: 4392032-7Report Type:Expedited (15-DaCompany Report #2012272
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Consumer	Oxycontin			
Hospitalization - Initial or Prolonged		Asthenia	Health	Tablets(Oxycodone Hydrochloride) Cr			
Other		Back Pain	Professional				
10 MG, BID Required		Drug Abuser	Other	Tablet	PS		
; 20 MG ; Intervention to 40 MG		Drug Toxicity					
Prevent Permanent Impairment/Damage		Erectile Dysfunction					
		Haematemesis		Diazepam(Diazepam)	SS		
		Haemorrhage		Oxazepam(Oxazepam)	SS		
		Headache		Temazepam(Temazepam)	SS		
		Iron Deficiency Anaemia		Lorazepam(Lorazepam)	SS		
		Mallory-Weiss Syndrome		Cannaboids(Cannabis)	SS		
		Respiratory Arrest		Diphenhydramine			
		Tachycardia		Hydrochloride(Diphen hydramine			
		Tooth Disorder		Hydrochloride)	SS		
		Tooth Extraction		Gabapentin(Gabapenti n)	SS		
				Claritin(Loratadine)	C		
				Zoloft(Sertraline Hydrochloride)	C		
				Altace(Ramipril)	C		
				Allopurinol	C		
				Depakote(Valproate Semisodium)	C		
				Soma (Carisoprodol)	C		
				Relafen (Nabumetone)	C		

Date:07/02/04ISR Number: 4392297-1Report Type:Expedited (15-DaCompany Report #2004042647
 Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Depression Insomnia	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL		Somnolence Suicidal Ideation Vision Blurred		Levothyroxine Sodium (Levothyroxine Sodium) Valsartan (Valsartan)	C C		

Date:07/02/04ISR Number: 4393857-4Report Type:Expedited (15-DaCompany Report #2004042632
Age: Gender:Male I/FU:I

Outcome	PT
Other	Central Nervous System Stimulation Feeling 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
ORAL		Hiv Test Positive Nervousness Sedation Tremor	Health Professional	Neurontin (Gabapentin) All Other Therapeutic Products	PS C		ORAL

Date:07/02/04ISR Number: 4417248-2Report Type:Expedited (15-DaCompany Report #2004048573
Age:2 DY Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death Neonatal Drug Exposure During Pregnancy	Consumer	Neurontin (Gabapentin) Oxcarbazepine (Oxcarbazepine) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Nadolol (Nadolol) Panadeine Co (Codeine Phosphate, Paracetamol)	PS SS C C C		PLACENTAL PLACENTAL

Date:07/06/04ISR Number: 4394499-7Report Type:Expedited (15-DaCompany Report #2004041566
Age:71 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Back Pain Blood Cholesterol Increased Condition Aggravated Drug Ineffective	Consumer	Neurontin (Tablets) (Gabapentin) Lipitor	PS		ORAL

10 MG (10 MG, 1 IN 1 D), ORAL	Feeling Abnormal Muscle Disorder Pharmaceutical Product Complaint	(Atorvastatin)	SS	ORAL
		Metoprolol (Metoprolol)	C	
		Esomeprazole (Esomeprazole)	C	
		Tocopherol (Tocopherom)	C	

Date:07/06/04ISR Number: 4394619-4Report Type:Expedited (15-DaCompany Report #2004041837
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Other	Anticonvulsant Drug Level Increased Depressed Level Of Consciousness	Consumer	Dilantin Suspension (Phenytoin Sodium) Neurontin (Gabapentin)	PS SS		
1200 MG (400 MG, 3 IN 1 D)	Feeling Abnormal Headache Ill-Defined Disorder Lethargy Pain Platelet Count Decreased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/06/04ISR Number: 4394622-4Report Type:Expedited (15-DaCompany Report #2003030361

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis	Consumer	Neurontin			
ORAL		Bipolar Disorder		(Gabapentin)	PS		ORAL
		Drug Hypersensitivity		Venlafaxine			
		Overweight		Hydrochloride			
		Tongue Geographic		(Venlafaxine			
				Hydrochloride)	SS		
				Anti-Diabetics			
				(Anti-Diabetics)	C		
				Estrogens			
				(Estrogens)	C		
				Thyroid (Thyroid)	C		

Date:07/06/04ISR Number: 4394651-0Report Type:Expedited (15-DaCompany Report #2004042614

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Affective Disorder	Consumer	Nardil (Phenelzine			
Initial or Prolonged		Anger		Sulfate)	PS		
		Anxiety		Neurontin			
		Bradyphrenia		(Gabapentin)	SS		
		Depression					
		Drug Ineffective					
		Memory Impairment					

Date:07/06/04ISR Number: 4394902-2Report Type:Expedited (15-DaCompany Report #2004042400

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Breast Disorder Male	Foreign	Neurontin			
200 MG (100		Nipple Neoplasm	Consumer	(Gabapentin)	PS		ORAL
MG, 2 IN 1		Refusal Of Treatment By	Company				
D), ORAL		Patient	Representative				

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Neurontin			
1800 MG		Dysphasia		(Gabapentin)	PS		ORAL
(TID), ORAL		Speech Disorder					
				Levetiracetam			
				(Levetiracetam)	C		
				Diazepam (Diazepam)	C		
				Codeine Sulfate			
				(Codeine Sulfate)	C		
				Peginterferon			
				Alfa-2a			
				(Peginterferon			
				Alfa-2a)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/07/04ISR Number: 4394386-4Report Type:Expedited (15-DaCompany Report #2004042860

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Consumer	Neurontin			
		Drug Ineffective		(Gabapentin)	PS		
100 MG (100							
MG, DAILY)		Hallucinations, Mixed					
		Nightmare		Interferon Beta			
		Suicidal Ideation		(Interferon Beta)	SS		
		Tearfulness					

Date:07/07/04ISR Number: 4395054-5Report Type:Expedited (15-DaCompany Report #2004043464

Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Computerised Tomogram	Health	Neurontin			
Other		Abnormal	Professional	(Gabapentin)	PS		
		Confusional State		Dilantin Suspension			
		Dysarthria		(Phenytoin Sodium)	SS		
		Memory Impairment		Phenobarbital			
				(Phenobarbital)	SS		
				Antiepileptics			
				(Antiepileptics)	SS		

Date:07/07/04ISR Number: 4395056-9Report Type:Expedited (15-DaCompany Report #2004036533

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression	Consumer	Zyrtec (Tablets)			
		Anxiety	Health	(Cetirizine)	PS		ORAL
5 MG (5 MG,							
DAILY), ORAL		Drug Ineffective	Professional				
		Educational Problem		Neurontin			
		Face Oedema		(Gabapentin)	SS		ORAL
ORAL							
		Migraine		Narine Repetabs			
		Nervousness		(Loratadine,			

Obsessive-Compulsive
 Disorder
 Personality Change
 Photophobia
 Restlessness
 Seasonal Allergy
 Sinus Congestion
 Sinus Disorder
 Sinus Headache

Pseudoephedrine
 Sulfate) SS

Date:07/07/04ISR Number: 4395064-8Report Type:Expedited (15-DaCompany Report #2004038283
 Age:75 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Arrhythmia Drug Ineffective	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG, 3 IN 1 D), ORAL		Feeling Abnormal	Professional	Hyoscyamine Sulfate (Hyoscyamine Sulfate) Librax (Chlordiazepoxide Hydrochloride,	C		

Freedom Of Information (FOI) Report

Clidinium Bromide) C
 Primidone
 (Primidone) C

Date:07/07/04ISR Number: 4395817-6Report Type:Expedited (15-DaCompany Report #2004043254
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (300 MG, 1 IN 1 D), ORAL		Tenosynovitis	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL

Enalapril Maleate C
 Furosemide C
 Insulin C
 Alendronate Sodium C

Date:07/08/04ISR Number: 4396039-5Report Type:Expedited (15-DaCompany Report #2004027685
 Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 400 MG (DAILY), ORAL		Anger Anxiety Asthenia	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Cognitive Disorder
 Confusional State
 Depression
 Disturbance In Attention
 Drug Dependence
 Drug Ineffective
 Drug Withdrawal Syndrome
 Dysstasia
 Fatigue
 Flat Affect
 Hyperhidrosis
 Hypotonia
 Insomnia

Nervous System Disorder
Peripheral Coldness
Speech Disorder
Treatment Noncompliance
Weight Decreased

Date:07/08/04ISR Number: 4396042-5Report Type:Expedited (15-DaCompany Report #2004029944
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Activities Of Daily Living Impaired	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
THREE TIMES A DAY, ORAL		Catatonia	Professional				
		Dry Mouth Dysphonia Eye Rolling Impaired Driving Ability Memory Impairment Somnolence		Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/08/04ISR Number: 4396044-9Report Type:Expedited (15-DaCompany Report #2004043537
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin			
Other		Completed Suicide		(Gabapentin)	PS		ORAL
ORAL		Injury					
		Pain					
		Suicidal Ideation					

Date:07/08/04ISR Number: 4396046-2Report Type:Expedited (15-DaCompany Report #2004043231
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser	Health	Neurontin			
		Euphoric Mood	Professional	(Gabapentin)	PS		
600 MG		Overdose					

Date:07/08/04ISR Number: 4397037-8Report Type:Expedited (15-DaCompany Report #2004032696
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Asthenia	Foreign	Neurontin			
Other		Coordination Abnormal	Health	(Gabapentin)	PS		ORAL
900 MG (300		Dizziness	Professional				
MG, 3 IN 1		Impaired Driving Ability					
D), ORAL							

Date:07/08/04ISR Number: 4397259-6Report Type:Expedited (15-DaCompany Report #2004038392
Age:82 YR Gender:Male I/FU:F

Outcome	PT
Death	Agnosia
Hospitalization -	Allodynia

Initial or Prolonged
Other

Alpha 2 Globulin
Increased
Amyotrophy
Anaemia
Arrhythmia
Arthralgia
Blood Immunoglobulin G
Decreased
Blood Pressure Systolic
Decreased
Burning Sensation
Cardiovascular Disorder
Carpal Tunnel Syndrome
Cervical Myelopathy
Cervical Spinal Stenosis
Demyelinating
Polyneuropathy
Disease Recurrence
Dyspnoea
Dysstasia
Fatigue
Heart Rate Increased
Hepatic Steatosis
Hypergammaglobulinaemia
Hypertension
Hypoaesthesia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypoalbuminaemia Hypokinesia Ill-Defined Disorder					
		Impaired Driving Ability Jaundice	Foreign Health	Neurontin (Gabapentin)	PS		
DAILY		Neoplasm Recurrence Neuralgia	Professional Company	Oxcarbazepine (Oxcarbazepine)	SS		
600 MG, DAILY		Pain In Extremity Paraesthesia Paraneoplastic Syndrome Peripheral Sensory Neuropathy Red Blood Cell Sedimentation Rate Increased Rheumatoid Factor Increased Serum Ferritin Decreased Sudden Death Tinel'S Sign Ventricular Extrasystoles Weight Increased	Representative	Tramadol (Tramadol) Omeprazole (Omeprazole) Zolpidem (Zolpidem) Thiamine Hydrochloride (Thiamine Hydrochloride) Pyridoxine Hydrochloride (Pyridoxine Hydrochloride) Prednisone (Prednisone) Paracetamol (Paracetamol) Diclofenac Sodium (Diclofenac Sodium)	C C C C C C C C		

Date:07/09/04ISR Number: 4395453-1Report Type:Expedited (15-DaCompany Report #2004043506
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Asthenia Dizziness	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Intervertebral Disc Protrusion		Fluoxetine (Fluoxetine) Raloxifene Hydrochloride (Raloxifene Hydrochloride) Lorazepam (Lorazepam)	C C C		

Simvastatin
(Simvastatin) C
Lisinopril
(Lisinopril) C
Paracetamol
(Paracetamol) C

Date:07/09/04ISR Number: 4395456-7Report Type:Expedited (15-DaCompany Report #2004029925
Age:31 YR Gender:Female I/FU:F

Outcome PT
Hospitalization - Aggression
Initial or Prolonged Alcoholism
Other Anxiety
Convulsion
Depersonalisation
Head Injury
Hypervigilance
Loss Of Consciousness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
1400 MG	(EVERY DAY),	Overdose Rash Sexual Assault Victim	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Skin Laceration Treatment Noncompliance	Professional				
				Gabapentin (Gabapentin)	SS		
				Carisoprodol (Carisoprodol)	SS		
				Sertraline Hydrochloride (Sertraline Hydrochloride)	C		
				Obetroln (Amfetamine Aspartate, Amfetamine Sulfate, Dexamfetamine Saccharate,	C		

Date:07/09/04ISR Number: 4395570-6Report Type:Expedited (15-DaCompany Report #2003030361
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Arthritis Bipolar Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Drug Hypersensitivity Glossitis Mood Swings Overweight Swollen Tongue Tongue Geographic		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS		
				Anti-Diabetics	C		
				Estrogens	C		
				Thyroid	C		

Date:07/09/04ISR Number: 4396022-XReport Type:Expedited (15-DaCompany Report #2004044358
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pruritus	Health	Neurontin			
300 MG (100		Therapeutic Response	Professional	(Gabapentin)	PS		
MG, 3 IN 1 D)		Unexpected	Company				
			Representative				

Date:07/12/04ISR Number: 4395454-3Report Type:Direct Company Report #CTU 222509
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthralgia		Neurontin 100 Mg	PS		ORAL
1-3 CAPSULES		Hypoaesthesia					
3 TIMES/D		Joint Range Of Motion					
ORAL		Decreased Paraesthesia Swelling					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/12/04ISR Number: 4395535-4Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 222508

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG AM		Fall		Neurontin 600 Mg	PS		ORAL
ORAL		Loss Of Consciousness					
600 MG PM		Off Label Use		Neurontin 100 Mg	SS		ORAL
ORAL							

Date:07/12/04ISR Number: 4396581-7Report Type:Expedited (15-DaCompany Report #2002068598
 Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (THREE TIMES DAILY), ORAL		Arthralgia Asthma Bronchitis Acute Condition Aggravated Dizziness Drug Interaction Nasopharyngitis Pyrexia	Foreign Study Health Professional	Gabapentin (Tablets) (Gabapentin) Carbamazepine (Carbamazepine) Clobazam (Clobazam) Theophylline (Theophylline) Pranlukast (Pranlukast) Salbutamol (Salbutamol) Budesonide (Budesonide) Tulobuterol Hydrochloride (Tulobuterol Hydrochloride) Methylprednisolone Sodium Succinate (Methylprednisolone Sodium Succinate)	PS C C C C C C C C C		ORAL

Aminophylline	
(Aminophylline)	C
Actit (Maltose,	
Sodium Acetate,	
Potassium Chloride,	
Sodium Chloride,	
Potassium Phosphate	C
Solita T	
(Electrolytes Nos)	C
Cefdinir (Cefdinir)	C
Acemetacin	
(Acemetacin)	C
Pentoxyverine	
(Pentoxyverine)	C
Sodium Chlorine	
(Sodium Chlorine)	C
Bromhexine	
Hydrochloride	
(Bromhexine	
Hydrochloride	C
Procaterol	
Hydrochloride	
(Procaterol	
Hydrochloride)	C
Hydrocortisone	

Freedom Of Information (FOI) Report

Sodium Succinate
 (Hydrocortisone
 Sodium Succinate) C
 Soldem 3a (Sodium
 Lactate, Potassium
 Chloride, Sodium
 Chloride,
 Carbohydrates Nos) C
 Diazepam (Diazepam) C
 Magnesium Oxide
 (Magnesium Oxide) C
 Cefcapene Pivoxil
 Hydrochloride
 (Cefcapene Pivoxil
 Hydrochloride) C

Date:07/12/04ISR Number: 4398300-7Report Type:Expedited (15-DaCompany Report #2004044295
 Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Brain Damage	Foreign	Neurontin			
Hospitalization - 800 MG (400 Initial or Prolonged MG, 2 IN 1D), Other ORAL		Pneumonia	Consumer	(Gabapentin)	PS		ORAL
		Pyrexia		Enalapril Maleate (Enalapril Maleate)	C		

Date:07/12/04ISR Number: 4398313-5Report Type:Expedited (15-DaCompany Report #2004044301
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 MG, 3 IN 1 D), ORAL		Asterixis	Foreign	Neurontin (Tablets)			
		Asthenia	Health	(Gabapentin)	PS		ORAL
		Encephalopathy	Professional				
		Metabolic Encephalopathy		Lisinopril			
		Toxic Induced					

Encephalopathy

(Lisinopril)	C
Acetylsalicylic Acid	
(Acetylsalicylic	
Acid)	C
Oxazepam (Oxazepam)	C
Buprenorphine	
Hydrochloride	
(Buprenorphine	
Hydrochloride)	C
Zidovudine	
(Zidovudine)	C
Lamivudine	
(Lamivudine)	C
Efavirenz	
(Efavirenz)	C
Pravastatin Sodium	
(Pravastatin Sodium)	C
Calcium Acetate	
(Calcium Acetate)	C
Alfuzosin	
(Alfuzosin)	C
Doxepin	
Hydrochloride	
(Doxepin	

Freedom Of Information (FOI) Report

Hydrochloride) C
 Esomeprazole
 (Esomeprazole) C

Date:07/12/04ISR Number: 4399233-2Report Type:Expedited (15-DaCompany Report #2004044191

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2400 MG (500 Other MG, QID	Accidental Overdose Confusional State Contusion Convulsion Drug Interaction Drug Toxicity Feeling Jittery Loss Of Consciousness Memory Impairment Mental Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
INTERVAL: EVERY DAY), ORAL			Atarax (Hydroxyzine Hydrochloride) Vistaril (Hydroxyzine Hydrochloride) Lithium Carbonate (Lithium Carbonate) Tiagabine Hydrochloride (Tiagabine Hydrochloride) Escitalopram (Escitalopram) Tetrabenazine (Tetrabenazine) Rofecoxib (Rofecoxib) Zolpidem Tartrate (Zolpidem Tartrate) Ramipril (Ramipril) Tolterodine L-Tartrate	SS SS SS C C C C C C		

(Tolterodine L-Tartrate)	C
Esomeprazole (Esomeprazole)	C
Atorvastatin (Atorvastatin)	C
Pioglitazone (Pioglitazone)	C

Date:07/12/04ISR Number: 4399295-2Report Type:Expedited (15-DaCompany Report #2004044330
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other 300 MG (300 MG, 1 IN 1 D), ORAL		Lung Infection Pseudomonal Retinal Oedema Vision Blurred Visual Acuity Reduced	Health Professional	Neurontin (Gabapentin) Rosiglitazone Maleate (Rosiglitazone)	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Maleate) C
 Glimepiride
 (Glimepiride) C
 Metformin
 (Metformin) C
 Lisinopril
 (Lisinopril) C

Date:07/13/04ISR Number: 4396865-2Report Type:Direct
 Age:13 YR Gender:Female I/FU:I

Company Report #CTU 222709

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening GENERIC	Abortion		Neurontin	PS		
Hospitalization - NEURONTIN	Adoption					
Initial or Prolonged PILLS	Congenital Anomaly					
Disability LITHIUM PILLS	Drug Exposure During		Lithium	SS		
Required HALDOL DEC	Pregnancy		Haldol	SS		
Intervention to SHOT	Injury					
Prevent Permanent RESPRIDAL	Pregnancy		Respridol	SS		
Impairment/Damage PILLS	Road Traffic Accident					

Date:07/13/04ISR Number: 4397793-9Report Type:Expedited (15-DaCompany Report #2004032764
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Aortic Valve Disease Cardiac Murmur	Consumer Health	Neurontin (Gabapentin)	PS		
300 MG (100 MG, TID, INTERVAL: EVER DAY)	Condition Aggravated	Professional				
	Drug Interaction					
	Grand Mal Convulsion					

ORAL

Calcium (Calcium) SS

ORAL

10 MEQ (10

Potassium Chloride
(Potassium Chloride) SS

MEQ, QD

INTERVAL:

EVERY DAY)

Magnesium
(Magnesium) SS

Selenium (Selenium) SS

Alendronate Sodium
(Alendronate Sodium) C

Digoxin (Digoxin) C

Levothyroxine Sodium
(Levothyroxine

Sodium) C

Warfarin Sodium
(Warfarin Sodium) C

Fluovastatin
(Fluovastatin) C

Famotidine
(Famotidine) C

Paracetamol
(Paracetamol) C

Diphhenhydramine
Hydrochloride

(Diphhenhydramine
Hydrochloride) C

Estrogens Conjugated
(Estrogens

Conjugated) C

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Freedom Of Information (FOI) Report

Furosemide
(Furosemide) C

Date:07/13/04ISR Number: 4397900-8Report Type:Expedited (15-DaCompany Report #2004013492
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bipolar Disorder Brain Damage Closed Head Injury Depressed Level Of Consciousness Drug Dependence Drug Withdrawal Syndrome Hallucination, Visual Intentional Self-Injury Nausea Pain Psychotic Disorder Stupor Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:07/13/04ISR Number: 4397901-XReport Type:Expedited (15-DaCompany Report #2004039225
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability							
Other		Confusional State Difficulty In Walking Disorientation Drug Ineffective Psychotic Disorder	Consumer Health Professional	Neurontin (Gabapentin)	PS		

Date:07/13/04ISR Number: 4397934-3Report Type:Expedited (15-DaCompany Report #2004039495
Age:68 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability							
		Difficulty In Walking	Consumer	Neurontin			

Other	Drug Ineffective	(Gabapentin)	PS	ORAL
ORAL	Pain	All Other Therapeutic Products	C	

Date:07/13/04ISR Number: 4398081-7Report Type:Expedited (15-DaCompany Report #2004044921
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Chest Discomfort Eye Swelling	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
300 MG (300 MG, DAILY INTERVAL: EVERYDAY), ORAL		Nausea Pruritus Steatorrhoea Swollen Tongue Vomiting	Professional	Baclofen Salbutamol	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/13/04ISR Number: 4398125-2Report Type:Expedited (15-DaCompany Report #2004044456

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Emphysema	Consumer	Neurontin (Gabapentin)	PS		ORAL
3000 MG (3 IN							
D), ORAL							
				Acetylsalicylic Acid	C		
				Glipizide	C		
				Gemfibrozil	C		
				Valsartan	C		

Date:07/13/04ISR Number: 4398131-8Report Type:Expedited (15-DaCompany Report #2004044965

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebral Cyst Excision	Consumer	Neurontin (Gabapentin)	PS		
				Paracetamol	C		

Date:07/13/04ISR Number: 4398134-3Report Type:Expedited (15-DaCompany Report #2004034121

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bedridden Cellulitis	Consumer	Neurontin (Gabapentin)	PS		
300 MG							
		Cystitis					
		Depression					
		Diabetes Mellitus					
		Dizziness					
		Drug Ineffective For					
		Unapproved Indication					
		Lung Neoplasm Malignant					
		Lymphoedema					
		Malaise					
		Muscle Spasms					
		Oedema Peripheral					

Pain
Unevaluable Event
Vaginal Pain
Vomiting

Date:07/13/04ISR Number: 4398135-5Report Type:Expedited (15-DaCompany Report #2004032769

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1600 MG (400 Other MG, 4 IN 1 D), ORAL	Delirium Middle Insomnia Oral Intake Reduced Pain Pain In Extremity Staphylococcal Infection Weight Decreased Weight Increased	Consumer Health Professional	Neurontin (Gabapentin) Metformin Hydrochloride Insulin Ramipril Potassium Chloride Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid,	PS C C C C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nicotinamide, C
 Calcium C
 Furosemide C

Date:07/14/04ISR Number: 4398249-XReport Type:Direct
 Age:72 YR Gender:Female I/FU:I

Company Report #CTU 222769

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 300 MG ORALLY		Abnormal Dreams		Neurontin 300 Mg			
Initial or Prolonged Disability		Anger		Park Davis/Pfizer	PS	Park Davis/Pfizer	ORAL
		Arrhythmia					
		Coordination Abnormal					
		Headache					
		Inflammation					
		Memory Impairment					
		Mood Altered					
		Muscle Rigidity					
		Nightmare					
		Oedema Peripheral					
		Skin Ulcer					
		Speech Disorder					
		Vision Blurred					

Date:07/14/04ISR Number: 4401591-7Report Type:Expedited (15-DaCompany Report #2004019177
 Age:42 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Aspiration	Consumer	Neurontin			
Other		Pneumonia	Health	(Gabapentin)	PS		
		Relapsing Polychondritis	Professional				

Date:07/14/04ISR Number: 4401632-7Report Type:Expedited (15-DaCompany Report #2004045177
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Death	Health	Neurontin			
300 MG (300			Professional	(Gabapentin)	PS		ORAL

D), ORAL

All Other
Therapeutic Products
(All Other
Therapeutic
Products) C

Date:07/14/04ISR Number: 4401837-5Report Type:Expedited (15-DaCompany Report #2004044348

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Feeling Cold	Consumer	Neurontin			
Other		Headache		(Gabapentin)	PS		
		Hyperhidrosis		Quetiapine Fumarate			
		Muscle Twitching		(Quetiapine			
		Musculoskeletal Stiffness		Fumarate)	SS		
		Nerve Injury		Risperidone			
		Suicidal Ideation		(Risperidone)	SS		
		Tardive Dyskinesia		Oxcarbazepine			
		Weight Decreased		(Oxcarbazepine)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Zolpidem Tartrate
(Zolpidem Tartrate) C

Date:07/15/04ISR Number: 4400685-XReport Type:Direct
Age:71 YR Gender:Female I/FU:I

Company Report #CTU 222891

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 80MG/M2 IV		Asthenia Fall		Taxol 80 Mg/M2 Iv Q Wk	PS		ORAL
QWK		Infection					
600 MG PO TID		Pneumonia		Neurontin 60 Mg Po Tid	SS		ORAL

Date:07/15/04ISR Number: 4403395-8Report Type:Expedited (15-DaCompany Report #2004028897
Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG (TID), ORAL		Difficulty In Walking Drug Effect Decreased	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
		Dysstasia					
		Pharmaceutical Product Complaint		Ibuprofen Budesonide Triamcinolone Acetonide Salmeterol Xinafoate Narine Repetabs (Loratadine, Pseudoephedrine Sulfate) Dyazide (Hydrochlorothiazide , Triamterene) Cimetidine Fluocinonide	C C C C C C C C C C		

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aortic Valve Disease Cardiac Murmur	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG, 3 IN 1 D), ORAL ORAL		Condition Aggravated Drug Effect Decreased Drug Interaction	Professional	Calcium (Calcium)	SS		ORAL
10 MEQ (10 MEQ, QD INTERVAL: EVERY DAY)		Grand Mal Convulsion Pharmaceutical Product Complaint		Potassium Chloride (Potassium Chloride)	SS		
				Magnesium (Magnesium)	SS		
				Selenium (Selenium)	SS		
				Alendronate Sodium (Alendronate Sodium)	C		
				Digoxin (Digoxin)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		

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Freedom Of Information (FOI) Report

Warfarin Sodium (Warfarin Sodium)	C
Fluvastatin (Fluvastatin)	C
Famotidine (Famotidine)	C
Paracetamol (Paracetamol)	C
Diphenhydramine Hydrochloride(Diphen hydramine Hydrochloride)	C
Estrogens Conjugated (Estrogens Conjugated)	C
Furosemide (Furosemide)	C

Date:07/15/04ISR Number: 4404810-6Report Type:Expedited (15-DaCompany Report #2004040067

Age:47 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG (600 Other MG, 3 IN 1 D), ORAL	Bronchospasm Gastrooesophageal Reflux Disease Odynophagia Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL			Olanzapine (Olanzapine)	SS		ORAL
ORAL			Aripiprazole (Aripiprazole)	SS		ORAL
			Valproate Semisodium Paroxetine Hydrochloride Clonazepam Diphenhydramine Hydrochloride	C C C C C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (300 Other MG, 3 IN 1 D) ORAL	Fatigue Headache Loss Of Consciousness Somnolence Tremor	Foreign Health Professional	Neurontin (Gabapentin) Carisoprodol Paracetamol Panadeine Co (Codeine Phosphate, Paracetamol) Diazepam	PS C C C C		ORAL

Outcome	PT
Hospitalization - Initial or Prolonged	Condition Aggravated Drug Ineffective

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pain Pancreatitis Chronic	Report Source	Product	Role	Manufacturer	Route
60 MG, Q12H			Health Professional Other	Ms Contin Tablets (Morphine Sulfate) Cr Tablet	PS		
				Neurontin (Gabapentin)	SS		
				Oxycontin (Oxycodone Hydrochloride)	C		
				Msir (Morphine Sulfate)	C		
				Lithium (Lithium)	C		
				Zoloft (Sertraline Hydrochloride)	C		
				Premarin (Estrogens Conjugated)	C		

Date:07/16/04ISR Number: 4404885-4Report Type:Expedited (15-DaCompany Report #2004040176
Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - (DAILY) Initial or Prolonged Other		Cardiac Failure Congestive	Health Professional	Neurontin (Gabapentin)	PS		
		Confusional State		Furosemide (Furosemide)	C		
		Disorientation		Potassium Chloride (Potassium Chloride)	C		
		Feeling Abnormal		Levothyroxine Sodium (Levothyroxine Sodium)	C		
		Gait Disturbance		Paroxetine Hydrochloride			
		Lung Neoplasm Malignant		Hydrochloride (Paroxetine Hydrochloride)	C		
		Mental Disorder		Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
		Miosis		Advicor - Slow Release (Lovastatin, Nicotinic Acid	C		
		Prostate Cancer					
		Pupillary Reflex Impaired					
		Tremor					

Acetylsalicylic Acid
(Acetylsalicylic
Acid) C
Oxycodone
Hydrochloride
(Oxycodone
Hydrochloride) C
Oxycocet (Oxycodone
Hydrochloride,
Paracetamol) C

Date:07/16/04ISR Number: 4404888-XReport Type:Expedited (15-DaCompany Report #2004041566
Age:71 YR Gender:Female I/FU:F

Outcome PT
Other Back Pain
Blood Cholesterol
Increased

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Freedom Of Information (FOI) Report

Dose	Duration	Condition Aggravated Drug Effect Decreased Feeling Abnormal	Report Source	Product	Role	Manufacturer	Route
1600 MG (800 MG, 2 IN 1 D), ORAL		Muscle Disorder Pharmaceutical Product Complaint Wrist Surgery	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
10 MG (10 MG, 1 IN 1 D), ORAL				Lipitor (Atorvastatin)	SS		ORAL
				Metoprolol (Metoprolol)	C		
				Esomeprazole (Esomeprazole)	C		
				Tocopherol (Tocopherol)	C		

Date:07/16/04ISR Number: 4405265-8Report Type:Expedited (15-DaCompany Report #2004033211
Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG, 3 IN 1 D), ORAL		Aggression Alcohol Use Anorexia Asocial Behaviour Catatonia Depression Drug Intolerance Eating Disorder Paranoia Psychotic Disorder Thinking Abnormal Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Trazodone (Trazodone)	C		
				Antihyperensives (Antihypertensives)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Autonomic Nervous System Imbalance	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
		Hallucination, Auditory	Professional	Risperidone (Risperidone)	C		
				Insulin (Insulin)	C		
				Oxazepam (Oxazepam)	C		
				Metformin Hydrochloride (Metformin Hydrochloride)	C		
				Gliclazide (Gliclazide)	C		
				Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
				Furosemide (Furosemide)	C		
				Amlodipine Besilate (Amlodipine Besilate)	C		

FDA - Adverse Event Reporting System (AERS)

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Lansoprazole
 (Lansoprazole) C
 Atorvastatin
 (Atorvastatin) C
 Atenolol (Atenolol) C
 Potassium Chloride
 (Potassium Chloride) C
 Ferrous Sulfate
 (Ferrous Sulfate) C
 Docusate Sodium
 (Docusate Sodium) C

Date:07/19/04ISR Number: 4402115-0Report Type:Direct
 Age:12 YR Gender:Female I/FU:I

Company Report #CTU 223051

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - INTRAVENTOUS	1250 MG IV Q	6	Stevens-Johnson Syndrome	Chloramphenicol 1 Gm	PS		
Initial or Prolonged	21 DAY			Streptomycin 1 Gm	SS		
INTRAVENTOUS	1 GM IV Q 12	48 HR		Toradol	SS		
INTRAVENTOUS	15MG IV X 1						
ONE DOSE				Gabapentin 100 Mg	SS		ORAL
100 MG PO TID	2 DAY						

Date:07/19/04ISR Number: 4402881-4Report Type:Expedited (15-DaCompany Report #2004042614
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Consumer	Nardil (Phenelzine Sulfate)	PS		
				Neurontin (Gabapentin)	SS		
				Lorazepam (Lorazepam)	SS		
				Clozapine (Clozapine)	C		

Pharmaceutical Product
Complaint

Lamotrigine
(Lamotrigine)

C

Date:07/19/04ISR Number: 4403613-6Report Type:Expedited (15-DaCompany Report #2004046622
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Neutropenia	Health Professional	Neurontin (Gabapentin)	PS		
UNKNOWN	UNKNOWN						
Other (UNKNOWN),							
UNKNOWN				...			C

Date:07/19/04ISR Number: 4405117-3Report Type:Expedited (15-DaCompany Report #2004041837
Age: Gender:Female I/FU:F

Outcome	PT
Life-Threatening	Adverse Event
Other	Anticonvulsant Drug Level Increased Feeling Abnormal Headache

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FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (400 MG, 3 IN 1 D)		Lethargy Pain Platelet Count Decreased	Consumer	Dilantin Suspension (Phenytoin Sodium) Neurontin (Gabapentin)	PS SS		

Date:07/20/04ISR Number: 4404436-4Report Type:Expedited (15-DaCompany Report #2004046062
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Other		Amnesia Anger Blood Glucose Decreased Blood Pressure Increased Circulatory Collapse Coma Convulsion Depressed Mood Diarrhoea Drug Ineffective Haemoglobin Decreased Hallucination, Auditory Hypermetropia Overdose Panic Attack Respiratory Rate Decreased Simple Partial Seizures Urinary Tract Infection	Consumer	Neurontin (Tablets) (Gabapentin) Lorazepam (Lorazepam) Morphine Sulfate (Morphine Sulfate) Morphine (Morphine) Temazepam (Temazepam) Valproate Semisodium (Valproate Semisodium)	PS SS C C C		ORAL

Date:07/20/04ISR Number: 4404516-3Report Type:Expedited (15-DaCompany Report #2004040168
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other	Asphyxia	Foreign	Neurontin		
600 MG (300	Cerebrovascular Accident	Consumer	(Gabapentin)	PS	ORAL
MG, 2 IN 1	Convulsion				
D), ORAL	Difficulty In Walking				
	Dizziness		All Other		
	Hypertension		Therapeutic Products	C	
			Rosiglitazone		
			Maleate	C	
			Metformin		
			Hydrochloride	C	

Date:07/21/04ISR Number: 4404576-XReport Type:Direct Company Report #CTU 223251
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Drug Ineffective		Neurontin	PS		
600 MG 2CAPS		General Physical Health					
3 TIMES A		Deterioration					
DAY, 300 MG							
2CAPS 3 TIMES							
A DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/21/04ISR Number: 4407114-0Report Type:Expedited (15-DaCompany Report #2004026318
Age:69 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Diabetes Mellitus	Consumer	Neurontin			
Initial or Prolonged	Difficulty In Walking		(Gabapentin)	PS		
Other	Eczema		Celecoxib			
	Gastrointestinal Disorder		(Celecoxib)	SS		
	Localised Infection		Insulin (Insulin)	SS		
	Loss Of Consciousness					
	Nerve Injury					
	Pain					
	Paraesthesia					
	Spinal Disorder					

Date:07/21/04ISR Number: 4407239-XReport Type:Expedited (15-DaCompany Report #2004046582
Age:17 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Abnormal Behaviour		Neurontin			
Initial or Prolonged	Conversion Disorder		(Gabapentin)	PS		ORAL
800 MG (400						
Other	Convulsion					
MG, 2 IN 1 D)						
	Dizziness					
ORAL						
	Fall		Carbamazepine			
	Head Injury		(Carbamazepine)	SS		
	Loss Of Consciousness					
	Treatment Noncompliance					

Date:07/21/04ISR Number: 4407475-2Report Type:Expedited (15-DaCompany Report #2004046607
Age:66 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Hypertensive Crisis	Foreign	Neurontin (Tablets)			
Initial or Prolonged		Consumer	(Gabapentin)	PS		ORAL
1800 MG (600						
MG, 3 IN 1						

D), ORAL

Date:07/21/04ISR Number: 4407479-XReport Type:Expedited (15-DaCompany Report #2004046365

Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State	Foreign	Neurontin			
900 MG (300		Dizziness	Health	(Gabapentin)	PS		ORAL
MG, 3 IN 1		Speech Disorder	Professional				
D), ORAL		Transient Ischaemic					
		Attack					

Date:07/21/04ISR Number: 4407670-2Report Type:Expedited (15-DaCompany Report #2004045177

Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Sudden Death	Health	Neurontin			
300 MG (300			Professional	(Gabapentin)	PS		ORAL
MG, 1 IN 1							
D), ORAL				Estradiol			
				(Estradiol)	C		
				All Other			

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Freedom Of Information (FOI) Report

Therapeutic Products
 (All Other
 Therapeutic
 Products) C
 Amlodipine Besilate
 (Amlodipine
 Besilate) C
 Levothyroxine
 (Levothyroxine) C
 Oxcarbazepine
 (Oxcarbazepine) C
 Olanzapine
 (Olanzapine) C
 Venlafaxine
 Hydrochloride
 (Venlafaxine
 Hydrochloride) C

Date:07/21/04ISR Number: 4407877-4Report Type:Expedited (15-DaCompany Report #2004046521
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1900 MG, ORAL Other		Pericardial Effusion Pericarditis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Zolpidem Tartrate (Zolpidem Tartrate)	C		
				Metformin Hydrochloride (Metformin Hydrochloride)	C		
				Atorvastatin (Atorvastatin)	C		
				Quinapril Hydrochloride (Quinapril Hydrochloride)	C		
				Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		

Date:07/22/04ISR Number: 4405153-7Report Type:Direct
Age:46 YR Gender:Female I/FU:I

Company Report #CTU 223371

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300 MG TID PO	Completed Suicide		Neurontin	PS		ORAL
				Tylenol # 3	C		
				Voltaren	C		

Date:07/22/04ISR Number: 4405219-1Report Type:Direct
Age:53 YR Gender:Male I/FU:I

Company Report #CTU 223351

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Required		Agitation		Gabapentin	PS		
Intervention to Prevent Permanent Impairment/Damage				Trazodone	C		
				Mirtazapine	C		
				Thiamine	C		
				Multivitamins/Zinc (Centrum)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Magnesium Hydroxide C
 Alohol/Mgoh/Simth C
 Acetaminophen C

Date:07/22/04ISR Number: 4408301-8Report Type:Expedited (15-DaCompany Report #2004040186

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (300 MG, 2 IN 1 D), ORAL		Arthralgia Dysgraphia Gait Disturbance Hypoaesthesia	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
30 MG (30 MG), ORAL		Pain In Extremity Spinal Cord Disorder		Procardia Xl (Nifedipine)	SS		ORAL
200 MG (200 MG, 1 IN 1 D)				Celebrex (Celecoxib) (Celecoxib)	SS		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:07/22/04ISR Number: 4408313-4Report Type:Expedited (15-DaCompany Report #2004046740

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other ORAL		Fracture Pain	Foreign Health	Neurotin (Gabapentin)	PS		ORAL
UNKNOWN	UNKNOWN	Postoperative Infection Road Traffic Accident	Professional	Ciprofloxacin (Ciprofloxacin)	SS		

Status Epilepticus

Tramadol
Hydrochloride
(Tramadol
Hydrochloride) SS

UNKNOWN UNKNOWN

Folic Acid (Folic
Acid) SS

UNKNOWN UNKNOWN

Phenprocoumon
(Phenprocoumon) C
Esomeprazole
(Esomeprazole) C
Paracetamol
(Paracetamol) C
Carbamazepine
(Carbamazepine) C

Date:07/22/04ISR Number: 4408343-2Report Type:Expedited (15-DaCompany Report #2004041539

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG, (300 MG, 2 IN 1 D), ORAL	Blood Insulin Decreased Haemorrhage Pancreas Transplant Post Procedural Complication Urine Amylase	Foreign Consumer	Neurotin (Tablets) (Gabapentin0	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/22/04ISR Number: 4409331-2Report Type:Expedited (15-DaCompany Report #2004046590

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Health	Neurontin (Tablets)			
800 MG		Drug Exposure During	Professional	(Gabapentin)	PS		
		Pregnancy		Phenytoin Sodium			
		Drug Exposure Via Breast		(Phenytoin Sodium)	C		
		Milk					
		Grand Mal Convulsion					
		Pharmaceutical Product					
		Complaint					
		Pregnancy					

Date:07/22/04ISR Number: 4409351-8Report Type:Expedited (15-DaCompany Report #2004048924

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Consumer	Neurontin			
2700 MG (300		Blood Testosterone		(Gabapentin)	PS		
MG)		Decreased					
		Disinhibition					
		Legal Problem					

Date:07/23/04ISR Number: 4409502-5Report Type:Expedited (15-DaCompany Report #2004047168

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaemia	Consumer	Neurontin			
900 MG (300		Blood Iron Decreased		(Gabapentin)	PS		ORAL
MG, 3 IN 1		Bone Marrow Disorder					
D), ORAL		Fatigue					
		Full Blood Count		Naproxen	C		
		Decreased		Fluoxetine			

Somnolence

Hydrochloride C
Alendronate Sodium C
Calcium C
Librax
(Chlordiazepoxide
Hydrochloride,
Clidinium Bromide) C

Date:07/23/04ISR Number: 4409514-1Report Type:Expedited (15-DaCompany Report #2004047491

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 800 MG (400 MG, 2 IN 1 D), ORAL		Arteriovenous Malformation Condition Aggravated Depression	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Dizziness Fatigue Headache Irritability		Paroxetine Hydrochloride (Paroxetine Hydrochloride)	SS		ORAL
		Pain Treatment Noncompliance		Estrogens Conjugated (Estrogens Conjugated)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/23/04ISR Number: 4409540-2Report Type:Expedited (15-DaCompany Report #2004047157

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Body Height Increased	Consumer	Neurontin			
600 MG (100		Emotional Disorder		(Gabapentin)	PS		ORAL
MG, 3 IN 1		Medication Tampering					
D), ORAL							

Date:07/26/04ISR Number: 4408561-3Report Type:Periodic Company Report #2004UW09339

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Rhabdomyolysis	Health	Crestor	PS		
10 MG QD PO							
Initial or Prolonged			Professional	Neurontin	SS		
300 MG				Plavix	C		
				Aspirin "Bayer"	C		
				Beta Blocker	C		

Date:07/26/04ISR Number: 4410486-4Report Type:Expedited (15-DaCompany Report #2004046633

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Cerebrovascular Accident	Foreign	Neurontin			
Initial or Prolonged		Coagulopathy	Health	(Gabapentin)	PS		ORAL
800 MG (400							
MG, 2 IN 1			Professional				
D), ORAL			Company				
			Representative	Acenocoumarol			
				(Acenocoumarol)	C		
				All Other			
				Therapeutic Products			
				(All Other			

Therapeutic
Products) C

Date:07/26/04ISR Number: 4410673-5Report Type:Expedited (15-DaCompany Report #2004047161
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Prolactin Increased Cystitis Interstitial	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Drug Ineffective Galactorrhoea Hypertrophy Breast Pain Weight Increased		All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:07/26/04ISR Number: 4410710-8Report Type:Expedited (15-DaCompany Report #2004046701
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abnormal Behaviour Dizziness	Foreign Health	Ziprasidone (Caps) (Ziprasidone)	PS		ORAL
160 MG (80 MG, 2 IN 1 D), ORAL		Epilepsy Mental Impairment	Professional Company				
1800 MG (600 MG, 3 IN 1		Scratch Skin Lesion	Representative	Gabapentin (Gabapentin)	SS		ORAL

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D), ORAL

300 MG, ORAL

ORAL

ORAL

Lamotrigine (Lamotrigine)	SS	ORAL
Ranitidine Hydrochloride (Ranitidine Hydrochloride)	SS	ORAL
Midazolam (Midazolam)	SS	ORAL
Dexchlorpheniramine Maleate (Dexchlorpheniramine Maleate)	SS	
Olanzapine (Olanzapine)	C	

Date:07/27/04ISR Number: 4412792-6Report Type:Expedited (15-DaCompany Report #2004032764

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (100 MG, 3 IN 1 D) ORAL ORAL		Cardiac Murmur Cardiac Valve Disease Condition Aggravated Convulsion Drug Interaction Herpes Virus Infection	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
10 MEQ (10 MEQ, DAILY)				Calcium (Calcium)	SS		ORAL
				Potassium Chloride (Potassium Chloride)	SS		
				Magnesium (Magnesium)	SS		
				Selenium (Selenium)	SS		
				Alendronate Sodium (Alendronate Sodium)	C		
				Digoxin (Digoxin)	C		
				Levothyroxine Sodium			

(Levothyroxine Sodium)	C
Warfarin Sodium (Warfarin Sodium)	C
Fluvastatin (Fluvastatin)	C
Famotidine (Famotidine)	C
Paracetamol (Paracetamol)	C
Diphenhydramine Hydrochloride	C
Estrogens Conjugated (Estrogens Conjugated)	C
Furosemide (Furosemide)	C

Date:07/27/04ISR Number: 4413026-9Report Type:Expedited (15-DaCompany Report #2004047826
 Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Transurethral Resection Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL				Celecoxib			

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(Celecoxib) C
 Furosemide
 (Furosemide) C
 Allopurinol
 (Allopurinol) C

Date:07/27/04ISR Number: 4413027-0Report Type:Expedited (15-DaCompany Report #2004047823
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG (100 MG, 2 IN 1 D), ORAL		Balance Disorder Blood Triglycerides Increased Body Temperature	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Increased Flatulence		Dilantin Suspension (Phenytoin Sodium)	SS		ORAL
		Partial Seizures Rash		Carbamazepine (Carbamazepine) Phenobarbital (Phenobarbital) Vitamins (Vitamins)	SS SS C		

Date:07/28/04ISR Number: 4413489-9Report Type:Expedited (15-DaCompany Report #2004040068
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2300 MG (1 IN 1 D),		Asthenia Cardiac Disorder Chest Pain Circulatory Collapse Convulsion Dizziness Headache Loss Of Consciousness Overdose Somatic Delusion	Consumer	Neurontin (Gabapentin)	PS		

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Psychotic Disorder	Health Professional Company Representative	Neurontin (Gabapentin) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) All Other Therapeutic Products (All Other Therapeutic Products)	PS C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/28/04ISR Number: 4413493-0Report Type:Expedited (15-DaCompany Report #2004048122

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Asthenia	Consumer	Neurontin			
Hospitalization - 900 MG (300	Colonic Haemorrhage		(Gabapentin)	PS		ORAL
Initial or Prolonged MG, 3 IN 1	Drug Interaction					
Other D), ORAL	Dyspnoea					
	Emphysema		All Other			
	Fall		Therapeutic Products	SS		
	Feeling Abnormal		Trazodone			
	Infrequent Bowel Movements		(Trazodone)	C		
	Knee Operation		Levothyroxine Sodium			
	Lung Disorder		(Levothyroxine Sodium)	C		
	Multi-Organ Disorder		Spironolactone			
	Urinary Tract Disorder		(Spironolactone)	C		
			Venlafaxine			
			Hydrochloride			
			(Venlafaxine Hydrochloride)	C		
			Lorazepam			
			(Lorazepam)	C		
			Tolterodine			
			L-Tartrate			
			(Tolterodine L-Tartrate)	C		
			Warfarin (Warfarin)	C		
			Hydrocodone			
			(Hydrocodone)	C		
			Risedronate Sodium			
			(Risedronate Sodium)	C		
			Esomeprazole			
			(Esomeprazole)	C		
			Diazepam (Diazepam)	C		
			Prednisone			
			(Prednisone)	C		
			Risperidone			
			(Risperidone)	C		

Date:07/28/04ISR Number: 4413785-5Report Type:Expedited (15-DaCompany Report #2004048907
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety Condition Aggravated Gun Shot Wound Injury Mental Disorder Pain Self Injurious Behaviour Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:07/28/04ISR Number: 4413932-5Report Type:Expedited (15-DaCompany Report #2004025636
Age:69 YR Gender:Male I/FU:F

Outcome	PT	Report Source
	Vertigo	Foreign Health Professional

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Dose	Duration	Company Representative	Product	Role	Manufacturer	Route
1800 MG, ORAL			Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Clopidogrel Sulfate	C		
			Verapamil Hydrochloride	C		
			Benazepril Hydrochloride	C		
			Gliclazide	C		
			Metformin Hydrochloride	C		
			Glyceryl Trinitrate	C		

Date:07/28/04ISR Number: 4413970-2Report Type:Expedited (15-DaCompany Report #2004048188
Age:17 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Condition Aggravated Dizziness	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG, (300 MG, 3 IN 1 D), ORAL		Grand Mal Convulsion Nausea	Professional Company Representative	Nitrazepam	C		

Date:07/28/04ISR Number: 4414106-4Report Type:Expedited (15-DaCompany Report #2004048216
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation Grand Mal Convulsion	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL			Professional Company				

Representative

Carbamazepine
(Carbamazepine)

C

Date:07/28/04ISR Number: 4414107-6Report Type:Expedited (15-DaCompany Report #2004048870

Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Arrhythmia	Foreign	Neurontin			
Initial or Prolonged	Urinary Tract Infection	Health	(Gabapentin)	PS		ORAL
300 MG (3 IN		Professional				
1 D), ORAL						

Date:07/29/04ISR Number: 4415010-8Report Type:Expedited (15-DaCompany Report #CH-JNJFOC-20040705129

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Drug Interaction	Foreign	Tramal (Tramadol			
	Post Procedural	Health	Hydrochloride)			
	Complication	Professional	Unspecified	PS		
	Postoperative Wound		Ciproxin			
	Complication		(Ciprofloxacin)	SS		
	Status Epilepticus		Gabapentin			
			(Gabapentin)	SS		
			Folic Acid (Folic			
			Acid)	SS		

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Marcoumar
 (Phenprocoumon) C
 Nexium
 (Esomeprazole) C
 Dafalgan
 (Paracetamol) C

Date:07/29/04ISR Number: 4415235-1Report Type:Expedited (15-DaCompany Report #2004049601
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bone Neoplasm Neurofibromatosis	Consumer	Neurontin (Gabapentin)	PS		
				Methadone (Methadone)	C		
				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
				Diazepam (Diazepam)	C		

Date:07/29/04ISR Number: 4415431-3Report Type:Expedited (15-DaCompany Report #2004027664
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Atrial Fibrillation	Consumer	Neurontin (Tablets)			
Other		Cardiac Operation		(Gabapentin)	PS		ORAL
2400 MG (800		Cardiac Pacemaker					
MG, 3 IN 1		Insertion					
D), ORAL		Coronary Artery Surgery		Donepezil			
		Decreased Appetite		Hydrochloride			
		Dementia Alzheimer'S Type		(Donepezil			
		Diarrhoea		Hydrochloride)	SS		ORAL
10 MG (10 MG,		Dizziness					
1 IN 1 D),		Hypoglycaemia					
ORAL							

Insomnia
Myocardial Infarction
Tricuspid Valve
Incompetence

Carbamazepine
(Carbamazepine) C
Warfarin Sodium
(Warfarin Sodium) C
Amiodarone
Hydrochloride
(Amiodarone
Hydrochloride) C
Digoxin (Digoxin) C
Furosemide
(Furosemide) C
Isosorbide
Mononitrate
(Isosorbide
Mononitrate) C
Potassium Chloride
(Potassium Chloride) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C
Centrum (Minerals
Nos, Vitamins Nos) C
Atorvastatin
(Atorvastatin) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C
 Magnesium
 (Magnesium) C
 Zinc (Zinc) C
 Metoprolol
 (Metoprolol) C
 Enalapril Maleate
 (Enalapril Maleate) C

Date:07/29/04ISR Number: 4443729-1Report Type:Periodic Company Report #USA-2004-0014564
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Overdose	Health Professional Other	Morphine Sulfate (Similar To Nda 19-516) (Morphine Sulfate)	PS		
				Fluoxetine	SS		
				Neurontin (Gabapentin)	SS		
				Trazodone	SS		
				Cannabis	SS		
				Alprazolam	SS		
				Temazepam`	SS		
				Oxazepam	SS		
				Metoprolol	SS		
				Dextromethorphan	SS		
				Acetaminophen (Paracetamol)	SS		
				Propoxyphene (Dextropropoxyphene)	SS		

Date:07/29/04ISR Number: 4443807-7Report Type:Periodic Company Report #USA-2003-0012038
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health	Ms Contin Tablets			

Loss Of Consciousness
Snoring

Professional
Other

(Morphine Sulfate) PS
Gabapentin SS
(Gabapentin)
Citalopram SS
(Citalopram)
Promethazine
(Promethazine) SS

Date:07/29/04ISR Number: 4444117-4Report Type:Periodic
Age:42 YR Gender:Female I/FU:I

Company Report #USA-2003-0011945

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Accidental Overdose	Health Professional Other	Morphine Sulfate (Similar To Nda 19-516) (Morphine Sulfate) Metahdone	PS		

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(Methadone)	SS
Propoxyphene	
(Dextropropoxyphene)	SS
Tramadol (Tramadol)	SS
Amitriptyline	
(Amitriptyline)	SS
Cyclobenzaprine	
(Cyclobenzaprine)	SS
Acetaminophen	
(Paracetamol)	SS
Celexa (Citalopram	
Hydrobromide)	SS
Effexor (Venlafaxine	
Hydrochloride)	SS
Neurontin	
(Gabapentin)	SS
Duragesic	C

Date:07/29/04ISR Number: 4444151-4Report Type:Periodic
 Age:41 YR Gender:Male I/FU:I

Company Report #USA-2003-0011953

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose Multiple Drug Overdose	Health Professional Other	Morphine Sulfate (Similar To Nda-19-516) (Morphine Sulfate) Fentanyl (Fentanyl) Diazepam (Diazepam) Gabapentin (Gabapentin) Lorazepam (Lorazepam) Metoclopramide (Metoclopramide) Promethazine (Promethazine) Venlafaxine (Venlafaxine)			
					PS		
					SS		
					SS		
					SS		
					SS		
					SS		
					SS		
					SS		

Date:07/29/04ISR Number: 4444312-4Report Type:Periodic
 Age:48 YR Gender:Female I/FU:I

Company Report #USA-2003-0011565

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse Multiple Drug Overdose	Health Professional Other	Morphine Sulfate (Similar To Nda 19-516) (Morphine Sulfate) Venlafaxine (Venlafaxine) Diphenhydramine (Diphenhydramine) Cannabis (Cannabis) Salicylates (Salicylates) Gabapentin (Gabapentin) Valproic Acid (Valproic Acid) Nicotine (Nicotine)	PS SS SS SS SS SS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/29/04ISR Number: 4444324-0Report Type:Periodic
 Age:43 YR Gender:Male I/FU:I

Company Report #USA-2003-0011709

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Health Professional Other	Morphine Sulfate (Similar To Nda 19516) (Morphine Sulfate) Hydrocodone Bitartrate (Similar To Ind 59175) (Hydrocodone Bitartrate) Acetaminophen (Paracetamol) Chlorpheniramine (Chlorphenamine) Cyclobenzaprine (Cyclobenzaprine) Ephedrine (Ephedrine) Gabapentin (Gabapentin) Fluconazole (Fluconazole) Promethazine (Promethazine) Meperidine Hydrochloride (Pethidine Hydrochloride) Carisoprodol (Carisoprodol) Meprobamate (Meprobamate) Setraline (Setraline) Codeine (Codeine) Metoprolol (Metoprolol)	PS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS		

Date:07/29/04ISR Number: 4444386-0Report Type:Periodic
 Age:54 YR Gender: I/FU:I

Company Report #2003UW12225

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Elavil Acetaminophen W/Hydrocodone Bitartrate Gabapentin	PS SS SS		

Date:07/29/04ISR Number: 4445347-8Report Type:Periodic Company Report #2003UW12264
 Age:34 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature Health Professional	Elavil Gabapentin Hydroxyzine	PS SS SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/30/04ISR Number: 4412074-2Report Type:Expedited (15-DaCompany Report #FR-BRISTOL-MYERS SQUIBB COMPANY-12652871
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Glucophage	PS	Bristol-Myers Squibb Company	ORAL
Hospitalization - Initial or Prolonged		Epilepsy		Aprovel	SS	Bristol-Myers Squibb Company	
		Ketosis		Furadantine	SS		ORAL
		Lactic Acidosis		Rifadine	SS		
		Septic Shock		Oflocet	SS		
				Hyperium	SS		
				Neurontin	SS		
				Ferritin	SS		

Date:07/30/04ISR Number: 4413017-8Report Type:Direct Company Report #CTU 223883
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dizziness		Diphenhydramine 50 Mg Q Hs Renewed 5/04	PS		
				Gabapentin 800 Mg Tid Increased 5/04	SS		
				Sertraline 200 Mg Daily Renewed 5/04	SS		
				Trazodone 100 Mg Q Hs Renewed 5/04	SS		
				Lisinopril 10 Mg Daily Renewed 5/04	SS		
				Diltiazem Sa 120 Mg Daily- Stopped 6/20/04	SS		
				Simvastatin	C		
				Accu-Chek Comfort Cv (Glucose) Test Strip	C		
				Omeprazole Sa	C		
				Insulin , Aspart, Hum	C		
				Levothyroxine Na			

(Synthroid) C
 Alcohol Prep Pad C
 Lancet, Techlite C

Date:07/30/04ISR Number: 4415668-3Report Type:Expedited (15-DaCompany Report #2004049605

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Migraine	Consumer	Neurontin (Gabapentin)	PS		
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		
				Oxycocet (Oxycodone Hydrochloride, Paracetamol)	SS		
				Cyclobenzaprine (Cyclobenzaprine)	SS		
				Naproxen Sodium (Naproxen Sodium)	SS		
				Midrid			

Freedom Of Information (FOI) Report

(Cidhloralphenazone,
 Isometheptene,
 Paracetamol) SS
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C
 Fluoxetine
 Hydrochloride
 (Fluoxetine
 Hydrochloride) C
 Diazepam (Diazepam) C
 Nabumetone
 (Nabumetone) C

Date:07/30/04ISR Number: 4416084-0Report Type:Expedited (15-DaCompany Report #2004033211
 Age:58 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG (300 MG, 3 IN 1 D), ORAL	Aggression Alcohol Use	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other		Antisocial Behaviour Aphagia Catatonia Depression Diet Refusal Drug Intolerance Paranoia Psychotic Disorder Thinking Abnormal Treatment Noncompliance Weight Decreased	Professional	Trazodone (Trazodone) Antihypertensives (Antihypertensives)	C C		

Date:07/30/04ISR Number: 4416085-2Report Type:Expedited (15-DaCompany Report #2004034119
 Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Device Failure	Health	Neurontin (Tablets)			

1800 MG (600
MG, 3 IN 1
D), ORAL

Weight Increased

Professional

(Gabapentin)

PS

ORAL

Furosemide
(Furosemide) C
Metformin
Hydrochloride
(Metformin
Hydrochloride) C
Pioglitazone
(Pioglitazone) C
Glimepiride
(Glimepiride) C
Rofecoxib
(Rofecoxib) C
Cetirizine
Hydrochloride
(Cetirizine
Hydrochloride) C
Ramipril (Ramipril) C
Zolpidem Tartrate

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Zolpidem Tartrate) C
 Amitriptyline
 (Amitriptyline) C
 Acetaylsalicylic
 Acid
 (Acetylsalicylic
 Acid) C
 Ultracet
 (Paracetamol,
 Tramadol
 Hydrochloride) C
 Atorvastatin
 (Atorvastatin) C

Date:08/02/04ISR Number: 4413190-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0340937A
 Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - UNKNOWN	Acute Generalised	Consumer	Zophren	PS	Glaxosmithkline	
Initial or Prolonged 250MG Three times per day	Exanthematous Pustulosis		Becilan	SS		ORAL
55 DAY	Aphthous Stomatitis					
	Dermatitis Bullous		Neurontin	SS		ORAL
INTRAVENOUS 110MG Monthly	Erythema		Taxotere	SS		
INTRAVENOUS 70MG Monthly	Inflammation		Caelyx	SS		
	Oedema Peripheral Pain In Extremity Pyrexia Rash Erythematous					

Date:08/02/04ISR Number: 4416059-1Report Type:Expedited (15-DaCompany Report #2004049208
 Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 10 MG (10 MG,	Blood Alkaline Phosphatase Increased	Foreign Health	Amlodipine (Amlodipine)	PS		ORAL

1 IN 1 D),	Pain In Extremity	Professional		
ORAL	Skin Ulcer			
600 MG (600			Gabaentin 9gabapentin)	SS ORAL
MG , 1 IN 1				
D), ORAL			Perindopril (Perindopril)	SS ORAL
2 MG (2 MG, 1				
IN 1 D), ORAL			Morphine Sulfate (Morphine Sulfate)	C
			Metformin (Metformin)	C
			Ibuprofen (Ibuprofen)	C
			Acetylsalicylic Acid(Acetylsalicylic Acid)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/02/04ISR Number: 4416069-4Report Type:Expedited (15-DaCompany Report #2004041002

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (3 IN 1 D), ORAL		Coordination Abnormal Fall	Foreign Health Professional Company Representative	Neurontin (Tablets) (Gabapentin) Tramadol/Acetaminoph en (Paracetamol Tramadol) Domperidone (Domperidone) Digoxin (Digoxin) Sotalol Hydrochloride (Sotalol Hydrochloride) Aldactazine (Altizide, Spironolactone) Fluindione (Fluindione)	PS C C C C C C		ORAL

Date:08/02/04ISR Number: 4416072-4Report Type:Expedited (15-DaCompany Report #2003002971

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG (100 MG, TID), ORAL		Circulatory Collapse Convulsion Gingivitis Neurodermatitis Pyrexia Therapeutic Response Unexpected	Foreign Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:08/02/04ISR Number: 4416074-8Report Type:Expedited (15-DaCompany Report #2004043254
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG (300 Other MG, 1 IN 1 D), ORAL	Dupuytren'S Contracture	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
			Enalapril Maleate (Enalapril Maleate)	C		
			Furosemide (Furosemide)	C		
			Insulin (Insulin)	C		
			Alendronate Sodium (Alendronate Sodium)	C		

Date:08/02/04ISR Number: 4416511-9Report Type:Expedited (15-DaCompany Report #2004049235
Age:94 YR Gender:Female I/FU:I

Outcome	PT
Death	Atrial Fibrillation
Other	Bronchitis Cardiac Failure Colon Cancer

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Disease Recurrence Intestinal Obstruction Somnolence	Report Source	Product	Role	Manufacturer	Route
INTRAVENOUS	(300 MG),		Foreign Health Professional	Neurontin (Gabapentin)	PS		
INTRAVENOUS			Company Representative				

Date:08/02/04ISR Number: 4417187-7Report Type:Expedited (15-DaCompany Report #2004049226
Age:47 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Headache Ovarian Neoplasm Pain Pharmaceutical Product Complaint	Report Source	Product	Role	Manufacturer	Route
800 MG (800 MG, 1 IN 1 D), ORAL			Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Raloxifene Hydrochloride (Raloxifene Hydrochloride)	C		
				Clonazepam (Clonazepam)	C		
				Mirtazapine (Mirtazapine)	C		
				Methadone (Methadone)	C		

Date:08/02/04ISR Number: 4417236-6Report Type:Expedited (15-DaCompany Report #2004049375
Age:49 YR Gender:Male I/FU:I

Outcome Dose Other	Duration	PT Chest Pain Dyspnoea Ventricular Hypertrophy	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin) Levetiracetam	PS		

Date:08/02/04ISR Number: 4417238-XReport Type:Expedited (15-DaCompany Report #2004049614
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Spinal Fusion Acquired	Consumer	Neurontin (Gabapentin)	PS		
				Vicodine (Hydrocodone Bitartrate, Paracetamol)	SS		

Date:08/02/04ISR Number: 4417260-3Report Type:Expedited (15-DaCompany Report #2004049251
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Back Injury Gun Shot Wound	Consumer	Neurontin (Tablets) (Gabapentin)	PS		
(800 MG),		Head Injury Intentional Self-Injury Legal Problem		Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		

(1 IN 2 D),

Freedom Of Information (FOI) Report

Antidepressants
(Antidepressants) C

Date:08/02/04ISR Number: 4417306-2Report Type:Expedited (15-DaCompany Report #2004048837

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anticonvulsant Drug Level	Consumer	Dilantin Kapseals (Phenytoin Sodium)	PS		ORAL
Other		Decreased					
300 MG (100		Arthritis					
MG, 3 IN 1		Blood Pressure Increased					
D), ORAL; 30		Carpal Tunnel Syndrome					
MG (30 MG, 1		Convulsion					
IN 1 D), ORAL		Drug Toxicity		Neurontin (Gabapentin)	SS		ORAL
ORAL		Fall					
		Fatigue		Cardura (Doxazosin Mesilate) (Doxazosin Mesilate)	SS		ORAL
6 MG (6 MG, 1		Hip Arthroplasty					
IN 1 D), ORAL		Memory Impairment					
		Mental Retardation					
		Severity Unspecified		Carbamazepine (Carbamazepine)	SS		
		Multiple Allergies		Oxcarbazepine (Oxcarbazepine)	SS		
		Nausea		Estradiol (Estradiol)	C		
		Neuropathy Peripheral		Nabumetone (Nabumetone)	C		
		Surgical Procedure		Fexofenadine Hydrochloride (Fexofenadine Hydrochloride)	C		
		Repeated		Acetylsalicylic Acid (Acetylsalicylic Acid)	C		

Date:08/03/04ISR Number: 4414082-4Report Type:Expedited (15-DaCompany Report #200412821FR
Age:60 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Cardiac Arrest		Rifadine	PS	Aventis	
Hospitalization -	Epilepsy				Pharmaceuticals Inc.	ORAL
Initial or Prolonged	Hepatic Failure		Oflocet	SS		ORAL
	Ketoacidosis		Glucophage	SS		ORAL
	Lactic Acidosis		Furadantine	SS		ORAL
	Respiratory Distress		Hyperium	SS		ORAL
	Septic Shock		Neurontin	SS		ORAL
			Aprovel	C		ORAL
			Ferritine	C		

Date:08/03/04ISR Number: 4417418-3Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
Age:44 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain Lower
Initial or Prolonged	Abdominal Pain Upper
Disability	Abnormal Behaviour
Other	Acute Sinusitis
	Anger

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Freedom Of Information (FOI) Report

Angioneurotic Oedema
Animal Bite
Ankle Fracture
Anorexia
Antinuclear Antibody
Positive
Anxiety
Arthralgia
Arthritis
Arthropod Bite
Asthenia
Asthma
Astigmatism
Atrial Flutter
Atrial Tachycardia
Back Pain
Bacteraemia
Balance Disorder
Blepharitis
Blepharospasm
Blister
Blood Albumin Decreased
Blood Glucose Increased
Blood Thyroid Stimulating
Hormone Decreased
Blood Urea Decreased
Brain Damage
Breast Mass
Bronchitis
Bronchospasm
Bruxism
Burning Sensation
Cardiovascular Disorder
Carpal Tunnel Syndrome
Cellulitis
Cerebral Disorder
Cervicitis
Change Of Bowel Habit
Chest Pain
Chromaturia
Cognitive Disorder
Condition Aggravated
Constipation
Contusion
Convulsion
Coordination Abnormal
Cough

Crying
Deafness
Depression
Diarrhoea
Diplopia
Disease Recurrence
Dissociative Disorder
Disturbance In Attention
Drug Hypersensitivity
Drug Withdrawal Syndrome
Dry Mouth
Dysarthria
Dysgeusia
Dyskinesia

Freedom Of Information (FOI) Report

Dyspepsia
Dysphagia
Dysphasia
Dyspnoea
Dysuria
Ear Pain
Ecchymosis
Enteritis Infectious
Eustachian Tube
Dysfunction
Eyelid Ptosis
Facial Bones Fracture
Fall
Fatigue
Feeling Abnormal
Feeling Jittery
Fibrocystic Breast
Disease
Flat Affect
Gastrointestinal Disorder
Gastrooesophageal Reflux
Disease
Gingival Pain
Glossodynia
Goitre
Granuloma
Haemorrhage
Hand Fracture
Head Injury
Headache
Heart Rate Increased
Heart Rate Irregular
Hordeolum
Hot Flush
Hypersomnia
Hypoglycaemia
Hypothyroidism
Impaired Healing
Imprisonment
Increased Upper Airway
Secretion
Increased Viscosity Of
Bronchial Secretion
Inflammation
Insomnia
Intentional Misuse
Intervertebral Disc

Degeneration
Intervertebral Disc
Disorder
Irritability
Irritable Bowel Syndrome
Joint Injury
Joint Sprain
Joint Swelling
Keratoconjunctivitis
Sicca
Laboratory Test Abnormal
Lacrimation Increased
Laryngitis
Loss Of Consciousness

Freedom Of Information (FOI) Report

Macroglossia
Malnutrition
Mammogram Abnormal
Mania
Mean Cell Haemoglobin
Increased
Meniere'S Disease
Menorrhagia
Mental Impairment
Mouth Breathing
Multiple Fractures
Muscle Spasms
Muscle Twitching
Myopia
Nasal Discomfort
Nasal Ulcer
Nausea
Nerve Injury
Nervous System Disorder
Nystagmus
Oedema Peripheral
Oesophageal Disorder
Otitis Media
Ovarian Cyst
Ovarian Cyst Ruptured
Ovarian Infection
Ovarian Mass
Pain
Pain In Extremity
Pco2 Decreased
Pelvic Inflammatory
Disease
Pharyngeal Injury
Pharyngitis
Photophobia
Pitting Oedema
Pollakiuria
Polyneuropathy
Propionibacterium
Infection
Pruritus
Psoriasis
Pyrexia
Radiculitis Lumbosacral
Radiculopathy
Rash Erythematous
Red Blood Cell Count

Decreased
Restless Legs Syndrome
Rhinitis Allergic
Road Traffic Accident
Sedation
Sensation Of Pressure
Sensory Disturbance
Sialoadenitis
Sinus Disorder
Sinusitis
Skin Candida
Skin Laceration
Sneezing
Somatisation Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1800 MG	(DAILY) ORAL	Somnolence				
		Speech Disorder				
		Stress				
		Suicidal Ideation	Neurontin			
		Supraventricular	(Gabapentin)	PS		ORAL
		Tachycardia				
		Swelling	Lamotrigine			
		Syncope	(Lamotrigine)	C		
		Tardive Dyskinesia	Lithium Carbonate			
		Tendonitis	(Lithium Carbonate)	C		
		Thermal Burn	Clonazepam			
		Thirst	(Clonazepam)	C		
		Thrombin Time Shortened	Metholphenidate			
		Tinnitus	Hydrochloride			
		Tongue Disorder	(Methylphenidate			
		Tongue Oedema	Hydrochloride)	C		
		Tooth Disorder	Levothyroxine Sodium			
		Tremor	(Levothyroxine			
		Tubo-Ovarian Abscess	Sodium)	C		
		Upper Respiratory Tract	Liothyroxine Sodium			
		Infection	(Liothyroxine			
		Urinary Tract Infection	Sodium)	C		
		Vaginal Candidiasis	Sertraline			
		Vaginal Discharge	Hydrochloride			
		Vaginal Haemorrhage	(Sertraline			
		Vertigo Positional	Hydrochloride)	C		
		Vestibular Neuronitis	Pilocarpine			
		Viral Infection	Hydrochloride			
		Viral Labyrinthitis	(Pilocarpine			
		Vision Blurred	Hydrochloride)	C		
		Visual Field Defect	Metoprolol Succinate			
		Weight Decreased	(Metoprolol			
		Weight Increased	Succinate)	C		
		White Blood Cell Count	Lansoprazole			
		Abnormal	(Lansoprazole)	C		
		White Blood Cell Count	Hyoscyamine Sulfate			
		Decreased	(Hyoscyamine			
		Wound Infection	Sulfate)	C		
		Wrist Fracture	Diltiazem			
			Hydrochloride			
			(Diltiazem			
			Hydrochloride)	C		
			Bupropion			
			Hydrochloride			

(Bupropion Hydrochloride)	C
Celebrex (Celecoxib)	C
Quetiapine Fumarate (Quetiapine Fumarate)	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C
Olanzapine	

Freedom Of Information (FOI) Report

(Olanzapine) C
 Omeprazole
 (Omeprazole) C
 Nasal Preparations
 (Nasal Preparations) C
 Nefazodone
 Hydrochloride
 (Nefazodone
 Hydrochloride) C
 Totolin
 (Guaifenesin,
 Phenylpropanolamine
 Hydrochloride) C
 Meclozine
 (Meclozine) C
 Fluticasone
 Propionate
 (Fluticasone
 Propionate) C
 Trimethobenzamide
 Hydrochloride
 (Trimethobenzamide
 Hydrochloride) C
 Respaire Sr 120
 (Guaifenesin,
 Pseudoephedrine
 Hydrochloride) C
 Risperidone
 (Risperidone) C

Date:08/03/04ISR Number: 4417420-1Report Type:Expedited (15-DaCompany Report #2004050556
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cerebrovascular Accident Hypoaesthesia	Consumer	Neurontin (Gabapentin)	PS		

Date:08/03/04ISR Number: 4417425-0Report Type:Expedited (15-DaCompany Report #2004049671
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Clostridium Colitis	Consumer	Neurontin			

Initial or Prolonged	Dehydration	(Gabapentin)	PS	ORAL
100 MG (100				
Other	Diabetes Mellitus			
MG), ORAL				
	Drug Ineffective	Prednisone		
	Haematoma	(Prednisone)	SS	
	Hepatic Enzyme Increased	Oxybutynin		
	Impaired Healing	(Oxybutynin)	C	
	Neuropathy			
	Pain In Extremity			
	Skin Atrophy			
	Wound			
	Wound Complication			

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Freedom Of Information (FOI) Report

Date:08/03/04ISR Number: 4417435-3Report Type:Expedited (15-DaCompany Report #2004049595

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG (600 MG, 2 IN 1 D), ORAL		Cardiac Disorder Coronary Artery Occlusion Diabetes Mellitus Drug Ineffective Lymphoma Myocardial Infarction Neoplasm Malignant Neuropathy	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Atenolol (Atenolol)	C		
				Isosorbide (Isosorbide)	C		
				Acetylsalicylic Acid)	C		
				Tocopherol (Tocopherol)	C		
				Pyridoxine Hydrochloride (Pyridoxine Hydrochloride)	C		
				Cyanocobalamin (Cyanocobalamin)	C		
				Insulin (Insulin)	C		

Date:08/03/04ISR Number: 4418013-2Report Type:Expedited (15-DaCompany Report #2004049712

Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG		Confusional State Parkinson'S Disease	Foreign Health	Neurontin (Gabapentin)	PS		
			Professional Company Representative	Lamaline (Belladonna Extract, Caffeine, Opium Tincture, Paracetamol)	C		
				Ketoprofen	C		
				All Other Therapeutic Products	C		

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Lamictal	PS	Glaxosmithkline	ORAL
UNKNOWN		Drug Interaction		Gabitril	SS		
UNKNOWN		Hypoglycaemia		Celexa	SS		
UNKNOWN		Vision Blurred		Neurontin	SS		
UNKNOWN				Xanax	SS		
UNKNOWN				Klonopin	SS		

Age:54 YR Gender:Male I/FU:I

Outcome	PT
Required	Abdominal Pain Upper
Intervention to	Cholestasis
Prevent Permanent	Chromaturia
Impairment/Damage	Hepatic Fibrosis
	Hepatotoxicity
	Inflammation
	Jaundice

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Liver Disorder Liver Function Test Abnormal	Report Source	Product	Role	Manufacturer	Route
INITIALLY 100 MG PO TID, DOSE GRADUALLY INCREASED				Gabapentin (Neurontin , Pfizer) 300 Mg Capsules	PS	Pfizer	ORAL

- Humalog C
- Lantus C
- Venlafaxine C
- Kcl C
- Pantoprazole C
- Furosemide C
- Metoprolol Xl C
- Valsartan Xl C
- Atorvastatin C
- Metformin/Glyburide C
- Rosiglitazone C

Date:08/04/04ISR Number: 4416324-8Report Type:Direct Company Report #CTU 224227
 Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG PO TID		Serotonin Syndrome		Gabapentin	PS		ORAL
Initial or Prolonged 100 MG PO Q		Urinary Tract Infection		Sertraline	SS		ORAL

DAILY

Date:08/04/04ISR Number: 4420016-9Report Type:Expedited (15-DaCompany Report #2004044456
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		No Adverse Drug Effect	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
3000 MG, ORAL			Professional	Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Glipizide (Glipizide)	C		
				Gemfibrozil (Gemfibrozil)	C		
				Valsartan (Valsartan)	C		

Date:08/04/04ISR Number: 4421882-3Report Type:Expedited (15-DaCompany Report #2004040264
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Dependence	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Drug Withdrawal Syndrome					
3200 MG, ORAL		Feeling Abnormal		Antihypertensives (Antihypertensives)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/04/04ISR Number: 4421915-4Report Type:Expedited (15-DaCompany Report #2004051390

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Accident	Consumer	Neurontin (Gabapentin)	PS		
		Inflammation		Celecoxib (Celecoxib)	SS		
		Movement Disorder					
		Pain					
		Spinal Cord Disorder					

Date:08/05/04ISR Number: 4421907-5Report Type:Expedited (15-DaCompany Report #2004036523

Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Abasia	Consumer	Neurontin (Tablets)			
Other		Disease Recurrence	Health	(Gabapentin)	PS		ORAL
3200 MG (800		Myalgia	Professional				
MG, 4 IN 1		Pain					
D), ORAL				Amitriptyline (Amitriptyline)	C		
				Pioglitazone (Pioglitazone)	C		
				Glibomet (Glibenclamide, Metformin Hydrochloride)	C		
				Simvastatin (Simvastatin)	C		
				Thyroid (Thyroid)	C		

Date:08/05/04ISR Number: 4422421-3Report Type:Expedited (15-DaCompany Report #2004044358

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Therapeutic Response	Health	Neurontin (Gabapentin)	PS		
		Unexpected	Professional				
300 MG (100							

MG 3 IN 1 D)

Company

Representative

Date:08/05/04ISR Number: 4423684-0Report Type:Expedited (15-DaCompany Report #2004018637

Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Condition Aggravated Difficulty In Walking	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
2700 MG (300 MG, 9 IN 1 D)		Polyneuropathy	Professional				

ORAL

Thioctic Acid (Thioctic Acid)	C
Furosemide (Furosemide)	C
All Other Therapeutic Products	C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/05/04ISR Number: 4423692-XReport Type:Expedited (15-DaCompany Report #2004046701
 Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	160 MG (80 MG, 2 IN 1 D) ORAL	Dizziness Pruritus	Foreign Health	Ziprasidone (Caps) (Ziprasidone)	PS		ORAL
		Scratch	Professional Company				
	1800 MG (600 MG, 3 IN 1 D), ORAL	Torsade De Pointes	Representative	Gabapentin (Gabapentin)	SS		ORAL
	300 MG, ORAL			Lamotrigine (Lamotrigine)	SS		ORAL
	ORAL			Ranitidine Hydrochloride (Ranitidine Hydrochloride)	SS		ORAL
	ORAL			Midazolam (Midazolam)	SS		ORAL
				Dexchlorpheniramine Maleate (Dexchlorpheniramine Maleate)	SS		
				Olanzapine (Olanzapine)	C		

Date:08/05/04ISR Number: 4423950-9Report Type:Expedited (15-DaCompany Report #2004048870
 Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Arrhythmia Bradycardia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL

Other Cardiac Arrest Professional All Other
 Osteomyelitis Therapeutic Products C
 Somnolence

Date:08/05/04ISR Number: 4423986-8Report Type:Expedited (15-DaCompany Report #2004051149
 Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Muscle Spasms Myalgia	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL			Professional	Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Bumetanide (Bumetanide)	C		
				Paracetamol (Paracetamol)	C		
				Codeine (Codeine)	C		
				Isosorbide Mononitrate (Isosorbide Mononitrate)	C		
				Tramadol (Tramadol)	C		
				Perindopril (Perindopril)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Omeprazole
(Omeprazole) C

Date:08/05/04ISR Number: 4423988-1Report Type:Expedited (15-DaCompany Report #2004051150

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hallucination, Visual Panic Attack	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
1200 MG (600 MG, 2 IN 1 D), ORAL							

Date:08/06/04ISR Number: 4421785-4Report Type:Expedited (15-DaCompany Report #2004051401

Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening Other ORAL		Abnormal Dreams Drug Withdrawal Syndrome Treatment Noncompliance	Foreign Health Professional	Gabapentin (Gabapentin) Venlafaxine (Venlafaxine) Buprenorphine (Buprenorphine)	PS C C		ORAL

Date:08/06/04ISR Number: 4421793-3Report Type:Expedited (15-DaCompany Report #CIP04001641

Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death Hospitalization - Initial or Prolonged 3 DOSES DAILY, ORAL		Cardiac Arrest Epilepsy Hepatic Failure Hepatocellular Damage Ketosis	Foreign Health Professional Other	Furadantine(Nitrofur antoin, Macrocrystals) Capsule	PS		ORAL

	Lactic Acidosis	Glucophage "Abic"		
	Respiratory Distress	(Metformin		
	Septic Shock	Hydrochloride)	SS	ORAL
3000 MG				
DAILY, ORAL				
		Oflocet (Ofloxacin)	SS	ORAL
2 DOSES				
DAILY, ORAL				
		Neurontin (Gabapentin		
)	SS	ORAL
2 DOSES				
DAILY, ORAL				
		Ferritin (Ferritin)	SS	
		Aprovel (Irbesartan)	SS	
		Rifadine (Rifampicin)	SS	ORAL
6 DOSES				
DAILY, ORAL				
		Hyperium (Rilmenidine		
)	C	ORAL
2 DOSES				
DAILY, ORAL				

Date: 08/06/04 ISR Number: 4422080-X Report Type: Expedited (15-Da Company Report #2004051416)

Age: Gender: Male I/FU: I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Insomnia	Consumer	Neurontin (Gabapentin)	PS		
				Alprazolam	C		
				Oxycodone			

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Freedom Of Information (FOI) Report

Hydrochloride
(Oxycodone
Hydrochloride) C
Pethidine
Hydrochloride
(Pethidine
Hydrochloride) C
Methadone
(Methadone) C
Esomeprazole
(Esomeprazole) C

Date:08/06/04ISR Number: 4422142-7Report Type:Expedited (15-DaCompany Report #2004051397
Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder Convulsion Speech Disorder	Health Professional	Dilantin Suspension (Phenytoin Sodium) Neurontin (Gabapentin)	PS SS		

Date:08/06/04ISR Number: 4422518-8Report Type:Expedited (15-DaCompany Report #2004021501
Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder Depression	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
300 MG		Drug Interaction	Professional				
(DAILY), ORAL		Fall Feeling Drunk		Escitalopram (Escitalopram)	SS		
20 MG		Hallucination, Visual					
(DAILY),		Migraine Nightmare Suicide Attempt		Allegra-D (Fexofenadine Pseudoephedrine Hydrochloride) Propacet (Dextropropoxyphene Napsilate,	C		

Paracetamol) C
 Pantoprazole C
 (Pantoprazole) C
 Docusate (Docusate) C

Date:08/06/04ISR Number: 4423382-3Report Type:Expedited (15-DaCompany Report #2004050911

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Anorexia Condition Aggravated	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
50 MG (1 IN 1 D) ORAL		Depression Diabetes Mellitus	Professional	Zoloft (Sertraline)	SS		ORAL
				Antihypertensives (Antihypertensives)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/04ISR Number: 4418602-5Report Type:Periodic
Age:52 YR Gender:Female I/FU:F

Company Report #US-GLAXOSMITHKLINE-A0413109A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Agitation	Consumer	Paxil	PS	Glaxosmithkline	ORAL
		Alopecia		Neurontin	SS		
300MG Four		Anger					
times per day	YR	Drug Withdrawal Syndrome		Wellbutrin Sr	SS	Glaxosmithkline	ORAL
150MG Per day	5 WK	Dysphonia		Trazodone	C		
		Hypersensitivity		Klonopin	C		
		Influenza Like Illness					
		Insomnia					
		Nausea					
		Paraesthesia					
		Performance Status					
		Decreased					
		Smoker					
		Somnolence					

Date:08/09/04ISR Number: 4424025-5Report Type:Expedited (15-DaCompany Report #2004044272
Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cytolytic Hepatitis	Foreign Health	Neurontin (Tabelts)	PS		ORAL
Other		Drug Interaction		(Gabapentin)			
1800 MG (3 IN		Prothrombin Time	Professional				
1 D), ORAL		Prolonged	Company Representative	Phenobarbital Sodium	SS		ORAL
				(Phenobarbital Sodium)			
300 MG (2 IN							
1 D), ORAL				Clonazepam	C		
				(Clonazepam)			
				Valproate Sodum	C		
				(Valproate Sodum)			
				Ceftazidime			
				Penahydrate			

(Ceftazidime Pentahydrate)	C
Fosfomicin (Fosfomicin)	C
Heparin-Fraction, Sodium Slat (Heparin-Fraction, Sodium Salt)	C
Spironolactone (Spironolactone)	C
Furosemide (Furosemide)	C
Amlodipine Besilate (Amlodipine Besilate)	C
Atenolol (Atenolol)	C
Cefotaxime Sodium (Cefotaxime Sodium)	C
Tramadol Hydrochloride (Tramadol Hydrochloride)	C
Pefloxacin Mesilate (Pefloxacinmesilate)	C
Domperidone	

Freedom Of Information (FOI) Report

(Domperidone) C
 Metronidazole
 (Metronidazole) C
 Amphotericin B
 (Amphotericin B) C
 Nystatin (Nystatin) C
 Paracetamol
 (Paracetamol) C
 Clonazepam
 (Clonazepam) C

Date:08/09/04ISR Number: 4424080-2Report Type:Expedited (15-DaCompany Report #2004042400

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Breast Cancer	Foreign	Neurontin			
Other		Nipple Disorder	Consumer	(Gabapentin)	PS		ORAL
200 MG (100		Nodule	Health				
MG, 2 IN 1D)		Refusal Of Treatment By	Professional				
, ORAL		Patient	Company				
		Treatment Noncompliance	Representative				

Date:08/09/04ISR Number: 4424254-0Report Type:Expedited (15-DaCompany Report #2004051408

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bladder Disorder	Consumer	Neurontin			
Other		Constipation		(Gabapentin)	PS		ORAL
3000 MG (300		Faecaloma					
MG), ORAL		Renal Cyst		Clonazepam			
		Urinary Retention		(Clonazepam)	SS		ORAL
300 MG, ORAL		Urinary Tract Infection		Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	SS		ORAL
60 MG, ORAL							

ORAL

Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS	ORAL
Quetiapine Fumarate (Quetiapine Fumarate)	C	
Valsartan (Valsartan)	C	
Macrogol (Macrogol)	C	

Date:08/09/04ISR Number: 4424258-8Report Type:Expedited (15-DaCompany Report #2004051359

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Hypersensitivity	Consumer	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/04ISR Number: 4424269-2Report Type:Expedited (15-DaCompany Report #2004051194
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain Dizziness	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG (1 D), ORAL		Visual Acuity Reduced					

Date:08/09/04ISR Number: 4424273-4Report Type:Expedited (15-DaCompany Report #2004042915
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthropathy Back Disorder Balance Disorder Bone Disorder Drug Ineffective	Consumer	Neurontin (Gabapentin) All Other Therapeutic Products (All Other Therapeutic Products)	PS C		

Date:08/09/04ISR Number: 4424283-7Report Type:Expedited (15-DaCompany Report #2004027040
Age:68 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Calcium Increased Delusion	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
2400 MG (600 MG, 4 IN 1 D), ORAL		Feeling Abnormal Hallucination, Visual	Professional				
SUBCUTANEOUS SUBCUTANEOUS	(1 IN 1 D),			Teriparatide (Teriparatide) Anaesthetics (Anaesthetics)	SS SS		

Morphine (Morphine)	SS
Enalapril	C
Nifedipine	C
One-A-Day (Ascorbic Acid, Cyanocobalamin, Ergocalciferol, Nicotianamide, Chlorphenamine Maleate	C
Calcium Magnesium Zinc	C
Risedronate Sodium	C
Amlodipine Besilate	C
Temazepam	C

Date:08/09/04ISR Number: 4424303-XReport Type:Expedited (15-DaCompany Report #2004040264
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Abuser	Consumer	Neurontin			
Other		Drug Withdrawal Syndrome	Health	(Gabapentin)	PS		ORAL
3200 MG, ORAL		Feeling Abnormal	Professional	Antihypertensives	C		
				All Other Therapeutic	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/04ISR Number: 4424333-8Report Type:Expedited (15-DaCompany Report #2004051029

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Anxiety	Consumer	Neurontin			
Other		Multiple Fractures Pain Skeletal Injury		(Gabapentin)	PS		

Date:08/09/04ISR Number: 4424338-7Report Type:Expedited (15-DaCompany Report #2004051193

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anaemia Drug Interaction	Health Professional	Neurontin (Gabapentin) Warfarin (Warfarin)	PS SS		

Date:08/09/04ISR Number: 4425006-8Report Type:Expedited (15-DaCompany Report #2004051197

Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 3600 MG (3 IN 1 D), ORAL		Cardiac Disorder Creatinine Renal Clearance Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Lisinopril (Lisinopril)	C		
				Atorvastatin (Atorvastatin)	C		
				Tenofibrate (Tenofibrate)	C		
				Ranitidine Hydrochloride (Ranitidine Hydrochloride)	C		
				Hydromorphone Hydrochloride (Hydromorphone Hydrochloride)	C		

Multivitamins
(Ascorbic Acid,
Ergocaliferol, Folic
Acid, Nocotinamide,
Panthenol, Retinol, C
Metoprolol Tartrate
(Metoprolol
Tartrate) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C
Sertraline
Hydrochloride
(Sertraline
Hydrochloride) C
Amlodipine Besilate
(Amlodipine
Besilate) C
Hydrochlorothiazide
(Hydrochlorothiazide
) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/09/04ISR Number: 4425011-1Report Type:Expedited (15-DaCompany Report #2004047157

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased Emotional Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (100 MG, 3 IN 1 D), ORAL		Growth Accelerated Pharmaceutical Product Complaint					

Date:08/09/04ISR Number: 4425012-3Report Type:Expedited (15-DaCompany Report #2004038647

Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coordination Abnormal Dyskinesia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
3200 MG, ORAL		Electroencephalogram Abnormal Eye Disorder Speech Disorder Vision Blurred	Professional	Citalopram Hydrobromide (Citalopram Hydrobromide) Alprazolam (Alprazolam) All Other Therapeutic Products	C C C		

Date:08/10/04ISR Number: 4422203-2Report Type:Expedited (15-DaCompany Report #US-ROCHE-376030

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Klonopin	PS	Roche	
UNKNOWN		Drug Interaction		Lamictal	I		
UNKNOWN		Hypoglycaemia		Neurontin	I		
UNKNOWN							

UNKNOWN Vision Blurred Xanax I
 UNKNOWN Gabitril I
 UNKNOWN Celexa I

Date:08/10/04ISR Number: 4425490-XReport Type:Expedited (15-DaCompany Report #2004052386
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Arthritis Spinal Column Stenosis	Consumer	Neurontin (Gabapentin)	PS		

Date:08/10/04ISR Number: 4425492-3Report Type:Expedited (15-DaCompany Report #2004052371
 Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 MG, 2 IN 1 D), ORAL		Blood Magnesium Decreased Blood Pressure Increased Dehydration Feeling Abnormal Flushing Oedema Peripheral	Consumer	Neurontin (Gabapentin) Furosemide (Furosemide) Metoprolol Succinate (Metoprolol Succinate) Quinapril Hydrochloride	PS C C		ORAL

Freedom Of Information (FOI) Report

(Quinapril	
Hydrochloride)	C
Digoxin (Digoxin)	C
Levothyroxine Sodium	
(Levothyroxine	
Sodium)	C
Simvastatin	
(Simvastatin)	C
Clopidogrel Sulfate	
(Clopidogrel	
Sulfate)	C
Diclofenac Sodium	
(Diclofenac Sodium)	C
Estrogens Esterified	
(Estrogens	
Esterified)	C
Acetylsalicylic Acid	
(Acetylsalicylic	
Acid)	C
Fish Oil (Fish Oil)	C
General Nutrients	
(General Nutrients)	C
All Other	
Therapeutic Products	
(All Other	
Therapeutic	
Products)	C
Calcium Citrate	
(Calcium Citrate)	C
Multivitamins	
(Ascorbic Acid,	
Ergocalciferol,	
Folic Acid,	
Nicotinamide,	C
Vitamin B (Vitamin	
B)	C
Ascorbic Acid	
(Ascorbic Acid)	C
Magnesium	
(Magnesium)	C

Date:08/10/04ISR Number: 4425814-3Report Type:Expedited (15-DaCompany Report #2004033685
 Age:43 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					

Death Other ORAL	Completed Suicide Drug Level Above Therapeutic Intentional Misuse Muscle Relaxant Drug Level Above Therapeutic Victim Of Homicide	Foreign Health Professional	Neurontin (Gabapentin) Baclofen (Baclofen)	PS SS	ORAL
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Date:08/10/04ISR Number: 4425842-8Report Type:Expedited (15-DaCompany Report #2004046607
 Age:66 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600		Hypertensive Crisis Visual Disturbance	Foreign Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL

MG, 3 IN 1

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Freedom Of Information (FOI) Report

D), ORAL

Insulin (Insulin) C
 Moduretic "Msd"
 (Amiloride
 Hydrochloride,
 Hydrochlorothiazide) C

Date:08/10/04ISR Number: 4426351-2Report Type:Expedited (15-DaCompany Report #A02200402328
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Agranulocytosis Iatrogenic Injury	Health Professional	Equanil - (Meprobamate) - Tablet - Unit Dose : Unknown	PS		ORAL
1 UNIT OD; 1							
MONTH - TIME							
TO ONSET	26	DAY					
10 MG OD; 1				Stilnox - (Zolpidem) - Tablet - 10 Mg	SS		ORAL
MONTH - TIME							
TO ONSET	26	DAY					
1 UNIT TID; 1				Phosphalugel - (Aluminium Phosphate Gel) - Unknown - Unit Dose : Unknown	SS		ORAL
MONTH - TIME							
TO ONSET	26	DAY					
26 DAY				Athymil - (Mianserin Hydrochloride) - Tablet - Unit Dose : Unknown	SS		ORAL
				Mopral - (Omeprazole) - Capsule - Unit Dose			

1 UNIT OD; 1 : Unknown SS

MONTH - TIME

TO ONSET 26 DAY

Neurontin -
 (Gabapentin) -
 Unknown - Unit Dose
 : Unknown SS ORAL

1 UNIT TID; 1

MONTH - TIME

TO ONSET 26 DAY

Date:08/11/04ISR Number: 4426669-3Report Type:Expedited (15-DaCompany Report #2004051426
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Consumer	Neurontin			
		Drug Interaction		(Gabapentin)	PS		
		Hypoglycaemia		Lamotrigine			
		Vision Blurred		(Lamotrigine)	SS		
				Alprazolam			
				(Alprazolam)	SS		
				Citalopram			
				Hydrobromide			
				(Citalopram			
				Hydrobromide)	SS		
				Clonazepam			
				(Clonazepam)	SS		

Freedom Of Information (FOI) Report

Tiagabine
Hydrochloride
(Tiagabine
Hydrochloride) C

Date:08/11/04ISR Number: 4426675-9Report Type:Expedited (15-DaCompany Report #2004051708
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (300 MG, 1 IN 1 D), ORAL		Difficulty In Walking Gait Disturbance Nerve Compression Pain In Extremity Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:08/11/04ISR Number: 4426676-0Report Type:Expedited (15-DaCompany Report #2004051692
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 800 MG (400 MG, 2 IN 1 D), ORAL		Asthenia Depression Feeling Abnormal Intentional Self-Injury Joint Injury Social Avoidant Behaviour Suicidal Ideation Therapy Non-Responder Treatment Noncompliance Weight Decreased	Consumer	Neurontin (Gabapentin) Carbamazepine (Carbamazepine) Oxcarbazepine (Oxcarbazepine) Levetiracetam (Levetiracetam)	PS SS SS SS		ORAL ORAL

Date:08/11/04ISR Number: 4426685-1Report Type:Expedited (15-DaCompany Report #2004047491
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Arteriovenous Malformation	Health Professional	Neurontin (Gabapentin)	PS		ORAL
800 MG (400 MG, 2 IN 1 D), ORAL		Depression					
		Dizziness					
		Fatigue Headache Irritability Pain		Paroxetine Hydrochloride (Paroxetine Hydrochloride)	SS		ORAL
ORAL		Treatment Noncompliance		Estrogens Conjugated (Estrogens Conjugated)	C		

Date:08/11/04ISR Number: 4426687-5Report Type:Expedited (15-DaCompany Report #2004040406
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dyskinesia Road Traffic Accident	Health Professional	Neurontin (Gabapentin)	PS		
900 MG (300 MG, 3 IN 1 D)			Company Representative				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/11/04ISR Number: 4426766-2Report Type:Expedited (15-DaCompany Report #2004052381

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness	Consumer	Neurontin (Gabapentin)	PS		
		Fibromyalgia		Rofecoxib (Rofecoxib)	SS		
		Spinal Cord Injury		All Other Therapeutic Products (All Other Therapeutic Products)	SS		
				All Other Therapeutic Products (All Other Therapeutic Products)	SS		
				Allergy Medication (Allergy Medication)	C		

Date:08/11/04ISR Number: 4427051-5Report Type:Expedited (15-DaCompany Report #2004052118

Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Anxiety	Consumer	Neurontin (Gabapentin)	PS		
Other		Completed Suicide					
		Medication Error					
		Pain					

Date:08/11/04ISR Number: 4427056-4Report Type:Expedited (15-DaCompany Report #2004038659

Age:38 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Acne	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Localised Infection	Health				
ORAL		Pharmaceutical Product Complaint	Professional	Omeprazole	C		
				Paracetamol	C		
		Pharyngeal Mass		Naproxen Sodium	C		

Scar
Staphylococcal Infection

Date:08/12/04ISR Number: 4427391-XReport Type:Expedited (15-DaCompany Report #2004032235
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Abasia	Foreign	Neurontin			
Other		Arthralgia	Health	(Gabapentin)	PS		ORAL
900 MG (300		Immobile	Professional				
MG, 3 IN 1		Weight Increased	Company				
D), ORAL			Representative	Hydrocortisone (Hydrocortisone)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/12/04ISR Number: 4427806-7Report Type:Expedited (15-DaCompany Report #2004052383
Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (400 Other MG, 3 IN 1 D), ORAL	Abdominal Pain Upper Constipation Diarrhoea Gastrointestinal Erosion Intestinal Obstruction Mastication Disorder Nausea Oral Discomfort Oral Fungal Infection Vitamin B12 Decreased	Consumer	Neurontin (Tablets) (Gabapentin) Clonazepam (Cloazepam) Venlafaxine Hydrochloride (Venlafaxine	PS C C		ORAL

Date:08/12/04ISR Number: 4427812-2Report Type:Expedited (15-DaCompany Report #2004026318
Age:69 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Diabetes Mellitus Difficulty In Walking Eczema Gastrointestinal Disorder Localised Infection Loss Of Consciousness Nerve Injury Pain Paraesthesia Spinal Operation	Consumer Health Professional	Neurontin (Gabapentin) Celecoxib (Celecoxib) All Other Therapeutic Products (All Other Therapeutic Products) Insulin (Insulin) .	PS SS SS SS SS		

Date:08/12/04ISR Number: 4427857-2Report Type:Expedited (15-DaCompany Report #2004052246
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization -	Atrophy	Consumer	Dilantin Suspension			

Initial or Prolonged	Body Height Decreased	(Phenytoin Sodium)	PS	ORAL
ORAL				
Other	Convulsion	Neurontin		
	Rash Generalised	(Gabapentin)	SS	ORAL
ORAL	Speech Disorder			

Date:08/13/04ISR Number: 4425032-9Report Type:Expedited (15-DaCompany Report #200412821FR

Age:60 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Cardiac Arrest		Rifadine	PS	Aventis	
Hospitalization -	Epilepsy				Pharmaceuticals Inc.	ORAL
Initial or Prolonged	Hepatic Failure		Oflocet	SS		ORAL
	Ketoacidosis		Glucophage	SS		ORAL
	Lactic Acidosis		Furadantine	SS		ORAL
	Respiratory Distress		Hyperium	SS		ORAL
	Septic Shock		Neurontin	SS		ORAL
			Aprovel	C		ORAL
			Ferritin	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/13/04ISR Number: 4425060-3Report Type:Expedited (15-DaCompany Report #PHRM2004FR02530
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Alanine Aminotransferase Increased		Nisis	PS	Novartis Sector: Pharma	ORAL
Other		Aspartate		Di-Antalvic	SS		ORAL
6 caps/day	7200 MIN	Aminotransferase		Augmentin	SS		ORAL
2 g/day	4320 MIN	Increased		Neurontin	SS		ORAL
3 DF/day		Blood Alkaline Phosphatase Increased Cholelithiasis Gamma-Glutamyltransferase Increased Hepatitis Cholestatic		Detensiel	C		ORAL

Date:08/13/04ISR Number: 4425099-8Report Type:Expedited (15-DaCompany Report #JP-JNJFOC-20040801088
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Health	Ofloxacin	PS		
OROPHARINGEAL Hospitalization - OROPHARINGEAL		Epilepsy	Professional	Rifadine	SS		
Initial or Prolonged OROPHARINGEAL		Hepatic Failure		Glucophage	SS		
OROPHARINGEAL		Hepatocellular Damage		Hyperium	SS		
OROPHARINGEAL		Ketoacidosis		Neurontin	SS		
OROPHARINGEAL		Lactic Acidosis		Furadantine	SS		
OROPHARINGEAL		Respiratory Distress		Aprovel	C		
OROPHARINGEAL		Septic Shock		Ferritin	C		

Date:08/13/04ISR Number: 4425415-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0341969A
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1G Twice per Initial or Prolonged day		Blood Alkaline Phosphatase Increased	Consumer	Augmentin	PS	Glaxosmithkline	ORAL
2UNIT Three times per day 5 DAY		Cytolytic Hepatitis Gamma-Glutamyltransferase Increased		Di Antalvic Neurontin	SS		ORAL
1UNIT Three times per day				Nisis	SS		ORAL
1UNIT per day				Detensiel	C		ORAL

Date:08/13/04ISR Number: 4428013-4Report Type:Expedited (15-DaCompany Report #2004052685
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Agitation Condition Aggravated Convulsion	Consumer	Neurontin (Gabapentin) Clonazepam (Clonazepam) Diazepam (Diazepam)	PS SS SS		

Date:08/13/04ISR Number: 4428016-XReport Type:Expedited (15-DaCompany Report #2004052217
Age:52 YR Gender:Female I/FU:I

Outcome	PT
Other	Abnormal Behaviour Activities Of Daily

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Living Impaired Blood Pressure Increased Condition Aggravated Dizziness	Report Source	Product	Role	Manufacturer	Route
300 MG (300 MG, 1 IN 1 D), ORAL AS NEEDED (5 MG), ORAL		Drug Interaction Fatigue Feeling Abnormal Hangover Hypersomnia Insomnia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Lumbar Puncture Abnormal Pain Pharmaceutical Product Complaint Renal Impairment Speech Disorder Thyroid Disorder Vision Blurred		Levothyroxine Sodium (Levothyroxine Sodium) Estrogens Conjugated (Estrogens Conjugated) Potassium Chloride (Potassium Chloride) Furosemide (Furosemide) Hyzaar (Hydrochlorothiazide , Losartan Potassium) Vicodin (Hydrocodone Bitartrate, Paracetamol)	C C C C		

Date:08/13/04ISR Number: 4428029-8Report Type:Expedited (15-DaCompany Report #2004032979
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG (600 MG, 3 IN 1 D), ORAL		Beta Haemolytic Streptococcal Infection Cellulitis Condition Aggravated	Consumer Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL

Drug Ineffective
Fasciitis
Headache
Ligament Disorder
Neuralgia
Neuropathy
Pain In Extremity
Pharmaceutical Product
Complaint

Estrogens Conjugated C
Levothyroxine Sodium C
All Other
Therapeutic Products C

Date:08/13/04ISR Number: 4428093-6Report Type:Expedited (15-DaCompany Report #2004052733
Age:47 YR Gender:Female I/FU:I

Outcome PT
Hospitalization - Abnormal Behaviour
Initial or Prolonged Agitation
Other Anger
Feeling Abnormal
Homicidal Ideation
Insomnia
Marital Problem
Paranoia
Relationship Breakdown
Thinking Abnormal

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Treatment Noncompliance Victim Of Spousal Abuse	Report Source	Product	Role	Manufacturer	Route
ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Methylphenidate Hydrochloride	SS		
				All Other Therapeutic Products	SS		

Date:08/13/04ISR Number: 4428094-8Report Type:Expedited (15-DaCompany Report #2004053486
Age: Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Arthralgia Pain In Extremity	Consumer	Neurontin (Gabapentin)	PS		
					Celecoxib (Celecoxib)	SS		
					Morphine Sulfate (Morphine Sulfate)	SS		
					Carisoprodol (Carisoprodol)	SS		
					Naproxen (Naproxen)	SS		
					Rofecoxib (Rofecoxib)	SS		
					Dextroprppoxyphene (Dextroprppoxyphene0	C		

Date:08/13/04ISR Number: 4428423-5Report Type:Expedited (15-DaCompany Report #HQWYE693302AUG04
Age:55 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 4 TABLETS			Cholestasis Cytolytic Hepatitis	Health Professional	Advil (Ibuprofen, Tablet)	PS		ORAL
DAILY ORAL	4	DAY		Other	Cortancyl (Prednisone,)	SS		

4	DAY	Doliprane (Paracetamol,)	SS
		Neurontin (Gabapentin,)	SS
		Triflucan (Fluconazole,)	SS
4	DAY	Temodal (Temozolomide)	C

Date:08/16/04ISR Number: 4426851-5Report Type:Expedited (15-DaCompany Report #PHNU2004DE00766
Age: Gender:Male I/FU:F

Outcome	PT
Disability	Abnormal Behaviour
Other	Alopecia
	Anosmia
	Aphasia
	Bradyphrenia
	Cerebellar Syndrome
	Chills
	Cogwheel Rigidity

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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
400 mg, QOD		Depression Difficulty In Walking	Health Professional	Tegretal	PS	Novartis Sector: Pharma	ORAL
UNKNOWN		Dyskinesia Electroencephalogram		Ergenyl "Sanofi-Synthelabo"	SS		
UNKNOWN		Abnormal		Neurontin	SS		
60mg/day		Erectile Dysfunction		Gabitril	SS		ORAL
100mg/day	549 DAY	Fall		Topamax	SS		ORAL
400 mg, QID		Fatigue		Carbabeta	SS		ORAL
400 mg, QID		Gait Disturbance		Carbabeta	SS		ORAL
1DF/day		General Physical Health		Movergan	SS		ORAL
.5 mg, QID		Deterioration Grand Mal Convulsion		Levodopa Parkotil	SS SS		ORAL ORAL
1mg/day		Hypokinesia		Parkotil	SS		ORAL
0.20mg/day		Liver Function Test		Parkotil	SS		ORAL
200 mg, QID		Abnormal		Comtess	SS		ORAL
50 mg, QID		Malaise Masked Facies		Amantadin Levodopa	SS SS		ORAL ORAL
62.5 mg, TID		Memory Impairment		Madopar	SS		ORAL
400 mg, TID		Mental Impairment		Carbabeta	SS		ORAL
UNKNOWN	3DF/Day	Micrographia		Zentropil	C		
UNKNOWN	1000-1500mg/d	Mobility Decreased		Keppra	C		
ay		Motor Dysfunction					
UNKNOWN		Movement Disorder Muscle Spasms		Nacom "Dupont Pharma"	C		

15mg/day	366 DAY	Muscle Twitching Myalgia	Tasmar Remergil	C C	ORAL
5mg/day		Nerve Conduction Studies Abnormal Neuroleptic Malignant Syndrome Paralysis	L-Dopa Levetiracetam Selegiline Phenhydan Keppra	C C C C C	
UNKNOWN	500 mg, BID	Parkinson'S Disease Performance Status	Lamictal "Glaxosmithkline"	 C	
UNKNOWN		Decreased Posture Abnormal Regressive Behaviour Salivary Hypersecretion Simple Partial Seizures Speech Disorder Tongue Paralysis Tremor			

Date:08/16/04ISR Number: 4429427-9Report Type:Expedited (15-DaCompany Report #2004044295
Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Brain Damage	Foreign	Neurontin			
Hospitalization - 800 MG (400 Initial or Prolonged MG, 2 IN 1 Other D), ORAL		Cerebrovascular Accident	Consumer	(Gabapentin)	PS		ORAL
		Diabetes Mellitus					
		Pneumonia					
		Pyrexia		Enalapril Maleate (Enalapril Maleate) Metformin Hydrochloride (Metformin Hydrochloride) Glibenclamide	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Glibenclamide) C

Date:08/17/04ISR Number: 4427760-8Report Type:Expedited (15-DaCompany Report #FR-BRISTOL-MYERS SQUIBB COMPANY-12652871
 Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Health	Glucophage	PS	Bristol-Myers Squibb	
Hospitalization - Initial or Prolonged		Epilepsy	Professional	Aprovel	SS	Bristol-Myers Squibb Company	ORAL
		Ketosis		Furadantine	SS		ORAL
		Lactic Acidosis		Rifadine	SS		
		Septic Shock		Oflocet	SS		
				Hyperium	SS		
				Neurontin	SS		
				Ferritin	SS		

Date:08/17/04ISR Number: 4429237-2Report Type:Direct Company Report #CTU 225059
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Depression		Neurontin 300mg			
		Insomnia		Parke-Davis	PS	Parke-Davis	ORAL
1 TWICE A DAY		Pain					
BY MOUTH		Pharmaceutical Product Complaint					

Date:08/17/04ISR Number: 4429828-9Report Type:Expedited (15-DaCompany Report #2004054371
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Arthritis	Consumer	Neurontin			
Other		Asthma		(Gabapentin)	PS		
		Mobility Decreased					
		Neuropathy					
		Thyroid Disorder					

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin			
Other		No Adverse Drug Effect	Consumer	(Gabapentin)	PS		
				Diclofenac Sodium			
				(Diclofenac Sodium)	SS		
				Propacet			
				(Dextropropoxyphene			
				Napsilate,			
				Paracetamol)	SS		
				Cyclobenzaprine			
				Hydrochloride			
				(Cyclobenzaprine			
				Hydrochloride)	SS		
				All Other			
				Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/17/04ISR Number: 4429838-1Report Type:Expedited (15-DaCompany Report #2004053609
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other 600 MG (300 MG, 2 IN 1 D), ORAL		Activities Of Daily Living Impaired Blister Burns Second Degree Dermatitis Exfoliative Insomnia Movement Disorder Pain Sunburn	Consumer	Neurontin (Gabapentin) Lansoprazole (Lansoprazole)	PS C		ORAL

Date:08/17/04ISR Number: 4429884-8Report Type:Expedited (15-DaCompany Report #2004051359
Age:84 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG, ORAL		Dizziness Feeling Abnormal Headache Hypersensitivity Nausea	Consumer	Neurontin (Gabapentin) Amlodipine Besilate (Amlodipine Besilate)	PS C		ORAL

Date:08/17/04ISR Number: 4429889-7Report Type:Expedited (15-DaCompany Report #2004053220
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2400 MG (600 MG, 4 IN 1 D), ORAL		Bone Pain Burning Sensation Drug Ineffective Multiple Sclerosis Pharmaceutical Product	Consumer	Neurontin (Gapabentin) Lisinopril	PS		ORAL

Complaint

(Lisinopril)

C

Date:08/17/04ISR Number: 4429890-3Report Type:Expedited (15-DaCompany Report #2004053222
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 150 MG (300 Initial or Prolonged MG), ORAL Other ORAL		Intentional Misuse	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Temazepam (Temazepam)	SS		ORAL
				Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		

Date:08/18/04ISR Number: 4429320-1Report Type:Direct
Age:53 YR Gender:Male I/FU:I

Company Report #CTU 225187

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
30 MG QHS		Drug Interaction		Methadone 10 Mg	PS		ORAL
ORAL		Overdose					
600 MG TID				Gabapentin 300 Mg	SS		ORAL
ORAL				Vicodin Prn	C		
				Aspirin	C		
				Docusate	C		
				Hydroxyzine	C		

Date:08/18/04ISR Number: 4429495-4Report Type:Direct Company Report #CTU 225157
 Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Asthenia		Gabapentin	PS		ORAL
300 MG TID PO 10 DAY							
Initial or Prolonged		Coordination Abnormal		Oxycontin	C		
		Fall		Oxycodone	C		
		Lumbar Radiculopathy		Lipitor	C		
		Urinary Tract Infection		Levaquin	C		
				Zantac	C		
				Advil	C		
				Xeloda	C		
				Reglan	C		
				Flomax	C		
				Loprox	C		
				Aranesp	C		

Date:08/18/04ISR Number: 4430582-5Report Type:Expedited (15-DaCompany Report #2004053846
 Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Difficulty In Walking	Foreign	Neurontin			

900 MG (300
 MG, 3 IN 1
 D), ORAL

Intervertebral Disc
 Protrusion

Consumer
 (Gabapentin)
 PS
 ORAL

Date:08/18/04ISR Number: 4430602-8Report Type:Expedited (15-DaCompany Report #2004051401
 Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Abnormal Dreams	Foreign	Gabapentin			
Other		Drug Withdrawal Syndrome	Health	(Gabapentin)	PS		ORAL
ORAL			Professional	Venlafaxine			
				(Venlafaxine)	C		
				Buprenorphine			
				(Buprenorphine)	C		

Date:08/18/04ISR Number: 4431228-2Report Type:Expedited (15-DaCompany Report #2004053447
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Exanthem	Foreign	Neurontin			
1800 MG (600		Leukocytoclastic	Health	(Gabapentin)	PS		ORAL
MG, 3 IN 1		Vasculitis	Professional				
D), ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Rofecoxib
 (Rofecoxib) C
 Fluvastatin Sodium
 (Fluvastatin Sodium) C

Date:08/18/04ISR Number: 4431920-XReport Type:Expedited (15-DaCompany Report #2004049614
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neuralgia	Consumer	Neurontin (Gabapentin)	PS		
		Spinal Fusion Acquired		Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		

Date:08/18/04ISR Number: 4432027-8Report Type:Expedited (15-DaCompany Report #2004048122
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 900 MG		Asthenia	Consumer	Neurontin (Gabapentin)	PS		ORAL
Initial or Prolonged (300MG,3 IN 1 Other D), ORAL		Depressed Level Of Consciousness	Professional				
		Drug Interaction		All Other Therapeutic Proucts (All Other Therapeutic Products)	SS		
		Dyspnoea		Trazodone (Trazodone	C		
		Dysuria		Levothyroxine Sodium (Levothyroxine Sodium)	C		
		Emphysema		Spiroonolactone (Spiroonolactone)	C		
		Fall		Venlafaxine Hydrochloride (Venlfxine Hydrochloride)	C		
		Feeling Abnormal					
		Knee Operation					
		Large Intestinal Haemorrhage					
		Lung Disorder					
		Respiratory Disorder					
		Treatment Noncompliance					

Lorazepam	
(Lorazepam)	C
Tolterodine-L-Tartrate	
(Tolterodine	
L-Tartrate)	C
Warfarin (Warfarin)	C
Hydrocodone	
(Hydrocodone)	C
Risedronate Sodium	
(Risedronate Sodium)	C
Esomeprazole	
(Esomeprazole)	C
Diazepam (Diazepam)	C
Prednisone	
(Prednisone)	C
Risperidone	
(Risperidone)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/18/04ISR Number: 4432030-8Report Type:Expedited (15-DaCompany Report #2004041004
 Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Anxiety Drug Hypersensitivity	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
2400 MG (800 MG, 3 IN 1 D), ORAL		Drug Ineffective Muscle Spasms	Professional				
		Pharmaceutical Product Complaint Post Procedural Complication Tremor Weight Decreased Weight Increased		Bupropion Hydrochloride (Bupropion Hydrochloride) Venlafaxine Hydrochloride (Venlafaxine Hydrochloride) Levothyroxine Sodium (Levothyroxine Sodium) Baclofen (Baclofen)	SS C C C		

Date:08/18/04ISR Number: 4432033-3Report Type:Expedited (15-DaCompany Report #2004047823
 Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Balance Disorder Blood Triglycerides	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
200 MG (100 MG, 2 IN 1 D), ORAL		Increased Body Temperature	Professional				
		Increased Flatulence		Dilantin Suspension (Phenytoin Sodium)	SS		ORAL
ORAL		Rash Simple Partial Seizures		Carbamazepine (Carbamazepine) Phenobarbital (Phenobarbital) Vitamins (Vitamins)	SS SS C		

Date:08/18/04ISR Number: 4432103-XReport Type:Expedited (15-DaCompany Report #2004053224
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cholestasis	Health	Gabapentin			
		Hepatic Fibrosis	Professional	(Gabapentin)	PS		ORAL
2700 MG (3 IN		Hepatic Trauma					
1D), ORAL		Hepatitis		Atorvastatin			
		Hepatobiliary Disease		(Atorvastatin)	SS		
		Hepatotoxicity		Metformin			
				(Metformin)	SS		
				Glibenclamide			
				(Glibenclamide)	SS		
				Rosiglitazone			
				(Rosiglitazone)	SS		

Date:08/18/04ISR Number: 4432125-9Report Type:Expedited (15-DaCompany Report #2004054409
Age: Gender:Female I/FU:I

Outcome	PT
Disability	Back Pain
	Gait Disturbance

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Freedom Of Information (FOI) Report

Dose	Duration	Neck Pain Pain Pain In Extremity	Report Source	Product	Role	Manufacturer	Route
300 MG			Consumer	Neurontin (Gabapentin)	PS		
				Metamucil (Glucose Monhydrate, Ispaghula Husk)	C		
				Herbal Nos/Vitamins Nos (Herbal Nos, Vitamins Nos)	C		

Date:08/18/04ISR Number: 4432141-7Report Type:Expedited (15-DaCompany Report #2004052685
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Tablets) (Gabapentin)	PS		
Hospitalization - Initial or Prolonged Other		Agitation Convulsion		Clonazepam (Clonazepam) Diazepam	SS SS		

Date:08/20/04ISR Number: 4432176-4Report Type:Expedited (15-DaCompany Report #2004054453
Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Nervous System Disorder Tremor	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
300 MG (300 MG, 1 IN 1 D), ORAL				Senna (Senna) Ferrous Fumarate (Ferrous Fumarate) Nitrazepam (Nitrazepam) Furosemide	C C C		

(Furosemide)	C
Gliclazide	
(Gliclazide)	C
Omeprazole	
(Omeprazole)	C
Amlodipine	
(Amlodipine)	C
Calcium Carbonate	
(Calcium Carbonate)	C
Risedronate Sodium	
(Risedronate Sodium)	C

Date:08/20/04ISR Number: 4432230-7Report Type:Expedited (15-DaCompany Report #2004056012
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cheyne-Stokes Respiration	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/20/04ISR Number: 4432554-3Report Type:Expedited (15-DaCompany Report #2004054849
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Ketosis Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (300 MG, 1 IN 1 D), ORAL							

Date:08/20/04ISR Number: 4432562-2Report Type:Expedited (15-DaCompany Report #2004037133
 Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other		Amnesia Drug Effect Decreased Intentional Misuse Irritable Bowel Syndrome	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
3200 MG (1600 MG, 2 IN 1 D), ORAL							

Abacavir Sulfate (Abacavir Sulfate)	C
Atazanavir Sulfate (Atazanavir Sulfate)	C
Stavudine (Stavudine)	C
Ritonavir (Ritonavir)	C
Desloratadine (Desloratadine)	C
Triamcinolone Acetonide (Triamcinolone Acetonide)	C
Aciclovir (Aciclovir)	C
Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid,	

Nicotinamide,

C

Date:08/20/04ISR Number: 4432566-XReport Type:Expedited (15-DaCompany Report #2004046590

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Health	Neurontin (Tablets)			
		Grand Mal Convulsion	Professional	(Gabapentin)	PS		
800 MG		Pharmaceutical Product		Phenytoin Sodium			
		Complaint		(Phenytoin Sodium)	C		
		Pregnancy					

Date:08/20/04ISR Number: 4432582-8Report Type:Expedited (15-DaCompany Report #2004054101

Age:56 YR Gender:Female I/FU:I

Outcome	PT
Other	Blood Pressure
	Fluctuation
	Dehydration
	Diplopia
	Disorientation
	Heart Rate Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Kidney Infection Rash	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1 D), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Hyzaar (Hydrochlorothiazide , Losartan Potassium) Hyoscyamine (Hyoscyamine) Oxycodone Hydrochloride (Oxycodone Hydrochloride) Fentanyl (Fentanyl) Alprazolam (Alprazolam) Prednisone (Prednisone) Warfarin (Warfarin)	C C C C C C C		

Date:08/20/04ISR Number: 4432595-6Report Type:Expedited (15-DaCompany Report #2004054705

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Bone Density Decreased Diabetes Mellitus Fibromyalgia Neoplasm Malignant Osteoarthritis Spinal Operation	Consumer	Neurontin (Gabapentin) Valdecoxib (Valdecoxib) Vicoprofen (Hydrocodone Bitartrate, Ibuprofen) All Other Therapeutic Products (All Other Therapeutic	PS SS SS		

Products)

SS

Date:08/20/04ISR Number: 4432651-2Report Type:Expedited (15-DaCompany Report #2004055008

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Consumer	Neurontin			
		Pain In Extremity		(Gabapentin)	PS		
				Hydrocortisone			
				(Hydrocortisone)	SS		

Date:08/23/04ISR Number: 4430820-9Report Type:Expedited (15-DaCompany Report #CH-BRISTOL-MYERS SQUIBB COMPANY-12670501

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

Outcome
Hospitalization -
Initial or Prolonged
Disability

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Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness Nystagmus		Methotrexate	PS	Bristol-Myers Squibb Company	
INTRATHECAL		Pain		Neurontin	SS		ORAL
5	DAY						

Date:08/23/04ISR Number: 4431210-5Report Type:Direct Company Report #CTU 225424
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 1/PER 12 HRS		Asthenia Coordination Abnormal Dizziness Dry Mouth Somnolence Speech Disorder Trismus Vision Blurred		Neurontin 300mg	PS		

Date:08/23/04ISR Number: 4432647-0Report Type:Expedited (15-DaCompany Report #2004054607
 Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other (300 MG), ORAL		Stevens-Johnson Syndrome Toxic Epidermal Necrolysis	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Atenolol (Atenolol) Metformin (Metfomrin) Potassium Chloride (Potassium Chloride) Nifedipine (Nifedipine) Initard (Insulin,	C C C C		

Insulin Injection,	
Isophane)	C
Calcitrol	
(Calcitrol)	C
All Otehr	
Therapeutic Rproducts	
(All Other	
Therapeutic	
Products)	C
Rofecoxib	
(Rofecoxib)	C
Tocopherol	
(Tocopherol)	C
Simvastatin	
(Simvastatin)	C
Probenecid	
(Probenecid)	C
Loratadine	
(Loratadine)	C
Perindopril	
(Perindopril)	C
Famotidine	
(Famotidine)	C
Gliclazide	

Freedom Of Information (FOI) Report

(Gliclazide) C
 Prochlorperazine
 (Prochlorperazine) C
 Paracetamol
 (Paracetamol) C
 Calcium (Calcium) C

Date:08/23/04ISR Number: 4432926-7Report Type:Expedited (15-DaCompany Report #2004048216
 Age:12 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Agitation Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL		Drug Ineffective Encephalopathy	Professional Company				
		Grand Mal Convulsion Psychomotor Retardation	Representative	Carbamazepine (Carbamazepine)	C		

Date:08/23/04ISR Number: 4433154-1Report Type:Expedited (15-DaCompany Report #2004051692
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Asthenia Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL
800 MG (400 MG, 2 IN 1 D), ORAL		Feeling Abnormal Intentional Self-Injury					
		Social Avoidant Behaviour Suicidal Ideation		Carbamazepine (Carbamazepine0)	SS		ORAL
ORAL		Therapeutic Response Decreased Treatment Noncompliance Weight Decreased		Oxcarbazepine (Oxcarbazepine) Levetiracetam (Levetiracetam)	SS SS SS		

Date:08/23/04ISR Number: 4433163-2Report Type:Expedited (15-DaCompany Report #2004054836
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthma	Consumer	Neurontin			
		Condition Aggravated		(Gabapentin)	PS		
		Fatigue		Geodon (Ziprasidone			
		Pain In Jaw		Hydrochloride)			
40 MG		Sexual Dysfunction		(Ziprasidone)	SS		
		Somnolence		Risperidone			
		Vision Blurred		(Risperidone)	SS		
				Paracetamol			
				(Paracetamol)	SS		
				Olanzapine			
				(Olanzapine)	SS		
				Aripiprazole			
				(Aripiprazole)	SS		

Date:08/23/04ISR Number: 4433164-4Report Type:Expedited (15-DaCompany Report #2004054695
Age:45 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Angina Pectoris
Initial or Prolonged	Hypoacusis
Other	Myocardial Infarction

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Freedom Of Information (FOI) Report

Dose	Duration	Stent Placement Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Glibenclamide (Glibenclamide)	C		

Date:08/23/04ISR Number: 4433166-8Report Type:Expedited (15-DaCompany Report #2004046521
Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 1900 MG (2 IN Initial or Prolonged 1 D), ORAL Other		Dyspnoea Pericardial Effusion Pericarditis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Zolpidem Tartrate (Zolpidem Tartrate)	C		
				Metformin Hydrochloride (Metformin Hydrochloride)	C		
				Atorvastatin (Atorvastatin)	C		
				Quinapril Hydrochloride (Quinapril Hydrochloride)	C		
				Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
				Quetiapine Fumarate (Quetiapine Fumarate)	C		

Date:08/23/04ISR Number: 4433168-1Report Type:Expedited (15-DaCompany Report #2004051708
Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Back Pain Difficulty In Walking	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (300 MG, 1 IN 1 D), ORAL		Gait Disturbance Headache Muscle Spasms Neck Pain Nerve Compression					

Date:08/23/04ISR Number: 4433175-9Report Type:Expedited (15-DaCompany Report #2004054882
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Upper Caesarean Section	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG, ORAL		Drug Exposure During Pregnancy Placental Disorder Therapeutic Agent Toxicity					

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FDA - Adverse Event Reporting System (AERS)

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Date:08/23/04ISR Number: 4433176-0Report Type:Expedited (15-DaCompany Report #2004054890

Age:1 DY Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Aspiration	Consumer	Neurontin			
Hospitalization -	Atrial Septal Defect		(Gabapentin)	PS		
TRANSPLACENTAL 300 MG, Initial or Prolonged PLACENTAL	Drug Exposure During					
Congenital Anomaly	Pregnancy					
Other	Feeding Problem In Newborn Gastrooesophageal Reflux Disease Hepatic Failure Neonatal Cardiac Failure Neonatal Disorder Neonatal Respiratory Distress Syndrome Premature Baby Respiratory Syncytial Virus Infection Thrombocytopenia Neonatal Tracheal Disorder Ventricular Septal Defect					

Date:08/23/04ISR Number: 4433177-2Report Type:Expedited (15-DaCompany Report #2004056021

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Acute Myeloid Leukaemia	Consumer	Neurontin			
Initial or Prolonged	Blood Disorder		(Gabapentin)	PS		ORAL
7200 MG (6 IN						
Other	Colitis					
1 D), ORAL	Dehydration		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		
			Topiramate (Topiramate)	C		
			Folic Acid (Folic Acid)	C		

Paracetamol	
(Paracetamol)	C
Metronidazole	
(Metronidazole)	C
Aciclovir	
(Aciclovir)	C
Levothyroxine Sodium	
(Levothyroxine Sodium)	C
Calcium Carbonate	
(Calcium Carbonate)	C
Lactulose	
(Lactulose)	C
Ciprofloxacin	
(Ciprofloxacin)	C
Bactrim	
(Sulfamethoxazole, Trimethoprim)	C

Freedom Of Information (FOI) Report

Date:08/24/04ISR Number: 4433273-XReport Type:Expedited (15-DaCompany Report #DRON00204002739

Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - DAILY PO	Duration Blood Glucose Increased	Health	Marinol (Dronabinol)	PS		ORAL
Initial or Prolonged SUBCUTANEOUS	200 IU QD SC Blood Magnesium Decreased	Professional	Aranesp (Aranesp)	SS		
INTRAVENOUS	1 MG QD IV Blood Sodium Decreased		Kytril (Granisetron)	SS		
20 MG BID PO	Confusional State Contusion		Oxycontin (Oxycodone Hydrochloride)	SS		ORAL
DAILY PO	Convulsion Eye Injury		Neurontin (Gabapentin)	SS		ORAL
DAILY PO	Face Injury Fall		Aricept (Donepezil Hydrochloride)	SS		ORAL
	Gastrointestinal		Morphine Sulfate			
	Haemorrhage		(Morphine Sulfate)	C		
	Haemolytic Anaemia		Chlorpromazine			
	Liver Transplant		(Chlorpromazine)	C		
	Rejection		Oxycodone			
	Mental Status Changes		(Oxycodone)	C		
	Metabolic Encephalopathy		Prograf (Prograf)	C		
	Nausea		Protonix (Protonix)	C		
	Pancytopenia		Prandin "Kuhn"			
	Vomiting		(Deflazacort)	C		
			Zoloft (Sertraline Hydrochloride)	C		
			Glipizide			
			(Glipizide)	C		
			Procardia			
			(Nifedipine)	C		
			Magnesium			
			(Magnesium)	C		
			Levaquin			
			(Levofloxacin)	C		
			Pentamidine			
			(Pentamidine)	C		
			Pericolace			
			(Pericolace)	C		
			Dulcolax (Bisacodyl)	C		
			Asa (Acetylsalicylic Acid)	C		

Lasix (Lasix)	C
Leukine	
(Sargramostim)	C
Dexamethasone	
(Dexamethasone)	C
Adriamycin	
(Doxorubicin)	C
Cisplatinum	
(Cisplatin)	C
Senecot (Senecot)	C
Inderal (Inderal)	C
Multivitamins	
(Multivitamins)	C
Insulin (Insulin)	C
Aldactone	
(Aldactone)	C
Megace (Megace)	C

FDA - Adverse Event Reporting System (AERS)

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Date:08/24/04ISR Number: 4433292-3Report Type:Expedited (15-DaCompany Report #2004054853
 Age:65 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 Other MG, 3 IN 1 D)	Intentional Misuse Irritability Mood Swings Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		
ILL-DEFINED DISORDER			Naproxen Sodium (Naproxen Sodium)	SS		
			Insulin (Insulin) All Othe Therapeutic Products (All Other Therapeutic Products (All Other Therapeutic	C C		

Date:08/24/04ISR Number: 4433294-7Report Type:Expedited (15-DaCompany Report #2004056018
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Amnesia Dysstasia Medical Device Complication Nerve Injury Sleep Disorder Due To General Medical Condition, Insomnia Type Weight Increased	Health Professional	Neurontin (Gabapentin) All Other Therapeutic Products (All Other Therapeutic Products)	PS SS		

Date:08/24/04ISR Number: 4433295-9Report Type:Expedited (15-DaCompany Report #2004054101
 Age:56 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Blood Pressure	Consumer	Neurontin			

900 MG (300	Fluctuation	(Gabapentin)	PS	ORAL
MG, 3 IN 1	Consciousness Fluctuating			
D), ORAL	Dehydration			
	Diplopia	Hyzaar		
	Disorientation	(Hydrochlorothiazide,		
	Dry Mouth	Losartan Potassium)	C	
	Fatigue	Hyoscyamine	C	
	Feeling Abnormal	Oxycodone		
	Heart Rate Increased	Hydrochloride		
	Kidney Infection	(Oxycodone		
	Pruritus Generalised	Hydrochloride)	C	
	Rash	Fentanyl (Fentanyl)	C	
	Somnolence	Alprazolam		
		(Alprazolam)	C	
		Prednisone		
		(Prednisone)	C	
		Warfarin (Warfarin)	C	
		All Other		
		Therapeutic Products		
		(All Other		
		Therapeutic		
		Products)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/24/04ISR Number: 4433303-5Report Type:Expedited (15-DaCompany Report #2004056093
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG, 3 IN 1 D), ORAL		Hallucination, Auditory Refusal Of Treatment By Patient Tinnitus	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
				Baclofen (Baclofen)	C		
				Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		

Date:08/24/04ISR Number: 4433362-XReport Type:Expedited (15-DaCompany Report #2004228700GB
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other INTRAMUSCULAR 3 MONTHS, 1ST INJ., INTRAMUSCULAR 600 MG, QD	150 MG, EVERY	Drug Interaction Unintended Pregnancy	Foreign Health Professional Other	Depo-Provera (Medroxy progesterone Acetate) Suspension, Sterile, 150mg/Ml	PS		
				Gabapentin Nm Pharma (Gabapentin) Capsule	SS		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 100 MG (100 Other MG), ORAL	Clostridium Colitis Dehydration Diabetes Mellitus Drug Ineffective Haematoma Hepatic Enzyme Increased Injury Neuropathy Pain Skin Atrophy Wound	Consumer Health Professional	Neurontin (Gabapentin) Prednisone (Prednisone) Oxybutynin	PS SS C		ORAL

Outcome	PT
Hospitalization - Initial or Prolonged Other	Abnormal Behaviour Asthenia Bronchitis Chest Pain Cognitive Disorder Electroencephalogram

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Dose	Duration	Abnormal Loss Of Consciousness Memory Impairment Overdose	Report Source	Product	Role	Manufacturer	Route
ORAL		Paraesthesia Personality Change	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Pyrexia Screaming Somnolence Stomach Discomfort Surgical Procedure Repeated Thinking Abnormal Vomiting		Tetracycline (Tetracycline) Erythromycin (Erythromycin) Oxycodone Hydrochloride (Oxycodone Hydrochloride) Morphine Sulfate (Morphine Sulfate0 Clonidine (Clonidine) Fentanyl Citrate (Fentanyl Citrate) Lansoprazole (Lansoprazole) Sucralfate (Sucralfate) Cevimeline Hydrochloride (Cevimeline Hydrochloride) Digoxin (Digoxin) Levothyroxine Sodium (Levothyroxine Sodium) Oxybutynin Hydrochloride (Oxybutynin Hydrochloride) Meloxicam (Meloxicam) Metaxalone (Metaxalone) Furosemide 9furosemide) Calcitonin, Salmon (Calcitonin, Salmon)	SS C		

Age:60 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Cardiac Arrest		Ofloxacin	PS		
OROPHARINGEAL	41 DAY					
Hospitalization -	Epilepsy		Rifadine	SS		
OROPHARINGEAL	41 DAY					
Initial or Prolonged	Hepatocellular Damage		Glucophage	SS		
OROPHARINGEAL						
OROPHARINGEAL	Ketoacidosis		Furadantine	SS		
	9 DAY					
OROPHARINGEAL	Lactic Acidosis		Neurontin	SS		
	41 DAY					
OROPHARINGEAL	Respiratory Distress		Hyperium	C		
	30 DAY					
OROPHARINGEAL	Septic Shock		Aprovel	C		
			Ferritin	C		

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Freedom Of Information (FOI) Report

Date:08/25/04ISR Number: 4432338-6Report Type:Expedited (15-DaCompany Report #GB-BRISTOL-MYERS SQUIBB COMPANY-12680518
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma		Holoxan	PS	Bristol-Myers Squibb Company	
INTRAVENOUS							DRIP
1 DAY							
INTRAVENOUS		4 DAY		Ketamine	SS		
INTRAVENOUS		4 DAY		Diamorphine	SS		
				Amitriptyline	SS		ORAL
				Gabapentin	SS		ORAL
				Movicol	C		

Date:08/25/04ISR Number: 4433588-5Report Type:Direct Company Report #CTU 225605
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
SEE ITEM B-5		Alopecia		Kenalog	PS		
SEE ITEM B-5		Dermatologic Examination Abnormal		Neurontin	SS		
				Prednisone	SS		
		Erythema		Methotrexate	SS		

Date:08/25/04ISR Number: 4433909-3Report Type:Expedited (15-DaCompany Report #2004056058
 Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Disorientation	Consumer	Neurontin (Tablets)			
600 MG (600		Dysstasia		(Gabapentin)	PS		ORAL
MG, 1 IN 1		Fall					
D), ORAL		Vision Blurred					
				Vicodin (Hydrocodone			

Bitartrate,
 Paracetamol) C
 Paracetamol
 (Paracetamol) C
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C

Date:08/25/04ISR Number: 4433991-3Report Type:Expedited (15-DaCompany Report #2004056325

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 900 MG (300 MG, 3 IN 1 D)		Complex Regional Pain Syndrome	Consumer	Neurontin (Gabapentin)	PS		
		Nerve Injury					
30 MG (30 MG, 1 IN 1 D)		Upper Limb Fracture		Morphine Sulfate (Morphine Sulfate)	SS		
(2 IN 1 D)				Macrogol (Macrogol)	SS		
(3 IN 1 D)				All Other Therapeutic Products)	SS		
(1 IN 1 D)				Thyroid Hormones (Thyroid Hormones)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/25/04ISR Number: 4434118-4Report Type:Expedited (15-DaCompany Report #2004055469

Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Hospitalization - ORAL	Initial or Prolonged	Epilepsy	Professional	Metformin Hydrohchloride (Metformin Hydrohchloride)	SS		ORAL
3 GRAM (1 IN 1 D), ORAL		Hepatic Failure Ketoacidosis Lactic Acidosis Respiratory Distress					
		Septic Shock		Rifampicin (Rifampicin)	SS		ORAL
ORAL				Nitrofurantoin (Nitrofurantoin)	SS		ORAL
ORAL				Ofloxacin (Ofloxacin)	SS		ORAL
ORAL				Rilmenidine (Rilmenidine)	C		ORAL
ORAL				Irbesartan (Irbesartan)	C		
				Ferrous Sulfate (Ferrous Sulfate)	C		
				Ascorbic Acid (Ascorbic Acid)	C		

Date:08/25/04ISR Number: 4434125-1Report Type:Expedited (15-DaCompany Report #2004056068

Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Interaction	Foreign Health	Gabapentin (Gabapentin)	PS		
600 MG (600 MG, 1 IN 1 D)			Professional	Medroxyprogesterone			

Acetate
(Medroxyprogesterone
Acetate) SS

INTRAMUSCULAR (150 MG, 1 IN

3 M),

INTRAMUSCULAR

Date:08/25/04ISR Number: 4434270-0Report Type:Expedited (15-DaCompany Report #2004056323

Age:53 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged (3 IN 1 D), ORAL	Agranulocytosis Bone Marrow Myelogram Abnormal Neutropenia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
(1 IN 1 D), ORAL			Meprobamate (Meprobamate)	SS		ORAL
(3 IN 1 D), ORAL			Aluminium Phosphate Gel (Aluminium Phosphate Gel)	SS		ORAL
(1 IN 1 D), ORAL			Mianserin Hydrochloride (Mianserin Hydrochloride)	SS		ORAL
			Omeprazole			

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Freedom Of Information (FOI) Report

(1 IN 1 D),	(Omeprazole)	SS	ORAL
ORAL			
(1 IN 1 D),	Zolpidem (Zolpidem)	SS	ORAL
ORAL			

Date:08/25/04ISR Number: 4434291-8Report Type:Expedited (15-DaCompany Report #2004056256
 Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Dysphagia	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Emotional Distress Mental Impairment Mood Altered Reflexes Abnormal Sleep Disorder					

Date:08/25/04ISR Number: 4434295-5Report Type:Expedited (15-DaCompany Report #2004056277
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 100 MG, ORAL		Acute Generalised Exanthematous Pustulosis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Aphthous Stomatitis Clostridial Infection Enterobacter Infection General Physical Health	Professional	Pyridoxine Hydrochloride (Pyridoxine Hydrochloride)	SS		
750 MG (25- MG, TID)		Deterioration					
		Inflammation Oedema Peripheral Pain In Extremity Rash Erythematous		Ondansetron Hydrochloride Docetaxel (Docetaxel)	SS SS		
INTRAVENOUS	110 MG						

(CYCLIC),
 INTRAVENOUS
 Skin Lesion
 Staphylococcal Infection

INTRAVENOUS 70 MG
 Doxorubicin Hydrochloride SS

(CYCLIC),
 INTRAVENOUS

Date:08/26/04ISR Number: 4433364-3Report Type:Expedited (15-DaCompany Report #PHRM2004FR02530
 Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	80 mg/day	Alanine Aminotransferase Increased		Nisis	PS	Novartis Sector: Pharma	ORAL
Other	6 caps/day	Aspartate		Di-Antalvic	SS		ORAL
	7200 MIN	Aminotransferase		Augmentin	SS		ORAL
	2 g/day	Increased		Neurontin	SS		ORAL
	3 DF/day	Blood Alkaline Phosphatase Increased Cholelithiasis Cholestasis Cytolytic Hepatitis Gamma-Glutamyltransferase Increased		Detensiel	C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/26/04ISR Number: 4433681-7Report Type:Periodic
Age:53 YR Gender:Male I/FU:I

Company Report #US-GLAXOSMITHKLINE-A0516125A

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
.5MG Per day	2 MON	Musculoskeletal Pain		Avodart	PS	Glaxosmithkline	ORAL
				Neurontin	SS		
				Prilosec	C		
				Zocor	C		
				Vicodin	C		

Date:08/26/04ISR Number: 4435751-6Report Type:Expedited (15-DaCompany Report #2004056095
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Insomnia	Consumer	Neurontin (Gabapentin)	PS		
900 MG (300 MG, 3 IN 1 D), UNKNOWN		Nausea Nightmare Post Procedural Complication		Hydrochlorothiazide (Hydrochlorothiazide) Potassium (Potassium)	C C		

Date:08/26/04ISR Number: 4435752-8Report Type:Expedited (15-DaCompany Report #2004049226
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache Ovarian Neoplasm	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
800 MG (800 MG, 1 IN 1 D), ORAL		Pain Pharmaceutical Product Complaint	Professional	Raloxifene			

Hydrochloride	
(Raloxifene Hydrochloride)	C
Clonazepam (Clonazepam)	C
Mirtazapine (Mirtazapine)	C
Methadone (Methadone)	C

Date:08/26/04ISR Number: 4435753-XReport Type:Expedited (15-DaCompany Report #2004056122
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dizziness	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL				All Other Therapeutic Products	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/26/04ISR Number: 4435763-2Report Type:Expedited (15-DaCompany Report #2004049375
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Health	Neurontin			
Other		Dyspnoea	Professional	(Gabapentin)	PS		
		Ventricular Hypertrophy		Levetiracetam			
				(Levetiracetam)	C		

Date:08/26/04ISR Number: 4435764-4Report Type:Expedited (15-DaCompany Report #2004048837
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anticonvulsant Drug Level	Consumer	Dilantin Kapseals			
Other		Above Therapeutic		(Phenytoin Sodium)	PS		ORAL
300 MG (100		Anticonvulsant Drug Level					
MG, 3 IN 1		Decreased					
D), ORAL; 30		Arthritis					
MG (30 MG, 1		Asthenia					
IN 1 D), ORAL		Blood Pressure Increased		Neurontin			
ORAL		Bradyphrenia		(Gabapentin)	SS		ORAL
		Carpal Tunnel Syndrome		Cardura (Doxazosin			
		Condition Aggravated		Mesilate) (Doxazosin			
		Convulsion		Mesilate)	SS		ORAL
6 MG (6 MG,1		Fall					
IN 1 D), ORAL		Fatigue		Carbamazepine			
		Hip Arthroplasty		(Carbamazepine)	SS		
		Hypersensitivity		Oxcarbazepine			
		Hypoaesthesia		(Oxcarbazepine)	SS		
		Injury		Estradiol			
		Memory Impairment		(Estradiol)	C		
		Neuropathy Peripheral		Nabumetone			
		Surgical Procedure		(Nabumetone)	C		
		Repeated		Fexofenadine			
				Hydrochloride			

(Fexofenadine
Hydrochloride) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C

Date:08/27/04ISR Number: 4437078-5Report Type:Expedited (15-DaCompany Report #2004056279
Age:30 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser Euphoric Mood	Health Professional Company Representative	Neurontin (Tablets) (Gabapentin)	PS		

Date:08/27/04ISR Number: 4437084-0Report Type:Expedited (15-DaCompany Report #2004057241
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Hypoaesthesia Intervertebral Disc Compression Spinal Deformity	Consumer	Neurontin (Gabapentin) Ibuprofen (Ibuprofen) Acetylsalicylic Acid	PS SS		

Freedom Of Information (FOI) Report

(Acetylsalicylic Acid) SS
 Naproxen Sodium (Naproxen Sodium) SS
 Vicodin (Hydrocodone Bitartrate, Paracetamol) SS
 Codeine (Codeine) SS

Date:08/27/04ISR Number: 4437105-5Report Type:Expedited (15-DaCompany Report #2004057028
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:08/27/04ISR Number: 4437107-9Report Type:Expedited (15-DaCompany Report #2004025426
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Creatinine Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other		Blood Pressure Diastolic					
800 MG (400 MG, 2 IN 1 D), ORAL		Decreased					
		Chest Pain					
		Constipation		Isosorbide Dinitrate			
		Dehydration					
		Delirium		(Isosorbide Dinitrate)	C		
		Diarrhoea		Salbutamol	C		
		Feeling Abnormal		(Salbutamol)	C		
		General Physical Health Deterioration		Furosemide	C		
		Pain		(Furosemide)	C		
		Psychotic Disorder		Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
				Methazolamide (Methazolamide)	C		
				Paroxetine			

Hydrochloride	
(Paroxetine	
Hydrochloride)	C
Irbesartan	
(Irbesartan)	C
Oxycocet (Oxycodone	
Hydrochloride,	
Paracetamol)	C
Fentanyl (Fentanyl)	C
Insulin Injection,	
Isophane (Insulin	
Injection, Isophane)	C
Paracetamol	
(Paracetamol)	C
Potassium Chloride	
(Potassium Chloride)	C
Prednisolone	
(Prednisolone)	C
Glyceryl Trinitrate	
(Glyceryl	
Trinitrate)	C

Freedom Of Information (FOI) Report

Loperamide
 Hydrochloride
 (Loperamide
 Hydrochloride) C
 Zolpidem Tartrate
 (Zolpidem Tartrate) C
 Docusate (Docusate) C
 Clonidine
 (Clonidine) C
 Lactulose
 (Lactulose) C
 Pantoprazole
 (Pantoprazole) C
 Metoprolol
 (Metoprolol) C
 Metolazone
 (Metolazone) C
 Amlodipine
 (Amlodipine) C
 Amitriptyline
 Hydrochloride
 (Amitriptyline
 Hydrochloride) C

Date:08/27/04ISR Number: 4437115-8Report Type:Expedited (15-DaCompany Report #2004058040
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Ineffective Injury	Consumer	Neurontin (Gabapentin)	PS		

Date:08/27/04ISR Number: 4437612-5Report Type:Expedited (15-DaCompany Report #2004PK01394
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 800 MG TID PO		Condition Aggravated Hyperkeratosis	Foreign Health	Beloc Zok Renagel	PS SS		ORAL
300 MG DAILY		Morphoea	Professional	Neurontin	SS		

Movement Disorder

Other

Phos-Ex

C

Einsalpha

C

Vigantoletten

C

Restex

C

Rohypnol

C

Marcoumar

C

Date:08/30/04ISR Number: 4438686-8Report Type:Expedited (15-DaCompany Report #2004051708

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

Outcome

PT

Other

Back Pain

Difficulty In Walking

Headache

Muscle Spasms

Neck Pain

Nerve Compression

Pain In Extremity

Pharmaceutical Product

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Freedom Of Information (FOI) Report

Complaint
Weight Decreased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG (300 MG, 1 IN 1 D), ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:08/30/04ISR Number: 4438688-1Report Type:Expedited (15-DaCompany Report #2004058034
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other		Deformity Drug Ineffective Ill-Defined Disorder Injury Off Label Use Pain Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:08/30/04ISR Number: 4438698-4Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
Age:44 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability Other	Acrochordon Actinic Keratosis Acute Sinusitis Albumin Globulin Ratio Decreased Alcoholism Angina Unstable Anxiety Aphasia Back Pain Bipolar Disorder Bladder Disorder Blood Calcium Decreased Blood Triglycerides

Increased
Blood Urea
Nitrogen/Creatinine Ratio
Increased
Bone Cyst
Brain Neoplasm
Bronchitis Acute
Cardiac Arrest
Carotid Artery Disease
Cerebrovascular Disorder
Chest Wall Mass
Cholecystitis
Cholelithiasis
Cognitive Disorder
Condition Aggravated
Condyloma Acuminatum
Conversion Disorder
Convulsion
Coordination Abnormal
Cyst
Deafness Unilateral

Freedom Of Information (FOI) Report

Depressed Level Of
Consciousness
Depression
Diabetes Mellitus
Difficulty In Walking
Diplopia
Dissociative Disorder
Disturbance In Attention
Drug Ineffective
Dysstasia
Ear Discomfort
Emotional Disorder
Enchondromatosis
Ependymoma
Exostosis
Exposure To Toxic Agent
Fall
Feeling Abnormal
Feeling Of Despair
Fibromyalgia
Folliculitis
Gastritis
Gastrointestinal Disorder
Postoperative
Gastroesophageal Reflux
Disease
Glioma
Gynaecomastia
Headache
Hepatic Steatosis
High Density Lipoprotein
Decreased
Hyperhidrosis
Hypersomnia
Impaired Driving Ability
Insomnia
Intervertebral Disc
Displacement
Intervertebral Disc
Protrusion
Joint Injury
Lower Gastrointestinal
Haemorrhage
Marital Problem
Memory Impairment
Meniscus Lesion
Mental Disorder

Monocyte Count Decreased
Monocyte Percentage
Increased
Musculoskeletal Stiffness
Myocardial Infarction
Nervous System Disorder
Neurodegenerative
Disorder
Obstructive Airways
Disorder
Oedema
Oedema Peripheral
Osteoarthritis
Osteopenia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration		Report Source	Product	Role	Manufacturer	Route
900 MG (1D), ORAL		Pain Paraesthesia Patellofemoral Pain	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Petit Mal Epilepsy	Professional				
		Pharyngolaryngeal Pain Plantar Fasciitis Post Procedural Complication Protein Total Decreased Scar Seborrhoeic Keratosis Sensory Disturbance Sick Sinus Syndrome Sinusitis Stress Suicidal Ideation Synovitis Tinnitus Tonsillar Disorder Vision Blurred Visual Disturbance Weight Decreased	Company Representative	Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin) Celecoxib (Celecoxi9b) Lansoprazole (Lansoprazole)	SS C C C C C		

Date:08/30/04ISR Number: 4438705-9Report Type:Expedited (15-DaCompany Report #2004058042
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Anxiety Carpal Tunnel Syndrome Completed Suicide Drug Ineffective Injury Neuropathy Peripheral Pain	Consumer	Neurontin (Gabapentin)	PS		

Date:08/30/04ISR Number: 4438719-9Report Type:Expedited (15-DaCompany Report #2004051359
Age:84 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Upper Dizziness	Consumer	Neurontin (Gabapentin)	PS		ORAL
(300 MG), ORAL		Drug Hypersensitivity Feeling Abnormal Headache Nausea		Amlodipine Besilate (Amlodipine Besilate)	C		

Date:08/30/04ISR Number: 4438955-1Report Type:Expedited (15-DaCompany Report #2004057101
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Disease Recurrence Hyperkeratosis Morphoea	Foreign Health Professional	Neurontin (Gabapentin) Metoprolol Succinate (Metoprolol Succinate) Sevelamer	PS SS		

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Freedom Of Information (FOI) Report

Hydrochloride (Sevelamer Hydrochloride)	SS
Calcium Acetate (Calcium Acetate)	C
Alfacalcidol (Alfacalcidol)	C
Colecalciferol (Colecalciferol)	C
Madopar (Benserazide Hydrochloride, Levodopa)	C
Flunitrazepam (Flunitrazepam)	C
Phenprocoumon (Phenprocoumon)	C

Date:08/30/04ISR Number: 4438958-7Report Type:Expedited (15-DaCompany Report #055-0945-M0100011
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia	Foreign	Gabapentin			
Hospitalization - 900 MG, ORAL		Cardiac Disorder	Consumer	(Gabapentin)	PS		ORAL
Initial or Prolonged		Cardiovascular Disorder		Cardiovascular			
Other		Cerebrovascular Accident		System Drugs			
		Gastrointestinal Disorder		(Cardiovascular			
		Metastases To		System Drugs)	C		
		Gastrointestinal Tract		Urologicals			
		Oedema Peripheral		(Urologicals)	C		
		Prostate Cancer		Antihypertensives			
		Metastatic		(Antihypertensives)	C		
				All Other			
				Therapeutic Products	C		

Date:08/30/04ISR Number: 4438959-9Report Type:Expedited (15-DaCompany Report #2004058288
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Cholelithiasis Cholestasis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL

Other	Cytolytic Hepatitis	Professional	Aporex (Dextropropoxyphene Hydrochloride, Paracetamol)	SS	ORAL
ORAL			Clavulin (Amoxicillin Trihydrate, Clavulanate Potassium)	SS	ORAL
2 GRAM, ORAL			Valsartan (Valsartan)	SS	ORAL
ORAL			Bisoprolol (Bisoprolol)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/30/04ISR Number: 4439490-7Report Type:Expedited (15-DaCompany Report #2004057129
 Age:28 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 1800 MG (600 Initial or Prolonged MG, 3 IN 1 D), ORAL	Essential Hypertension Gastric Disorder Generalised Oedema Grand Mal Convulsion	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
1 IN 1 D, ORAL	Hepatic Enzyme Increased Pulmonary Oedema Tachycardia Tachypnoea		Udramil (Trandolapril, Verapamil Hydrochloride)	SS		ORAL
10 MG (10 MG, 1 IN 1 D), ORAL			Torasemide (Torasemide)	SS		ORAL
			Clindamycin Hydrochloride (Clindamycin Hydrochloride) Irbesartan (Irbesartan) Moxonidine (Moxonidine) Insulin (Insulin) Insulin Injection, Isophane (Insulin Injection, Isophane)	SS C C C C		

Date:08/30/04ISR Number: 4439506-8Report Type:Expedited (15-DaCompany Report #2003002971
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Analgesic Effect	Foreign	Neurontin			

300 MG (100
MG, TID),
ORAL

Atopy
Circulatory Collapse
Condition Aggravated
Convulsion
Gingivitis
Pyrexia
Therapeutic Response
Unexpected

Consumer
Health
Professional

(Gabapentin)
PS
ORAL

Date:08/30/04ISR Number: 4439646-3Report Type:Expedited (15-DaCompany Report #USA-2004-0016457
Age:51 YR Gender:Male I/FU:I

Outcome
Hospitalization -
Initial or Prolonged

PT
Blood Glucose Increased
Blood Magnesium Decreased
Blood Sodium Decreased
Confusional State
Contusion
Convulsion
Eye Disorder
Face Injury
Fall
Mental Status Changes
Nausea
Pancytopenia

Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
20 MG, QL2H		Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		
5 MG, Q4H			Oxyir Capsules 5 Mg(Oxycodone Hydrochloride, Oxycodone Hydrochloride) Ir	SS		
INTRAVENOUS	1 MG,		Aranesp(Darbepoetin)	SS		
INTRAVENOUS			Kytril(Granisetron)	SS		
2 MG; 4 MG			Prograft(Tacrolimus)	SS		
40 MG, DAILY			Protonix(Pantoprazole)	SS		
81 MG, DAILY			Acetylsalicylic Acid(Acetylsalicylic Acid)	SS		
400 MG			Gabapentin(Gabapentin)	SS		
			Magnesium(Magnesium)	SS		
			Senokot(Senna Fruit)	SS		
			Donepezil(Donepezil)	SS		
			Inderal(Propranolol Hydrochloride)	SS		
			Dronabinol(Dronabinol)	SS		
			Megace(Megestrol Acetate)	SS		
			Insulin(Insulin)	SS		
			Aldactone(Spironolactone)	SS		
			Ambien(Zolpidem Tartrate)	SS		
			Chlorpromazine(Chlorpromazine)	SS		

promazine)	C
Glipizide(Glipizide)	C
Procardia	C
Pentamidine(Pentamid ine)	C
Peri-Colace (Sennoside A+B, Docusate Sodium) Tablet	C
Dulcolax	C
Dexamethasone(Dexame thasone)	C
Cis-Platinum(Cisplat in)	C
Zoloft(Sertraline Hydrochloride)	C
Leukine(Sargramostim)	C
Adriamycin(Doxorubic in)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4439193-9Report Type:Expedited (15-DaCompany Report #2004047157
 Age:14 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Medication Error Pharmaceutical Product	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (100 MG, 3 IN 1 D), ORAL		Complaint	Professional				

Date:08/31/04ISR Number: 4439198-8Report Type:Expedited (15-DaCompany Report #2004056277
 Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 100 MG, ORAL		Acute Generalised Exanthematous Pustulosis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
750 MG (250 MG, 3 IN 1 D)		Aphthous Stomatitis C-Reactive Protein Increased Clostridial Infection	Professional	Pyridoxine Hydrochloride (Pyridoxine Hydrochloride)	SS		
INTRAVENTOUS (CYCLIC), INTRAVENTOUS	110 MG	Enterobacter Infection Inflammation Neutrophil Count Increased Oedema Peripheral Pain In Extremity Pyrexia		Ondansetron Hydrochloride (Ondansetron Hydrochloride) Docetaxel (Docetaxel)	SS SS		
INTRAVENTOUS	70 MG	Rash Rash Erythematous Scar Skin Lesion Staphylococcal Infection		Doxorubicin Hydrochloride (Doxorubicin Hydrochloride)	SS		

(CYCLIC),

INTRAVENOUS

Date:08/31/04ISR Number: 4439230-1Report Type:Expedited (15-DaCompany Report #2004056058

Age:84 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Balance Disorder Disorientation	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
600 MG (600 MG, 1 IN 1 D), ORAL		Fall Gait Disturbance Hearing Impaired Vision Blurred		Vicodin (Hydrocodone Bitartrate, Paracetamol) Paracetamol (Paracetamol) All Other Therapeutic Products (All Other Therapeutic Products)	C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/31/04ISR Number: 4439240-4Report Type:Expedited (15-DaCompany Report #2004057138

Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Confusional State	Consumer	Neurontin			
Initial or Prolonged	Contusion	Health	(Gabapentin)	PS		
Other	Convulsion	Professional	Aricept (Donepezil			
	Eye Injury		Hydrochloride)	SS		
	Fall		Oxycodone			
	Mental Status Changes		Hydrochloride			
	Nausea		(Oxycodone			
	Pancytopenia		Hydrochloride)	SS		
40 MG (20 MG,						
2 IN 1 D)	Soft Tissue Disorder					
	Swelling		Darbepoetin Alfa			
	Vomiting		(Darbepoetin Alfa)	SS		
SUBCUTANEOUS	(200 MCG),					
SUBCUTANEOUS						
			Dronabinol			
			(Dronabinol)	SS		
			Granisetron			
			(Granisetron)	SS		
1 MG (1 MG, 1						
IN 1 D)						
			Zolpidem Tartrate			
			(Zolpidem Tartrate)	C		
			Sertraline			
			Hydrochloride	C		
			Glipizide			
			(Glipizide)	C		
			Nifedipine			
			(Nifedipine)	C		
			Magnesium			
			(Magnesium)	C		
			Levofloxacin			
			(Levofloxacin)	C		
			Pentamidine			
			(Pentamidine)	C		
			Bisacodyl			
			(Bisacodyl)	C		
			Acetylsalicylic Aicd			
			(Acetylsalicylic			
			Acid)	C		

Furosemide	C
(Furosemide)	
Sargramostim	
(Sargramostim)	C
Dexamethasone	
(Dexamethasone)	C
Granisetron	
(Granisetron)	C
Morphine Sulfate	
(Morphine Sulfate)	C
Chlorpromazine	
(Chlorpromazine)	C
Tacrolimus	
(Tacrolimus)	C
Pantoprazole	
(Pantoprazole)	C
Doxorubicin	
(Doxorubicin)	C
Cisplatin	
(Cisplatin)	C
Deflazacort	
(Deflazacort)	C

Freedom Of Information (FOI) Report

Senna Fruit (Senna Fruit)	C
Propranolol Hydrochloride (Propranolol Hydrochloride)	C
Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotinamide, Insulin (Insulin) Spironolactone (Spironolactone)	C
Megestrol Acetate (Megestrol Acetate)	C
Repaglinide (Repaglinide)	C

Date:08/31/04ISR Number: 4439767-5Report Type:Expedited (15-DaCompany Report #2004058545
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Withdrawal Syndrome Dyskinesia	Foreign Health Professional	Neurontin (Gabapentin)	PS		
900 MG (300 MG, 3 IN 1 D)							

Date:08/31/04ISR Number: 4440209-4Report Type:Expedited (15-DaCompany Report #2004058317
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Inappropriate Antidiuretic Hormone Secretion	Foreign Health Professional	Neurontin (Gabapentin) All Other Therapeutic Products (All Other Therapeutic Products)	PS C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 300 MG PO Q Initial or Prolonged 4-5 HRS	Coma		Neurontin	PS		ORAL

[PRIOR TO
ADMISSION]

Codeine	C
Valium	C
Xanax	C
Glyburide	C
Metformin	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/01/04ISR Number: 4439730-4Report Type:Direct
 Age:32 YR Gender:Male I/FU:I

Company Report #CTU 226064

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Anxiety		Neurontin	PS		ORAL
2 TIMES DAILY						
Hospitalization -	Confusional State					
ORAL						
Initial or Prolonged	Suicidal Ideation					

Date:09/01/04ISR Number: 4441814-1Report Type:Expedited (15-DaCompany Report #2004056323
 Age:53 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Agranulocytosis	Foreign	Neurontin			
Initial or Prolonged	Iatrogenic Injury	Health	(Gabapentin)	PS		ORAL
UNK (3 IN 1						
D), ORAL	White Blood Cell Count	Professional				
	Decreased		Meprobamate			
UNK (1 IN 1			(Meprobamate)	SS		ORAL
D), ORAL						
			Aluminium Phosphate			
UNK (3 IN 1			Gel (Aluminium			
D), ORAL			Phosphate Gel)	SS		ORAL
UNK (1 IN 1			Mianserin			
D), ORAL			Hydrochloride			
			(Mianserin			
UNK (1 IN 1			Hydrochloride)	SS		ORAL
D), ORAL						
			Omeprazole			
UNK (1 IN 1			(Omeprazole)	SS		ORAL
D), ORAL						

UNKN (1 IN 1

D), ORAL

Date:09/01/04ISR Number: 4442015-3Report Type:Expedited (15-DaCompany Report #2004058552

Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Choking	Consumer	Neurontin			
300 MG (300		Diarrhoea		(Gabapentin)	PS		ORAL
MG, 1 IN 1		Dysgeusia					
D), ORAL		Dyspnoea					
		Feeling Abnormal		Prednisone	C		
		Gait Disturbance		Escitalopram	C		
		Nausea		Ultracet			
		Stomach Discomfort		(Paracetamol,			
				Tramadol			
				Hydrochloride)	C		
				Alprazolam	C		
				Rofecoxib	C		

Date:09/01/04ISR Number: 4443168-3Report Type:Expedited (15-DaCompany Report #2004059024

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis	Consumer	Neurontin			
600 MG		Knee Arthroplasty		(Gabapentin)	PS		
		Osteoporosis		Fentanyl	C		
				Ibuprofen	C		
				Rofecoxib	C		

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Freedom Of Information (FOI) Report

Helianthus Tuberosus C
 All Other
 Therapeutic Products C

Date:09/01/04ISR Number: 4443172-5Report Type:Expedited (15-DaCompany Report #2004052685
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Agitation Condition Aggravated Convulsion	Consumer	Neurontin (Tablets) (Gabapentin) Clonazepam (Clonazepam) Diazepam (Diazepam)	PS SS SS		

Date:09/01/04ISR Number: 4443176-2Report Type:Expedited (15-DaCompany Report #2004058193
 Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 600 MG (300 Initial or Prolonged MG, 2 IN 1 Other D), ORAL		Agitation Anger Crying Drug Ineffective Intentional Self-Injury Memory Impairment Skin Laceration Suicide Attempt	Consumer	Neurontin (Gabapentin) Bupropion Hydrochloride Chlordiazepoxide Tizanidine Hydrochloride Vicodin (Hydrocodone Bitartrate, Paracetamol) Depo-Estradiol (Chlorobutanol, Cottonseed Oil, Estradiol Cipionate) Vitamins Vicodin (Hydrocodone Bitartrate, Paracetamol)	PS C C C C C C C		ORAL

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Pain Tachycardia	Consumer	Dilantin Suspension (Phenytoin Sodium) Neurontin (Tablets) (Gabapentin)	PS SS		ORAL
600 MG (100							
MG , 1 IN 1							
D) ORAL							
				Irbesartan	C		
				Insulin	C		
				Levothyroxine Sodium	C		
				Lansoprazole	C		
				Rosuvastatin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/02/04ISR Number: 4444227-1Report Type:Expedited (15-DaCompany Report #A03200402800

Age:51 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Asthenia	Health	Ambien - Zolpidem			
Initial or Prolonged	Confusional State	Professional	Tartrate - Tablet	PS		ORAL
ORAL						
	Contusion		Aranesp -			
	Convulsion		Darbepoetin Alfa -			
	Face Injury		Solution	SS		
SUBCUTANEOUS	SUBCUTANEOUS					
	Fall		Marinol - Dronabinol	SS		
	Fatigue		Kytril - Granisetron			
	Gastrointestinal		- Solution -1 Mg	SS		
INTRAVENOUS	1 MG QD -					
	Haemorrhage					
INTRAVENOUS						
	Haemolytic Anaemia					
NOS						
	Liver Transplant		Oxycontin -			
	Rejection		Oxycodone			
	Metabolic Encephalopathy		Hydrochloride - 20			
	Pancytopenia		Mg	SS		
20 MG Q12HR						
	Swelling		Neurontin -			
			Gabapentin	SS		
			Aricept - Donepezil			
			Hydrochloride	SS		
			Morphine Sulfate	C		
			Chlorpromazine	C		
			Oxycodone	C		
			Tacrolimus	C		
			Pantoprazole	C		
			Deflazacort	C		
			Sertraline			
			Hydrochloride	C		
			Glipizide	C		
			Nifedipine	C		
			Magnesium	C		
			Levofloxacin	C		
			Pentamidine	C		
			Peri-Colace	C		
			Bisacodyl	C		
			Acetylsalicylic Acid	C		
			Furosemide	C		
			Sargramostim	C		

Dexamethasone	C
Doxorubicin	C
Cisplatin	C
Senna Fruit	C
Propranolol	
Hydrochloride	C
Multivitamins	C
Insulin	C
Spironolactone	C
Megestrol Acetate	C

Date:09/02/04ISR Number: 4444582-2Report Type:Expedited (15-DaCompany Report #2004051692

Age: Gender:Female I/FU:F

Outcome	PT
Other	Anxiety
	Asthenia
	Depression
	Feeling Abnormal
	Joint Injury
	Self-Medication

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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 Laceration Social Avoidant Behaviour Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		ORAL
800 MG (400 MG, 2 IN 1 DAY), ORAL		Therapy Non-Responder Treatment Noncompliance Weight Decreased					
				Carbamazepine (Carbamazepine)	SS		ORAL
				Oxcarbazepine (Oxcarbazepine)	SS		
				Levetiracetam (Levetiracetam)	SS		

Date:09/02/04ISR Number: 4444584-6Report Type:Expedited (15-DaCompany Report #2004058557
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 400 MG (400 MG), ORAL		Multiple Sclerosis Teeth Brittle Therapeutic Response Decreased Tooth Disorder Tooth Extraction Tooth Injury Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Certizine Hydrochloride (Cetirizine Hydrochloride)	C		
				Ranitidine Hydrochloride (Ranitidine Hydrochloride)	C		
				Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride)	C		
				Morphine Sulfate (Morphine Sulfate)	C		
				Interferon Beta (Interferon Beta)	C		

Loperamide
Hydrochloride
(Loperamide
Hydrochloride) C
Oxybutynin
(Oxybutynin) C
Tizanidine
Hydrochloride
(Tizanidine
Hydrochloride) C

Date:09/02/04ISR Number: 4444634-7Report Type:Expedited (15-DaCompany Report #2004058684
Age:54 YR Gender:Female I/FU:I

Outcome PT
Other Arrhythmia
Cerebrovascular Accident
Condition Aggravated
Crying
Depressed Mood
Depression
Diarrhoea

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Disorientation Dizziness Fall	Report Source	Product	Role	Manufacturer	Route
150 MG (1 IN 1 D), ORAL (SEE IMAGE)		Feeling Abnormal Feeling Drunk Homicidal Ideation	Consumer	Zoloft (Sertraline)	PS		ORAL
UNK (1 IN 1 D); ORAL (SEE IMAGE)		Hypertension Hyperventilation Kidney Infection Multiple Sclerosis		Neurontin (Gabapentin)	SS		ORAL
		Nausea Neuropathic Pain Neuropathy Pain Stupor Suicidal Ideation Thinking Abnormal Urine Abnormality Vomiting Projectile		Potassium Chloride Losartan Potassium Lansoprazole Metoclopramide Salbutamol	C C C C C		

Date:09/03/04ISR Number: 4440511-6Report Type:Expedited (15-DaCompany Report #PHBS2004US11398
Age:16 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Depressed Level Of Consciousness	Report Source Health Professional	Product Baclofen	Role PS	Manufacturer Novartis Sector: Pharma	Route
UNKNOWN		Dizziness Tachycardia		Tizanidine Hydrochloride	SS		
UNKNOWN				Nortriptyline Hydrochloride	SS		
UNKNOWN				Gabapentin	SS		
UNKNOWN				Phenelzine	SS		

UNKNOWN

Ketorolac SS

UNKNOWN

Bethanechol SS

UNKNOWN

Olanzapine SS

Date:09/03/04ISR Number: 4443031-8Report Type:Periodic

Company Report #2004UW11098

Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Cholesterol Increased Drug Interaction	Health Professional	Crestor Neurontin	PS SS		

Date:09/03/04ISR Number: 4444773-0Report Type:Expedited (15-DaCompany Report #2004043254

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Dupuytren'S Contracture Tenosynovitis	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (300 Other MG, 1 IN 1							

D), ORAL

Enalapril Maleate (Enalapril Maleate)	C
Furosemide (Furosemide)	C
Insulin (Insulin)	C
Alendronate Sodium	

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Freedom Of Information (FOI) Report

(Alendronate Sodium) C

Date:09/03/04ISR Number: 4446010-XReport Type:Expedited (15-DaCompany Report #2004040068

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Consumer	Neurontin			
		Cardiac Disorder		(Gabapentin)	PS		
2300 MG (1 IN		Chest Pain					
1 D)		Circulatory Collapse					
		Convulsion					
		Delusion					
		Dizziness					
		Electrocardiogram					
		Abnormal					
		Emotional Distress					
		Headache					
		Loss Of Consciousness					
		Pain					
		Palpitations					
		Psychosomatic Disease					

Date:09/03/04ISR Number: 4446098-6Report Type:Expedited (15-DaCompany Report #2004053609

Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Activities Of Daily	Consumer	Neurontin			
Other		Living Impaired	Health	(Gabapentin)	PS		ORAL
600 MG (300		Burns Second Degree	Professional				
MG, 1 IN 1		Insomnia					
D), ORAL		Movement Disorder		Lansoprazole			
		Pain		(Lansoprazole)	C		
		Photosensitivity Reaction		Budesonide			
		Sunburn		(Budesonide)	C		
				Loratadine			
				(Loratadine)	C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Bone Pain	Consumer	Neurontin			
Other		Colon Cancer		(Gabapentin)	PS		
		Drug Ineffective		Vicodin (Hydrocodone			
		Pain		Bitartrate,			
				Paracetamol)	SS		
				Methocarbamol			
				(Methocarbamol)	SS		
				Ibuprofen			
				(Ibuprofen)	SS		
				Paracetamol			
				(Paracetamol)	SS		

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Freedom Of Information (FOI) Report

Date:09/07/04ISR Number: 4445048-6Report Type:Expedited (15-DaCompany Report #2004059951
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arrhythmia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
200 MG (100 MG, 2 IN 1 D),		Chest Pain	Professional				

Date:09/07/04ISR Number: 4445053-XReport Type:Expedited (15-DaCompany Report #2004059979
Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Exanthem Haematoma	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
		Pruritus Generalised		Candesartan Cilexetil (Candesartan Cilexetil)	C		
				Paracetamol (Paracetamol)	C		

Date:09/07/04ISR Number: 4445061-9Report Type:Expedited (15-DaCompany Report #2004057238
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Drug Effect Decreased Drug Interaction	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
12 MG (12 MG, 1 IN 1 D),		International Normalised Ratio Decreased	Professional	Warfarin	SS		ORAL
ORAL		Pulmonary Embolism		Nicotine	C		

Date:09/07/04ISR Number: 4446007-XReport Type:Expedited (15-DaCompany Report #2004047157
 Age:14 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Effect Decreased Emotional Disorder	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (100 MG, 3 IN 1 D), ORAL		Growth Accelerated Medication Error Pharmaceutical Product Complaint	Professional				

Date:09/07/04ISR Number: 4446230-4Report Type:Expedited (15-DaCompany Report #2004002059
 Age:60 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Other	Abasia Accidental Overdose Coma Convulsion Drug Effect Decreased Dyskinesia Electroencephalogram Abnormal Facial Bones Fracture

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Event	Report Source	Product	Role	Manufacturer	Route
		Fall Fatigue Head Injury					
300 MG		Headache Memory Impairment	Consumer Health	Doxepin (Caps) (Doxepin)	PS		ORAL
(DAILY), ORAL		Migraine	Professional				
900 MG		Myoclonus Overdose		Neurontin (Gabapentin)	SS		ORAL
(DAILY), ORAL		Pain					
30 MG (BID),		Scar Somnolence		Phenobarbital (Phenobarbital)	SS		ORAL
ORAL		Surgery					
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Omeprazole (Omeprazole)	C		
				Verapamil Hydrochloride (Verapamil Hydrochloride)	C		
				Temazepam (Temazepam)	C		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		
				Baclofen (Baclofen)	C		
				Carisoprodol (Carisoprodol)	C		
				Parafon Forte (Chlorzoxazone, Paracetamol)	C		
				Hydroxyzine Embonate (Hydroxyzine Embonate)	C		
				Prochlorperazine Edisylate (Prochlorperazine			

Edisylate)	C
Sumatriptan	
Succinate	
(Sumatriptan	
Succinate)	C
Dyazide	
(Hydrochlorothiazide	
, Triamterene)	C
Pravastatin Sodium	
(Pravastatin Sodium)	C
Tegaserod	
(Tegaserod)	C
Percodan	
(Acetylsalicylic	
Acid, Caffeine,	
Phenacetin,	
Oxycodone	C
Vicodin	
(Paracetamol,	
Hydrocodone	
Bitartrate)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/07/04ISR Number: 4446332-2Report Type:Expedited (15-DaCompany Report #2004061469

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Murmur Depression Hepatitis C Memory Impairment Staphylococcal Infection Stress Suicide Attempt	Consumer	Neurontin (Gabapentin) Lithium (Lithium)	PS C		

Date:09/07/04ISR Number: 4446414-5Report Type:Expedited (15-DaCompany Report #2004059988

Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Porphyria	Health Professional	Neurontin (Gabapentin)	PS		

Date:09/07/04ISR Number: 4446430-3Report Type:Expedited (15-DaCompany Report #2004054705

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bone Density Decreased Diabetes Mellitus Fibromyalgia Osteoarthritis Recurrent Cancer Somnolence	Consumer	Neurontin (Gabapentin) Valdecoxib (Valdecoxib) Vicoprofen (Hydrocodone Bitartrate, Ibuprofen) All Other Therapeutic Products (All Other Therapeutic	PS SS SS		ORAL
900 MG (300							
MG , 3 NI 1							
D), ORAL							

Products)

SS

Date:09/07/04ISR Number: 4446432-7Report Type:Expedited (15-DaCompany Report #2004060294

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Neuropathy Peripheral	Consumer	Neurontin			
Other		Peripheral Occlusive Disease		(Gabapentin)	PS		
				Ultracet			
				(Paracetamol, Tramadol Hydrochloride)	SS		
				Carisoprodol			
				(Carisoprodol)	SS		
				Sodium Fluoride			
				(Sodium Fluoride)	SS		
				All Other Therapeutics			
				Products (All Other Therapeutic			
				Products)	SS		
				Propacet			

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Freedom Of Information (FOI) Report

(Dextropropoxyphene
Napsilate,
Paracetamol) SS
Morphine C

Date:09/07/04ISR Number: 4446434-0Report Type:Expedited (15-DaCompany Report #2004060932
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Appendicectomy	Consumer	Neurontin			
Other		Bladder Operation		(Gabapentin)	PS		ORAL
ORAL		Hysterectomy		Celebrex (Celecoxib)	SS		ORAL
200 MG (200		Muscle Twitching					
MG, 1 IN 1		Post Procedural					
D), ORAL		Complication		Aciclovir	C		
		Post Procedural Pain		Pantoprazole	C		
		Treatment Noncompliance		Diazepam	C		
				Quetiapine Fumarate	C		
				Fluoxetine			
				Hydrochloride	C		

Date:09/07/04ISR Number: 4446439-XReport Type:Expedited (15-DaCompany Report #2004046062
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Amnesia	Consumer	Neurontin (Tablets)			
Initial or Prolonged		Anger	Health	(Gabapentin)	PS		ORAL
ORAL		Blood Glucose Decreased	Professional	Lorazepam			
Other		Blood Pressure Increased		(Lorazepam)	SS		
		Circulatory Collapse		Morphine Sulfate			
		Coma		(Morphine Sulfate)	C		
		Convulsion		Morphine (Morphine)	C		
		Diarrhoea		Temazepam			
		Drug Ineffective		(Temazepam)	C		
		Dysgeusia		Valproate Semisodium			
		Feeling Abnormal		(Valproate			
		Haemoglobin Decreased		Semisodium)	C		

Hallucination
Heart Rate Increased
Hyperhidrosis
Hypermetropia
Overdose
Panic Attack
Respiratory Rate
Decreased
Urinary Tract Infection
Visual Disturbance

Date:09/07/04ISR Number: 4446443-1Report Type:Expedited (15-DaCompany Report #2004054371

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Arthritis	Consumer	Neurontin			
Disability		Asthma		(Gabapentin)	PS		
Other		Neuropathy Peripheral Thyroid Disorder					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/07/04ISR Number: 4446487-XReport Type:Expedited (15-DaCompany Report #2004060467

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetes Mellitus Gastric Disorder Neuropathic Arthropathy Osteoporosis	Consumer	Neurontin (Gabaentin) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Methadone (Methadone) Acetylsalicylic Acid (Acetylsalicylic Acid) Ibuprofen (Ibuprofen) Etodolac (Etodolac) Vicodin (Hydrochodone Bitartrate, Paracetamol)	PS SS SS SS SS SS C		

Date:09/08/04ISR Number: 4446583-7Report Type:Expedited (15-DaCompany Report #2004060018

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Potassium Decreased Blood Prolactin Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Disease Recurrence Malaise Nausea Tic Vomiting		Estrogens Conjugated Fentanyl Torasemide All Other Therapeutic Products	C C C C		

Date:09/08/04ISR Number: 4446584-9Report Type:Expedited (15-DaCompany Report #2004060005

Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Aggression	Consumer	Neurontin	
1800 MG (600	Balance Disorder		(Gabapentin)	PS
MG, 3 IN 1 D)	Coordination Abnormal			
	Delirium		Midodrine	
	Dizziness		Hydrochloride	C
	Dry Mouth		Vitamins	C
	Fatigue			
	Mood Altered			
	Somnolence			

Date:09/08/04ISR Number: 4446589-8Report Type:Expedited (15-DaCompany Report #2004057028
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Interaction	Health Professional	Neurontin (Gabapentin)	PS		
				Tricyclic Antidepressants	SS		
				All Other Therapeutic Products	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/08/04ISR Number: 4446662-4Report Type:Expedited (15-DaCompany Report #2004047168

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG, 3 IN 1 D), ORAL		Anaemia Blood Iron Decreased Platelet Count Decreased White Blood Cell Count Decreased	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Naproxen (Naproxen) Fluoxetine Hydrochloride (Fluoxdetine Hydrochloride) Alendronate Sodium (Alendronate Sodium) Calcium (Calcium) Librax (Chlordiazepoxide Hydrochloride, Clidinium Bromide)	C C C C		

Date:09/09/04ISR Number: 4448333-7Report Type:Expedited (15-DaCompany Report #2004055469

Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - ORAL Initial or Prolonged 3 GRAM (1 IN 1 D), ORAL (1 D), ORAL		Cardiac Arrest Grand Mal Convulsion Hepatic Failure Ketoacidosis Lactic Acidosis Respiratory Distress Septic Shock	Foreign Health Professional	Neurontin (Gabapentin) Metformin Hydrochloride (Metformin Hydrochloride) Rifampicin (Rifampicin) Nitrofurantoin	PS SS SS		ORAL ORAL ORAL

(1 D), ORAL	(Nitrofurantoin)	SS	ORAL
(1 D), ORAL	Ofloxacin (Ofloxacin)	SS	ORAL
(1 D), ORAL	Rilmenidine (Rilmenidine)	C	ORAL
	Irbesartan (Irbesartan)	C	
	Ferrous Sulfate (Ferrous Sulfate)	C	
	Ascorbic Acid (Ascorbic Acid)	C	

Date:09/09/04ISR Number: 4448563-4Report Type:Expedited (15-DaCompany Report #8739
Age:42 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Areflexia
Initial or Prolonged	Back Pain
Other	Cerebellar Syndrome
	Intervertebral Disc Disorder
	Nervous System Disorder
	Neuropathic Pain
	Nystagmus
	Pain In Extremity

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Freedom Of Information (FOI) Report

Dose	Duration	Paraesthesia Quadriparesis Sensory Disturbance	Report Source	Product	Role	Manufacturer	Route
100 MG TID		Stem Cell Transplant Vestibular Disorder	Foreign Other	Methotrexate	PS		
				Methotrexate	SS		
				Methotrexate	SS		
				Methotrexate	SS		
				Methotrexate	SS		
				Methotrexate	SS		
				Cytarabine	SS		
				Etoposide	SS		
				Neurontin	SS		
				Cyclophosphamide	C		
				Doxorubicin	C		
				Vincristine	C		
				Prednisone	C		
				Rituximab	C		
Carmustine	C						

Date:09/09/04ISR Number: 4448669-XReport Type:Expedited (15-DaCompany Report #2004058288
 Age:70 YR Gender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNK (3 IN 1 Other D), ORAL		Cholelithiasis Cholestasis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
				Aporex (Dextropropoxyphene Hydrochloride, Paracetamol)	SS		ORAL
UNK (3 IN 1 D), ORAL		Cytolytic Hepatitis	Professional	Clavulin (Amoxicillin Trihydrate, Clavulanate Potassium)	SS		ORAL

2 GRAM (2 IN

1 D), ORAL

Valsartan
(Valsartan) SS ORAL

UNK, ORAL

Bisoprolol
(Bisoprolol) C

Date:09/09/04ISR Number: 4448685-8Report Type:Expedited (15-DaCompany Report #2004061409
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Toxicity	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
1800 MG (600							
MG, 3 IN 1							

D), ORAL

Rofecoxib
(Rofecoxib) C

Diclofenac Sodium
(Diclofenac Sodium) C

Domperidone
(Domperidone) C

Warfarin (Warfarin) C

Cetirizine
(Cetirizine) C

Freedom Of Information (FOI) Report

Lansoprazole
 (Lansoprazole) C
 Paracetamol
 (Paracetamol) C
 Morphine Sulfate
 (Morphine Sulfate) C
 Fentanyl (Fentanyl) C
 Amitriptyline
 (Amitriptyline) C
 Cisplatin
 (Cisplatin) C

Date:09/09/04ISR Number: 4448808-0Report Type:Expedited (15-DaCompany Report #2004061141
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Failure Cough	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (1 IN 1 D), ORAL		Dyspnoea					
		Gastric Operation		Co-Diovan (Hydrochlorothiazide , Valsartan)	C		
				Rofecoxib	C		
				Premarin (Estrogens Conjugated)	C		

Date:09/09/04ISR Number: 4448858-4Report Type:Expedited (15-DaCompany Report #2004061114
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 20 MG (20 MG, Other 1 IN 1 D), ORAL		Cataract Coronary Artery Occlusion Disease Progression Drug Hypersensitivity Feeling Hot	Consumer	Lipitor (Atorvastatin)	PS		ORAL
				Neurontin			

300 MG (300	Glaucoma	(Gabapentin)	SS	ORAL
MG, 1 IN 1	Gout			
D), ORAL	Localised Infection			
	Movement Disorder	Vancomycin		
	Myalgia	(Vancomycin)	SS	
INTRAVENOUS	INTRAVENOUS			
	Pain	Cefazolin Sodium		
	Pain In Extremity	(Cafazolin Sodium)	SS	
INTRAVENOUS	INTRAVENOUS			
	Paraesthesia	Metoprolol Tartrate		
	Postoperative Infection	(Metoprolol		
	Pruritus	Tartrate)	C	
	Rash	Metformin		
		Hydrochloride		
		(Metformin		
		Hydrochloride)	C	
		Novolin 20/80		
		(Insulin Human,		
		Insulin Isophane,		
		Human Biosynthetic)	C	
		Furosemide		
		(Furosemide)	C	
		Potassium Chloride		
		(Potassium Chloride)	C	
		Nicotinic Acid		
		(Nicotinic Acid)	C	

Freedom Of Information (FOI) Report

Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C
 Lisinopril
 (Lisinopril) C

Date:09/09/04ISR Number: 4448941-3Report Type:Expedited (15-DaCompany Report #2004061152
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2700 MG (300 Other MG, 9 IN 1 D), ORAL		Back Pain Balance Disorder Breast Cancer Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL
40 MG (20 MG, 2 IN 1 D), ORAL		Lung Neoplasm Malignant Suicidal Ideation		Bextra (Valdecoxib)	SS		ORAL
400 MG (200 MG, 2 IN 1 D), ORAL				Celebrex (Celecoxib)	SS		ORAL
				All Other Therapeutic Products (All Other Therapeutic Products)	SS		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Losartan Potassium (Losartan Potassium)	C		
				Femara (Letrozole)	C		
				Lortab (Hydrocodone Bitartrate, Paracetamol)	C		
				Doxazosin Mesilate			

(Doxazosin Mesilate)	C
Valsartan	
(Valsartan)	C
Glipizide	
(Glipizide)	C
Metformin	
Hydrochloride	
(Metformin	
Hydrochloride)	C
Pioglitazone	
(Pioglitazone)	C
Cyclobenzaprine	
Hydrochloride	
(Cyclobenzaprine	
Hydrochloride)	C
Diazepam (Diazepam)	C
Multivitamins	
(Ascorbic Acid,	
Ergocalciferol,	
Folic Acid,	
Nicotinamide,	C
Tocopherol	
(Tocopherol)	C
Ocuvite (Ascorbic	
Acid, Retinol,	

D), ORAL

Cardiac Failure
Congestive
Cerebral Atrophy
Embolic Cerebral
Infarction
Somnolence

Company

Representative

Antibiotics
(Antibiotics)

C

Date:09/10/04ISR Number: 4459575-9Report Type:Periodic
Age:50 YR Gender:Female I/FU:I

Company Report #2004215321US

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Hypersensitivity Rash Generalised	Consumer	Bextra (Valdecoxib) Tablet	PS		
100 MG, QD				Neurontin (Gabapentin)	SS		
				Depakote (Valproate Semisodium)	C		
				Klonopin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/13/04ISR Number: 4447776-5Report Type:Expedited (15-DaCompany Report #200412821FR
 Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Rifadine	PS	Aventis	
Hospitalization - Initial or Prolonged		Cardiac Failure				Pharmaceuticals Inc.	ORAL
		Diabetic Complication		Oflocet	SS		ORAL
		Diabetic Foot		Glucophage	SS		ORAL
		Epilepsy		Furadantine	SS		ORAL
		Hepatic Failure		Hyperium	SS		ORAL
		Hypertension		Neurontin	SS		ORAL
		Ketoacidosis		Aprovel	C		ORAL
		Laboratory Test Abnormal		Fero-Grad	C		
		Lactic Acidosis					
		Respiratory Distress					
		Septic Shock					
		Urinary Tract Infection					

Date:09/13/04ISR Number: 4449263-7Report Type:Expedited (15-DaCompany Report #KII-2004-0013288
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Coma	Study Consumer Health Professional	Morphine Sulfate (Similar To Nda 19-516) (Morphine Sulfate)	PS		ORAL
ORAL			Other	Gabapentin (Gabapentin)	SS		ORAL
ORAL				Carisoprodol (Carisoprodol)	SS		ORAL
ORAL				Clonidine (Clonidine)	SS		ORAL

Date:09/13/04ISR Number: 4449379-5Report Type:Expedited (15-DaCompany Report #2004061433
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Osteoarthritis Osteoporosis Rheumatoid Arthritis	Consumer	Neurontin (Gabepentin)	PS		

Date:09/13/04ISR Number: 4449498-3Report Type:Direct Company Report #CTU 226922
 Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Agitation Anxiety Arthralgia Condition Aggravated Confusional State Depression Impulsive Behaviour Panic Attack Suicidal Ideation Suicide Attempt		Neurontin 600 Bextra Provil Lamictal Effector Wellbrutin	PS C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/13/04ISR Number: 4450743-9Report Type:Direct
Age:51 YR Gender:Female I/FU:I

Company Report #CTU 226946

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Anxiety		Lexapro 10 Mg	PS		
1 / DAY							
		Burning Sensation		Neurontin 300 Mg			
		Disorientation		Pfizer	SS	Pfizer	
2 / DAY AS							
NEEDED		Medication Error					
		Pain In Extremity					
		Pruritus					

Date:09/14/04ISR Number: 4450260-6Report Type:Expedited (15-DaCompany Report #2004061467
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Foreign	Gabapentin			
300 MG (300		Convulsion	Health	(Gabapentin)	PS		ORAL
MG, 1 IN 1							
		Tongue Haemorrhage	Professional				
D), ORAL							

Date:09/14/04ISR Number: 4450262-XReport Type:Expedited (15-DaCompany Report #2004062069
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase	Foreign	Neurontin			
1800 MG (600		Increased	Health	(Gabapentin)	PS		
MG, 3 IN 1 D)							
		Aspartate	Professional				
		Aminotransferase	Company				
		Increased	Representative				
		Gamma-Glutamyltransferase					
		Increased					
		Liver Disorder					

Date:09/14/04ISR Number: 4450377-6Report Type:Expedited (15-DaCompany Report #2004054607

Age:58 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Stevens-Johnson Syndrome	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other	300 MG ORAL	Toxic Epidermal Necrolysis	Professional	Atenolol (Atenolol)	C		
				Metformin (Metformin)	C		
				Potassium Chloride (Potassium Chloride)	C		
				Nifedipine (Nifedipine)	C		
				Initard (Insulin, Insulin Injection, Isophane)	C		
				Calcitriol (Calcitriol)	C		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		
				Rofecoxib (Rofecoxib)	C		

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Freedom Of Information (FOI) Report

Tocopherol	
(Tocopherol)	C
Simvastatin	
(Simvastatin)	C
Probenecid	
(Profenecid)	C
Loratadine	
(Loratadine)	C
Perindopril	
(Perindopril)	C
Famotidine	
(Famotidine)	C
Gliclazide	
(Glicazide)	C
Prochlorperazine	
(Prochlorperazine)	C
Paracetamol	
(Paracetamol)	C
Calcium (Calcium)	C
All Other	
Therapeutic Products	C

Date:09/14/04ISR Number: 4450512-XReport Type:Expedited (15-DaCompany Report #2004061440

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Back Pain	Consumer	Neurontin (Tablets)			
Initial or Prolonged	Bone Neoplasm		(Gabapentin)	PS		ORAL
1800 MG (600						
Other	Burning Sensation					
MG, 3 IN 1						
	Coma					
	Cyst		Tiagabine			
	Hallucination		Hydrochloride			
	Hepatic Function Abnormal		(Tiagabine			
	Metabolic Disorder		Hydrochloride)	C		
	Nerve Root Lesion		Cyclobenzaprine			
	Neuralgia		(Cyclobenzaprine)	C		
	Night Sweats		Estrogens Conjugated			
	Pain In Extremity		(Estrogens			
	Spinal Disorder		Conjugated)	C		
	Suicidal Ideation		Zolpidem Tartrate			
			(Zolpidem Tartrate)	C		

D), ORAL

Paroxetine	
Hydrochloride	
(Paroxetine	
Hydrochloride)	C
Tizanidine	
(Tizanidine)	C
All Other	
Therapeutic Products	
(All Other	
Therapeutics	
Products)	C
Seretide Mite	
(Fluticasone	
Propionate,	
Salmeterol	
Xinafoate)	C
Salbutamol	
(Salbutamol)	C
Epinephrine	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Epinephrine) C

Date:09/14/04ISR Number: 4450513-1Report Type:Expedited (15-DaCompany Report #2004061130
 Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG (600 MG, 3 IN 1 D), ORAL		Dysphagia Vomiting	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL

All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) SS
 Warfarin Sodium
 (Warfarin Sodium) C
 Megestrol Acetate
 (Megestrol Acetate) C
 Sucralfate
 (Sucralfate) C
 Esomeprazole
 (Esomeprazole) C
 Fentanyl (Fentanyl) C
 Metoprolol
 (Metoprolol) C
 Docusate Sodium
 (Docusate Sodium) C

Date:09/14/04ISR Number: 4450514-3Report Type:Expedited (15-DaCompany Report #2004051416
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Insomnia	Consumer Health Professional	Neurontin (Gabapentin) All Other Therapeutic Products (All Other Therapeutic	PS		

Products)	C
Alprazolam	
(Alprazolam)	C
Oxycodone	
Hydrochloride	
(Oxycodone	
Hydrochloride)	C
Pethidine	
Hydrochloride	
(Pethidine	
Hydrochloride)	C
Methadone	
(Methadone)	C
Esomeprazole	
(Esomeprazole)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/14/04ISR Number: 4450515-5Report Type:Expedited (15-DaCompany Report #2004061447

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Insomnia Jaundice	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Loss Of Control Of Legs Multiple Sclerosis Nausea Pain Pain In Extremity Skin Discolouration		Baclofen (Baclofen) Ibuprofen (Ibuprofen) Glatiramer Acetate (Glatiramer Acetate)	C C C		

Date:09/14/04ISR Number: 4450641-0Report Type:Expedited (15-DaCompany Report #2004048924

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 2700 MG (300 MG, UNKNOWN)		Abnormal Behaviour Blood Testosterone Decreased Legal Problem	Consumer	Neurontin (Gabapentin)	PS		

Date:09/14/04ISR Number: 4450658-6Report Type:Expedited (15-DaCompany Report #2004063259

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Chest Pain Oesophageal Ulcer	Consumer	Neurontin (Gabapentin) Motrin (Ibuprofen) Rofecoxib (Rofecoxib)	PS SS SS		

Date:09/14/04ISR Number: 4450660-4Report Type:Expedited (15-DaCompany Report #2004061446

Age:82 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1200 MG (400 MG,3 IN 1 D),	Drug Interaction Somnolence	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
ORAL				Oxycodone (Oxycodone)	SS		
				Digoxin (Digoxin)	C		
				Lisinopril (Lisinopril)	C		
				Hydrochlorothiazide (Hydrochlorothiazide)	C		
				Baclofen (Baclofen)	C		
				Warfarin Sodium (Warfarin Sodium)	C		
				Melatonin (Melatonin)	C		
				Paracetamol (Paracetamol)	C		
				Simvastatin (Simvastatin)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/14/04ISR Number: 4450661-6Report Type:Expedited (15-DaCompany Report #2004061448

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Nephrolithiasis	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:09/14/04ISR Number: 4450663-XReport Type:Expedited (15-DaCompany Report #2004051708

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (300MG,1 IN 1 D),ORAL		Back Pain Headache Muscle Spasms Neck Pain Nerve Compression Walking Aid User Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:09/15/04ISR Number: 4450267-9Report Type:Expedited (15-DaCompany Report #USA-2004-0016457

Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 20 MG, Q12H		Asthenia Blood Glucose Increased Blood Magnesium Decreased Blood Sodium Decreased	Health Professional Other	Oxycontin Tablets (Oxycondone Hydrochloride) Cr Tablet	PS		
5 MG, Q4H		Confusional State Contusion Convulsion Face Injury Fall Fatigue		Oxyir Capsules 5mg (Oxycodone Hydrochloride, (Oxycodone Hydrochloride) Ir	SS		
				Aranesp			

		Mental Status Changes	(Darbepoetin)	SS
		Nausea	Kytril (Granisetron)	SS
INTRAVENOUS	1 MG,			
		Pancytopenia		
INTRAVENOUS				
		Swelling	Prograf (Tacrolimus)	SS
SEE IMAGE				
		Vomiting	Protonil (Pantoprazole)	SS
40 MG DAILY				
			Acetylsalicylic Acid (Acetylsalicylic Acid)	SS
81 MG, DAILY				
			Gabapentin (Gabapentin)	SS
			Magnesium (Magnesium)	SS
400 MG				
			Senokot (Sennoside A+B)	SS
			Donepezil (Donepezil)	SS
			Inderal (Propranolol Hydrochloride)	SS
			Dronabinol (Dronabinol)	SS
			Megace (Megestrol Acetate)	SS
			Insulin (Insulin)	SS
			Aldactone	

Freedom Of Information (FOI) Report

(Spironolactone)	SS
Ambien (Zolpidem Tartrate)	SS
Chlorpromazine (Chlorpromazine)	C
Glipizide (Glipizide)	C
Procardia	C
Pentamidine (Pentamidine)	C
Peri-Colace (Sennoside A+B, Docusate Sodium)	C
Dulcolax	C
Dexamethasone (Dexamethasone)	C
Cis-Platinum (Cisplatin)	C
Zoloft (Sertraline Hydrochloride)	C
Leukine (Sargramostim)	C
Adriamycin (Doxorubicin)	C

Date:09/15/04ISR Number: 4451367-XReport Type:Expedited (15-DaCompany Report #2003113288
 Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Buttock Pain	Health	Neurontin			
Other		Dizziness	Professional	(Gabapentin)	PS		ORAL
300 MG		Ejaculation Disorder					
(DAILY), ORAL		Myocardial Infarction		Trimipramine			
		Somnolence		(Trimipramine)	C		
		Urinary Incontinence		Acetylsalicylic Acid			
		Visual Disturbance		(Acetylsalicylic Acid)	C		
				All Other Therapeutic Products			
				(All Other Therapeutic Products)	C		
				Furosemide			

(Furosemide)	C
Carvedilol	
(Carvedilol)	C
Captopril	
(Captopril)	C
Ibuprofen	
(Ibuprofen)	C
Paracetamol	
(Paracetamol)	C
Arthrotec	
(Diclofenac Sodium, Misoprostol)	C
Dextropropoxyphene Hydrochloride	
(Dextropropoxyphene Hydrochloride)	C
Theragran (Vitamins Nos)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Thiamine
Hydrochloride
(Thiamine
Hydrochloride) C

Date:09/15/04ISR Number: 4451427-3Report Type:Expedited (15-DaCompany Report #2004062380
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia Arthropathy Chondropathy Fall	Consumer	Neurontin (Gabapentin)	PS		

Date:09/15/04ISR Number: 4451445-5Report Type:Expedited (15-DaCompany Report #2004062834
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthropathy Back Disorder Musculoskeletal Disorder	Consumer	Neurontin (Gabapentin) Ibuprofen (Ibuprofen) Paracetamol (Paracetamol)	PS SS C		

Date:09/15/04ISR Number: 4451451-0Report Type:Expedited (15-DaCompany Report #2004062831
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia Back Pain Osteoarthritis Pain In Extremity	Consumer	Neurontin (Gabapentin) Oxycocet (Oxycodone Hydrochloride, Paracetamol) Paroxetine Hydrochloride (Paroxetine Hydrochloride)	PS SS SS		

Rofecoxib	
(Rofecoxib)	SS
Methylphenidate	
Hydrochloried	
(Methylphenidate	
Hydrochloride)	SS
Fentanyl (Fentanyl)	SS

Date:09/15/04ISR Number: 4451615-6Report Type:Expedited (15-DaCompany Report #2004062323

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Withdrawal Syndrome	Consumer	Neurontin			
		Economic Problem		(Gabapentin)	PS		ORAL
6400 MG, ORAL				Fentanyl (Fentanyl)	C		
				Benzocaine			
				(Benzocaine)	C		
				Clonidine			
				(Clonidine)	C		

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Freedom Of Information (FOI) Report

Clopidogrel Sulfate (Clopidogrel Sulfate)	C
Temazepam (Temazepam)	C
Trazodone (Trazodone)	C
Imipramine (Imipramine)	C
Levothyroxine Sodium (Levothyroxine Sodium)	C
Risperidone (Risperidone)	C

Date:09/15/04ISR Number: 4451622-3Report Type:Expedited (15-DaCompany Report #2004061480
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Disease Recurrence	Foreign	Neurontin			
Other		Fall	Health	(Gabapentin)	PS		
		Hip Fracture	Professional	Amlodipine Besilate			
		Vertigo	Company	(Amlodipine Besilate)	C		
			Representative	Paroxetine Hydrochloride			
				(Paroxetine Hydrochloride)	C		
				All Other Theapeutic Products (All Other Therapeutic Products)	C		

Date:09/15/04ISR Number: 4452003-9Report Type:Direct Company Report #CTU 227202
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - PO [PRIOR TO Initial or Prolonged ADMISSION]		Asthenia		Neurontin	PS		
		Chills					

Nausea

Zetia C
Hydrocodone C
Celebrex C
Mirapex C

Date:09/15/04ISR Number: 4452071-4Report Type:Direct
Age:53 YR Gender:Female I/FU:I

Company Report #CTU 227269

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 300 MG Hospitalization - Initial or Prolonged	Suicidal Ideation		Neurontin	PS	Parke-Davis	

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/16/04ISR Number: 4452502-XReport Type:Expedited (15-DaCompany Report #2004036067

Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, 3 IN 1 D), ORAL		Thrombocytopenia	Foreign Health Professional Company Representative	Gabapentin (Gabapentin) Torasemide (Torasemide) Spironolactone (Spironolactone) Acetylsalicylic Acid (Acetylsalicylic Acid) Omeprazole (Omeprazole) Alprazolam (Alprazolam)	PS C C C C C		ORAL

Date:09/16/04ISR Number: 4452504-3Report Type:Expedited (15-DaCompany Report #2004042400

Age:65 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200 MG (100 MG, 2 IN 1 D), ORAL		Breast Mass Nipple Disorder	Foreign Consumer Health Professional Company Representative	Neurontin (Gabapentin) Limaprost (Limaprost)	PS C		ORAL

Date:09/16/04ISR Number: 4452506-7Report Type:Expedited (15-DaCompany Report #2004062853

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Gait Disturbance Pancreatic Atrophy	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
200 MG (100 MG, 2 IN 1D), ORAL				Clomipramine (Clomipramine)	C		

Date:09/16/04ISR Number: 4452510-9Report Type:Expedited (15-DaCompany Report #2004062825
Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Cardiac Failure Drug Interaction Generalised Oedema	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Risperidone (Risperidone) All Other Therapetutic Products (All Other Therapeutic Products)	PS SS C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/16/04ISR Number: 4452849-7Report Type:Expedited (15-DaCompany Report #2004064412

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin	PS		
Other		Amnesia Cerebrovascular Accident Completed Suicide Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress Suicidal Ideation Suicide Attempt		(Gabapentin)			

Date:09/16/04ISR Number: 4452850-3Report Type:Expedited (15-DaCompany Report #2004064444

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin	PS		
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Suicidal Ideation Suicide Attempt		(Gabapentin)			

Date:09/16/04ISR Number: 4452851-5Report Type:Expedited (15-DaCompany Report #2004064438

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin	PS		
Other		Amnesia Cerebrovascular Accident Completed Suicide Hepatic Failure		(Gabapentin)			

Hypertension
Injury
Myocardial Infarction
Renal Failure
Stress
Suicidal Ideation
Suicide Attempt

Date:09/16/04ISR Number: 4452852-7Report Type:Expedited (15-DaCompany Report #2004064433

Age: Gender:Female I/FU:I

Outcome

PT

Other

Amnesia

Cerebrovascular Accident

Completed Suicide

Emotional Distress

Hepatic Failure

Hypertension

Injury

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Freedom Of Information (FOI) Report

Dose	Duration	Myocardial Infarction Renal Failure Suicidal Ideation	Report Source	Product	Role	Manufacturer	Route
		Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4452857-6Report Type:Expedited (15-DaCompany Report #2004063422
Age: Gender:Male I/FU:I

Outcome Dose Other	Duration	PT Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Suicidal Ideation Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4452858-8Report Type:Expedited (15-DaCompany Report #2004064409
Age: Gender:Male I/FU:I

Outcome Dose Other	Duration	PT Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Suicidal Ideation Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Neurontin			
Other		Cerebrovascular Accident		(Gabapentin)	PS		
		Completed Suicide					
		Emotional Distress					
		Hepatic Failure					
		Hypertension					
		Injury					
		Myocardial Infarction					
		Renal Failure					
		Suicidal Ideation					
		Suicide Attempt					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/16/04ISR Number: 4452860-6Report Type:Expedited (15-DaCompany Report #2004064395

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4452861-8Report Type:Expedited (15-DaCompany Report #2004064430

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4452862-XReport Type:Expedited (15-DaCompany Report #2004064427

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure	Consumer	Neurontin (Gabapentin)	PS		

Hypertension
Injury
Myocardial Infarction
Renal Failure
Suicidal Ideation
Suicide Attempt

Date:09/16/04ISR Number: 4452863-1Report Type:Expedited (15-DaCompany Report #2004064423

Age: Gender:Male I/FU:I

Outcome

PT

Other

Amnesia

Cerebrovascular Accident

Completed Suicide

Emotional Distress

Hepatic Failure

Hypertension

Injury

Myocardial Infarction

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Renal Failure Suicidal Ideation Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4452890-4Report Type:Expedited (15-DaCompany Report #2004063419
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress					

Date:09/16/04ISR Number: 4452891-6Report Type:Expedited (15-DaCompany Report #2004063317
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress					

Date:09/16/04ISR Number: 4452892-8Report Type:Expedited (15-DaCompany Report #2004063298
Age: Gender:Male I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Mental Disorder Myocardial Infarction Renal Failure Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:09/16/04ISR Number: 4452893-XReport Type:Expedited (15-DaCompany Report #2004063456

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4452894-1Report Type:Expedited (15-DaCompany Report #2004063455

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4452895-3Report Type:Expedited (15-DaCompany Report #2004063449

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension	Consumer	Neurontin (Gabapentin)	PS		

Injury
Myocardial Infarction
Renal Failure

Date:09/16/04ISR Number: 4452899-0Report Type:Expedited (15-DaCompany Report #2004052246

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Atrophy Body Height Decreased	Consumer Health	Dilantin Suspension (Phenytoin Sodium)	PS		ORAL
Other ORAL	Condition Aggravated Convulsion Rash Generalised Speech Disorder	Professional	Neurontin (Gabapentin)	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/16/04ISR Number: 4452900-4Report Type:Expedited (15-DaCompany Report #2004062829
 Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Arachnoiditis Back Pain	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1600 MG (800 MG, 2 IN 1 D), ORAL		Disease Recurrence Economic Problem Impaired Healing Migraine Stress Treatment Noncompliance Upper Limb Fracture		Tizanidine (Tizanidine) Carisoprodol (Carisoprodol) Temazepam (Temazepam) Metahdone (Methadone)	C C C C		

Date:09/16/04ISR Number: 4452903-XReport Type:Expedited (15-DaCompany Report #2004056093
 Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Disease Recurrence Hallucination, Auditory	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL		Treatment Noncompliance	Professional	Oxycodone Hydrochloride (Oxycodone Hydrochloride) Baclofen (Baclofen) Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Alopecia Chest Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 2 IN 1 D); ORAL (SEE IMAGE) (1 IN 1 D); ORAL (SEE IMAGE)		Drug Effect Decreased Drug Interaction Dyspnoea Headache Heart Rate Increased Neck Pain		Zoloft (Sertraline)	SS		ORAL
UNKNOWN	(SEE IMAGE)			Escitalopram (Escitalopram)	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Post Procedural Complication	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Transurethral Prostatectomy Urinary Incontinence	Professional	Celecoxib (Celecoxib) Furosemide	C		

Freedom Of Information (FOI) Report

(Furosemide) C
 Allopurinol
 (Allopurinol) C

Date:09/16/04ISR Number: 4452918-1Report Type:Expedited (15-DaCompany Report #2004054101
 Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG, (300 MG, 3 IN 1 D), ORAL		Blood Pressure Fluctuation Dehydration Depressed Level Of Consciousness Diplopia Disorientation Fatigue	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
75 MG (75 MG, 1 IN 1 D)		Heart Rate Increased Kidney Infection Pruritus Generalised Rash Somnolence		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride0 Hyzaar (Hydrochlorothiazide , Losartan Potassium) Hysocyamine (Hyoscyamine) Oxycodone Hydrochloride (Oxycodone Hydrochloride) Fentanyl (Fentanyl) Alprazolam (Alprazolam) Prednisone (Prednisone) Warfarin (Warfarin) Lansoprazole (Lansoprazole) Triamcinolone Acetonide (Triamcinolone Acetonide)	SS C C C C C C C C C C C		

Date:09/16/04ISR Number: 4453026-6Report Type:Expedited (15-DaCompany Report #2004063404
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Neurontin			
Other		Cerebrovascular Accident		(Gabapentin)	PS		
		Completed Suicide					
		Hepatic Failure					
		Hypertension					
		Injury					
		Myocardial Infarction					
		Renal Failure					
		Stress					
		Suicidal Ideation					
		Suicide Attempt					

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Freedom Of Information (FOI) Report

Date:09/16/04ISR Number: 4453027-8Report Type:Expedited (15-DaCompany Report #2004063401

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4453028-XReport Type:Expedited (15-DaCompany Report #2004063398

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4453029-1Report Type:Expedited (15-DaCompany Report #2004063396

Age: Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Hepatic Failure	Consumer	Neurontin (Gabapentin)	PS		

Hypertension
Injury
Mental Disorder
Myocardial Infarction
Renal Failure
Stress
Suicidal Ideation
Suicide Attempt

Date:09/16/04ISR Number: 4453030-8Report Type:Expedited (15-DaCompany Report #2004063452
Age: Gender:Female I/FU:I

Outcome PT
Other Amnesia
Cerebrovascular Accident
Completed Suicide
Hepatic Failure
Hypertension
Injury

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Dose	Duration	Myocardial Infarction Renal Failure Stress	Report Source	Product	Role	Manufacturer	Route
		Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4453031-XReport Type:Expedited (15-DaCompany Report #2004063426
Age: Gender:Male I/FU:I

Outcome Dose Other	Duration	PT Amnesia Cerebrovascular Accident Completed Suicide Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress Suicidal Ideation Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4453032-1Report Type:Expedited (15-DaCompany Report #2004063429
Age: Gender:Male I/FU:I

Outcome Dose Other	Duration	PT Amnesia Cerebrovascular Accident Completed Suicide Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress Suicidal Ideation Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Neurontin	PS		
Other		Cerebrovascular Accident		(Gabapentin)			
		Completed Suicide					
		Hepatic Failure					
		Hypertension					
		Injury					
		Myocardial Infarction					
		Renal Failure					
		Stress					
		Suicidal Ideation					
		Suicide Attempt					

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Freedom Of Information (FOI) Report

Date:09/16/04ISR Number: 4453039-4Report Type:Direct
 Age:17 YR Gender:Female I/FU:I

Company Report #CTU 227330

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Convulsion		Neurontin	PS		
Initial or Prolonged		Depression		Lexipro	SS		
Other		Drug Ineffective					
		Off Label Use					

Date:09/16/04ISR Number: 4453172-7Report Type:Expedited (15-DaCompany Report #2004063314
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Neurontin			
		Cerebrovascular Accident		(Gabapentin)	PS		
		Emotional Distress					
		Hepatic Failure					
		Hypertension					
		Injury					
		Myocardial Infarction					
		Suicidal Ideation					
		Suicide Attempt					

Date:09/16/04ISR Number: 4453174-0Report Type:Expedited (15-DaCompany Report #2004063301
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Neurontin			
		Cerebrovascular Accident		(Gabapentin)	PS		
		Emotional Disorder					
		Hepatic Failure					
		Hypertension					
		Injury					
		Mental Disorder					
		Myocardial Infarction					
		Renal Failure					
		Stress					
		Suicidal Ideation					
		Suicide Attempt					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Neurontin			
Other		Cerebrovascular Accident		(Gabapentin)	PS		
		Completed Suicide					
		Emotional Distress					
		Hepatic Failure					
		Hypertension					
		Injury					
		Myocardial Infarction					
		Renal Failure					
		Stress					
		Suicidal Ideation					
		Suicide Attempt					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/16/04ISR Number: 4453178-8Report Type:Expedited (15-DaCompany Report #2004064418

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4453179-XReport Type:Expedited (15-DaCompany Report #2004064406

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4453180-6Report Type:Expedited (15-DaCompany Report #2004063324

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident	Consumer	Neurontin (Gabapentin)	PS		

Completed Suicide
Emotional Distress
Hepatic Failure
Hypertension
Injury
Myocardial Infarction
Renal Failure
Stress
Suicidal Ideation
Suicide Attempt

Date:09/16/04ISR Number: 4453182-XReport Type:Expedited (15-DaCompany Report #2004063291
Age: Gender:Female I/FU:I

Outcome PT
Other Amnesia
Cerebrovascular Accident
Completed Suicide
Emotional Distress

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Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
		Hepatic Failure Hypertension Injury	Consumer	Neurontin (Gabapentin)	PS		
		Myocardial Infarction Renal Failure Stress Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4453185-5Report Type:Expedited (15-DaCompany Report #2004063284

Age: Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT	Consumer	Neurontin (Gabapentin)	PS	
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS	

Date:09/16/04ISR Number: 4453189-2Report Type:Expedited (15-DaCompany Report #2004063278

Age: Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT	Consumer	Neurontin (Gabapentin)	PS	
Other		Amnesia Cerebrovascular Accident Completed Suicide Emotional Distress Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress	Consumer	Neurontin (Gabapentin)	PS	

Suicidal Ideation
Suicide Attempt

Date:09/16/04ISR Number: 4453191-0Report Type:Expedited (15-DaCompany Report #2004063320
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Neurontin			
Other		Cerebrovascular Accident		(Gabapentin)	PS		
		Completed Suicide					
		Emotional Distress					
		Hepatic Failure					
		Hypertension					
		Injury					
		Myocardial Infarction					
		Renal Failure					
		Suicidal Ideation					
		Suicide Attempt					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/16/04ISR Number: 4453192-2Report Type:Expedited (15-DaCompany Report #2004063307

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4453193-4Report Type:Expedited (15-DaCompany Report #2004063300

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4453194-6Report Type:Expedited (15-DaCompany Report #2004063395

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Cerebrovascular Accident Completed Suicide Hepatic Failure	Consumer	Neurontin (Gabapentin)	PS		

Hypertension
Injury
Myocardial Infarction
Renal Failure
Stress
Suicidal Ideation
Suicide Attempt

Date:09/16/04ISR Number: 4453198-3Report Type:Expedited (15-DaCompany Report #2004063392
Age: Gender:Male I/FU:I

Outcome PT
Other Amnesia
Cerebrovascular Accident
Completed Suicide
Emotional Distress
Hepatic Failure
Hypertension
Injury

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Mental Disorder Myocardial Infarction Renal Failure	Report Source	Product	Role	Manufacturer	Route
		Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4453199-5Report Type:Expedited (15-DaCompany Report #2004063304
Age: Gender:Male I/FU:I

Outcome Dose Other	Duration	PT Amnesia Cerebrovascular Accident Completed Suicide Emotional Disorder Hepatic Failure Hypertension Injury Mental Disorder Myocardial Infarction Renal Failure Suicidal Ideation Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin)	PS		

Date:09/16/04ISR Number: 4453203-4Report Type:Expedited (15-DaCompany Report #2004063285
Age: Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Amnesia Cerebrovascular Accident Completed Suicide Hepatic Failure Hypertension Injury Myocardial Infarction Renal Failure Stress Suicidal Ideation Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin)	PS		

Outcome	PT
Hospitalization -	Asthenia
Initial or Prolonged	Convulsion
	Fall
	Fatigue
	Gastrointestinal
	Haemorrhage
	Haemolytic Anaemia
	Head Injury
	Liver Transplant
	Rejection
	Mental Status Changes
	Metabolic Encephalopathy
	Nausea
	Neuropathy
	Pancytopenia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Swelling Tremor Vomiting	Report Source	Product	Role	Manufacturer	Route
UNKNOWN		1	DAY	Granisetron Hydrochloride	PS	Roche	
SUBCUTANEOUS				Darbepoetin Alfa	SS		
UNKNOWN				Dronabinol	SS		
UNKNOWN		31	DAY	Oxycodone	SS		
UNKNOWN				Gabapentin	SS		
UNKNOWN				Donepezil Hydrochloride	SS		
UNKNOWN				Zolpidem Tartrate	SS		
55	DAY			Morphine Sulfate	C		
				Chlorpromazine	C		
				Oxyir	C		
				Tacrolimus	C		
				Pantoprazole	C		
				Prandin	C		
				Sertraline Hydrochloride	C		
				Glipizide	C		
				Nifedipine	C		
				Magnesium	C		
8	DAY			Levofloxacin	C		
				Pentamidine	C		
				Pericolace	C		
				Bisacodyl	C		
				Acetylsalicylic Acid	C		
INTRAVENOUS		1	DAY	Frusemide	C		
SUBCUTANEOUS		6	DAY	Sargamostim	C		
INTRAVENOUS		1	DAY	Dexamethasone	C		
				Doxorubicin	C		
				Cisplatin	C		
				Senna Fruit	C		

Propranolol C
 Multivitamins C
 Insulin C
 Spironolactone C
 Megestrol Acetate C

Date:09/17/04ISR Number: 4452830-8Report Type:Expedited (15-DaCompany Report #2004057138
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Asthenia Confusional State Contusion Convulsion Fall Fatigue Mental Status Changes Nausea Pancytopenia	Consumer Health Professional	Neurontin (Gabapentin) Aricept (Donepezil Hydrochloride) (Donepezil) Oxycodone Hydrochloride (Oxycodone Hydrochloride)	PS SS SS		
40 MG (20 MG , 2 IN 1 D)		Skull X-Ray Abnormal					
SUBCUTANEOUS	(200 MCG),	Swelling Vomiting		Darbepoetin Alfa (Darbepoetin Alfa)	SS		
SUBCUTANEOUS				Dronabinol (Dronabinol) Granisetron	SS		

Freedom Of Information (FOI) Report

1 MG (1 MG, 1

IN 1 D)

(Granisetron) SS

Zolpidem Tartrate (Zolpidem Tartrate)	C
Sertraline Hydrochloride(Sertra line Hydrochloride)	C
Glipizide (Glipizide)	C
Nifedipine (Nifedipine)	C
Magenesium (Magnesium)	C
Levofloxacin (Levofloxacin)	C
Pentamidine (Pentamidine)	C
Peri-Colace (Casanthranol, Docusate Sodium)	C
Bisacodyl (Bisacodyl)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C
Furosemide (Furosemide)	C
Sargramostim (Sargramostim)	C
Dexamethasone (Dexamethasone)	C
Granisetron (Granisetron)	C
Morphine Sulfate (Morphine Sulfate)	C
Chlorpromazine (Chlorpromazine)	C
Tacrolimus (Tacrolimus)	C
Pantoprazole (Pantoprazole)	C
Doxorubicin (Doxorubicin)	C
Cisplatin(Cisplatin)	C
Deflazacort (Deflazacort)	C

Senna Fruit (Sena Fruit)	C
Propranolol Hydrochloride (Propranolol Hydrochloride)	C
Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotinamide,	C
Insulin (Insulin)	C
Spirolactone (Spirolactone)	C
Megestrol Acetate (Megestrol Acetate)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Repaglinide
(Repaglinide) C

Date:09/17/04ISR Number: 4454523-XReport Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 227473

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Generic Gabapentin	PS		
300 MG 11			Drug Ineffective				
TABS/DAY			Pharmaceutical Product				
			Complaint				

Date:09/17/04ISR Number: 4455250-5Report Type:Expedited (15-DaCompany Report #2004063250
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability			Foreign Health	Neurontin (Gabapentin)	PS		
Other							
1200 MG (1 D)			Somnolence				
			Vertigo	Dextropropoxyphene (Dextropropoxyphene)	C		
			Professional Company Representative	Rofecoxib (Rofecoxib)	C		

Date:09/17/04ISR Number: 4455491-7Report Type:Expedited (15-DaCompany Report #2004063492
Age:83 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabaqentin)	PS		ORAL
Other							
300 MG (300			Asthenia				
MG, 1 IN 1			Difficulty In Walking				
			Fall				
D), ORAL			Hypoacusis	Clopidogrel Sulfate (Clopidogrel)			

Sulfate) C
 Simvastatin C
 (Simvastatin) C
 Ramipril (Ramipril) C
 Vitamins (Vitamins) C

Date:09/17/04ISR Number: 4455492-9Report Type:Expedited (15-DaCompany Report #2004063490
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1600 MG (400 Other MG, 4 IN 1 D), ORAL	Hot Flush Hyperhidrosis Marital Problem Mental Disorder Mid-Life Crisis Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Erythromycin (Erythromycin) Citalopram Hydrobromide (Citalopram Hydrobromide)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/17/04ISR Number: 4455512-1Report Type:Expedited (15-DaCompany Report #2004063309

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),	Asthenia Cardiac Operation	Consumer	Neurontin (Gabapentin)	PS		

All Other Therapeutic Products (All Other Therapeutic Products)	C		
Moexipril Hydrochloride (Moexipril Hydrochloride)	C		
Pravastatin Sodium (Pravastatin Sodium)	C		
Os-Cal (Calcium, Ergocalciferol)	C		
Centrum Silver (Ascorbic Acid, Calcium, Minerals Nos, Retinol, Tocopheryl Acetate, Tocopherol (Tocopherol)	C		
Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
Celecoxib (Celecoxib)	C		

Date:09/17/04ISR Number: 4455548-0Report Type:Expedited (15-DaCompany Report #2004064627

Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Dose 5 MG (5 MG, 1		Back Pain	Consumer	Norvasc (Amlodipine)	PS		ORAL

Initial or Prolonged IN 1 D), ORAL Other (3 IN 1 D), ORAL	Drug Effect Decreased Drug Ineffective Insomnia Pain In Extremity Pollakiuria Prostate Cancer	Neurontin (Gabapentin)	SS	ORAL
(1 D), ORAL		Tolterodine L-Tartrate (Tolterodine L-Tartrate)	SS	ORAL
		Tamsulosin (Tamsulosin) Terazosin Hydrochloride (Terazosin Hydrochloride)	SS C	

Date:09/20/04ISR Number: 4454665-9Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 227587

Outcome PT
Other Arthritis
Carpal Tunnel Syndrome

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Interaction	Report Source	Product	Role	Manufacturer	Route
75 MG 2 / DAY		Neuralgia Panic Reaction Suicidal Ideation Thinking Abnormal		Effexor Xr 75 Mg Wyeth	PS	Wyeth	ORAL
ORAL				Neurontin 100 Mg Pfizer	SS	Pfizer	ORAL
200 MG 3 / DAY	ORAL			Clonazepam	C		
				Omeprazole	C		
				Lipitor	C		
				Lisinopril	C		
				Zestril	C		

Date:09/20/04ISR Number: 4455873-3Report Type:Expedited (15-DaCompany Report #2004054371
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Disability Other		Arthritis Asthma Neuropathy Peripheral Thyroid Disorder	Consumer Health Professional	Neurontin (Gabapentin)	PS		

Date:09/20/04ISR Number: 4455878-2Report Type:Expedited (15-DaCompany Report #2004061130
Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dysphagia Pharmaceutical Product	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D),	ORAL	Complaint Vomiting					

All Other

Therapeutic Products (All Other Therapeutic Products)	SS
Warfarin Sodium (Warfarin Sodium)	C
Megestrol Acetate (Megestrol Acetate)	C
Sucralfate (Sucralfate)	C
Esomeprazole (Esomeprazole)	C
Fentanyl (Fentanyl)	C
Metoprolol (Metoprolol)	C
Docusate Sodium (Docusate Sodium)	C

Date:09/20/04ISR Number: 4455883-6Report Type:Expedited (15-DaCompany Report #2004048907
Age: Gender:Female I/FU:F

Outcome PT
Other Anxiety
Bipolar Disorder
Condition Aggravated

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Gun Shot Wound Pain Polytraumatism	Consumer	Neurontin (Gabapentin)	PS		

Date:09/20/04ISR Number: 4459077-XReport Type:Direct Company Report #CTU 227676
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Confusional State Drug Level Increased Hypernatraemia Oxygen Saturation Decreased Pneumonia Aspiration Respiratory Rate Decreased Suicide Attempt		Benztropine Phenytoin Gabapentin Gabapentin Benztropine Phenytoin Quetiapine	PS SS SS C C C C		

Date:09/21/04ISR Number: 4454501-0Report Type:Expedited (15-DaCompany Report #JP-JNJFOC-20040801088
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death OROPHARINGEAL		Cardiac Arrest 41 DAY	Health	Ofloxacin	PS		
Hospitalization - OROPHARINGEAL		Epilepsy 41 DAY	Professional	Rifadine	SS		
Initial or Prolonged OROPHARINGEAL		Hepatic Failure Hepatocellular Damage 30 DAY		Glucophage Hyperium	SS SS		
OROPHARINGEAL		Ketoacidosis 41 DAY		Neurontin	SS		
OROPHARINGEAL		Lactic Acidosis 9 DAY		Furadantine	SS		
OROPHARINGEAL		Respiratory Distress		Aprovel	C		
OROPHARINGEAL		Septic Shock		Ferrograd	C		

Date:09/21/04ISR Number: 4456549-9Report Type:Expedited (15-DaCompany Report #2004059979
Age:82 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Exanthem	Foreign	Neurontin			
Initial or Prolonged	Pruritus	Consumer	(Gabapentin)	PS		ORAL
ORAL			Candesartan Cilexetil (Candesartan Cilexetil)	C		
			Paracetamol (Paracetamol)	C		

Date:09/21/04ISR Number: 4456551-7Report Type:Expedited (15-DaCompany Report #2004059951
Age:50 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Death	Arrhythmia	Foreign	Neurontin			
Other	Cardio-Respiratory Arrest	Health	(Gabapentin)	PS		ORAL
200 MG (100	Chest Pain	Professional				
MG, 2 IN 1	Insomnia					
D), ORAL	Paraesthesia		Mycophenolate Mofetil (Mycophenolate			

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Mofetil)	C
Prednisone	
(Prednisone)	C
Ciclosporin	
(Ciclosporin)	C
Calcium Carbonate	
(Calcium Carbonate)	C
Ranitidine	
Hydrochloride	
(Ranitidine	
Hydrochloride)	C
Isoniazid	
(Isoniazid)	C
Bactrim	
(Sulfamethoxazole,	
Trimethoprim)	C

Date:09/21/04ISR Number: 4456633-XReport Type:Expedited (15-DaCompany Report #2004063495
 Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Consumer	Neurontin			
		Coordination Abnormal		(Gabapentin)	PS		ORAL
600 MG (600		Fall					
MG, 1 IN 1		Movement Disorder					
D), ORAL							

Atorvastatin	
(Atorvastatin)	C
Nifedipine	
(Nifedipine)	C
Metformin	
Hydrochloride/Rosigl	
itazone (Metformin	
Hydrochloride,	
Rosiglitazone)	C
Glibenclamide	
(Glibenclamide)	C
Metoprolol	
(Metoprolol)	C
Ramirpil (Ramipril)	C
Folic Acid (Folic	
Acid)	C
Magnseium	

(Magnesium) C
Multivitamins
(Ascorbic Acid,
Egocarciferol, Folic
Acid, Nicotinamide,
Panthenol, Retinol, C

Date:09/21/04ISR Number: 4456636-5Report Type:Expedited (15-DaCompany Report #2004051151
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG (600 MG, 3 IN 1 D), ORAL		Crohn'S Disease Drug Ineffective Nerve Degeneration Osteosclerosis Pain	Consumer	Neurontin (Gabapentin) Celecoxib (Celecoxib) Rofecoxib	PS SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Rofecoxib) C

Date:09/21/04ISR Number: 4456638-9Report Type:Expedited (15-DaCompany Report #2004065245
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Abdominoplasty Blood Glucose False	Consumer	Neurontin (Gabapentin)	PS		ORAL
150 MG (150 MG, 1 IN 1 D), ORAL		Positive Blood Glucose Increased Condition Aggravated Diabetic Neuropathy		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS		ORAL
		Hysterectomy					
		Insomnia		Metformin (Metformin)	C		
				Salbutamol (Salbutamol)	C		
				Pantoprazole (Pantoprazole)	C		
				Prazyne (Diaste, Pancreatin, Pepsin, Simeticone)	C		
				Celecoxib (Celecoxib)	C		

Date:09/21/04ISR Number: 4457066-2Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040903625
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - ORAL		Acidosis Acute Respiratory	Literature Health	Acetaminophen (Acetaminophen)	PS		ORAL
Initial or Prolonged Other ORAL		Distress Syndrome Alanine Aminotransferase Increased	Professional	Olanzapine (Olanzapine)	SS		ORAL
				Buspirone			

ORAL	Aspartate	(Buspirone)	SS	ORAL
ORAL	Aminotransferase Increased	Paroxetine (Paroxetine)	SS	ORAL
ORAL	Blood Calcium Decreased Cardiac Arrest	Gabapentin (Gabapentin)	SS	ORAL
ORAL	Coagulopathy Coma	Levothyroxine (Levothyroxine)	SS	ORAL
ORAL	Compartment Syndrome Exposure To Toxic Agent	Estrogens (Estrogens)	SS	ORAL
ORAL	Haemodialysis Labile Blood Pressure Renal Failure	Medication (All Other Therapeutic Products)	SS	ORAL
ORAL	Toxicologic Test Abnormal			

Date:09/22/04ISR Number: 4457381-2Report Type:Expedited (15-DaCompany Report #2004052386
Age:63 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL Other	Cerebrovascular Accident Hearing Impaired Spinal Column Stenosis	Consumer	Neurontin (Gabapentin) Hydrocodone (Hydrocodone) Paracetamol (Paracetamol)	PS C C		ORAL

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Freedom Of Information (FOI) Report

Date:09/22/04ISR Number: 4457462-3Report Type:Expedited (15-DaCompany Report #2004064324

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin (Gabapentin)	PS		

Date:09/22/04ISR Number: 4457470-2Report Type:Expedited (15-DaCompany Report #2004065293

Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abasia Anorexia	Consumer	Gabapentin (Gabapentin)	PS		ORAL
ORAL Other		Aphagia Back Disorder Back Pain Burning Sensation Difficulty In Walking Fatigue Feeling Abnormal Gastric Disorder Gastritis Hypoaesthesia Listless Malaise Nausea Nervous System Disorder Restlessness Throat Cancer Treatment Noncompliance Vomiting Weight Decreased		Glibenclamide (Glibenclamide)	C		

Date:09/22/04ISR Number: 4457777-9Report Type:Expedited (15-DaCompany Report #2004065273

Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abasia	Foreign	Gabapentin			

200 MG (100
MG, 2 IN 1D),
ORAL

Pleurothotonus	Health	Gabapentin)	PS	ORAL
Wheelchair User	Professional			

Rivastigmine Tartrate (Rivastigmine Tartrate0	C
Ramipril (Ramipril)	C
Calcium (Calcium)	C
Oxazepam (Oxazepam)	C
Warfarin Sodium (Warfarin Sodium)	C
Fluticasone Propionate (Fluticasone Propionate)	C
Salbutamol (Salbutamol)	C
Pantoprazole (Pantoprazole)	C
Allopurinol (Allopurinol)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/22/04ISR Number: 4457784-6Report Type:Expedited (15-DaCompany Report #2004065288

Age:84 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Condition Aggravated Disorientation Fall Gait Disturbance Nervous System Disorder Parkinson'S Disease Somnolence Vertigo Wrist Fracture	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Madopar (Benserazide Hydrochloride, Levodopa) Paroxetine Hydrochloride (Paroxetine Hydrochloride) Furosemide (Furosemide) Alprazolam (Alprazolam) Oemprazole(Omeprazol e) Diltazem Hydrochloride (Diltiazem Hydrochloride) Ginko Biloba Extract (Ginko Biloba Extract)	PS C C C C C		

Date:09/22/04ISR Number: 4458069-4Report Type:Expedited (15-DaCompany Report #2002069039

Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG (DAILY), ORAL	Anxiety Condition Aggravated Disease Recurrence	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
150 MG (1D), ORAL	Grand Mal Convulsion Loss Of Consciousness Myoclonic Epilepsy		Clozapine (Clozapine)	C		ORAL
	Petit Mal Epilepsy Urinary Incontinence		Venlafaxine (Venlafaxine)	C		

Panadeine Co
(Codeine Phosphate,
Paracetamol) C

Date:09/23/04ISR Number: 4461200-8Report Type:Expedited (15-DaCompany Report #2004067531

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Anxiety	Consumer	Neurontin			
Other		Completed Suicide		(Gabapentin)	PS		ORAL
ORAL		Drug Abuser		All Other			
		Drug Interaction		Therapeutic Products			
		Pain		(All Other			
				Therapeutic			
				Products)	SS		
				Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	SS		

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Freedom Of Information (FOI) Report

Date:09/23/04ISR Number: 4461209-4Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040903683

Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression	Literature	Extra Strength			
Hospitalization -		Anion Gap Increased	Health	Tylenol			
Initial or Prolonged		Blood Bicarbonate	Professional	(Acetaminophen)	PS		ORAL
ORAL							
Other		Decreased		Citalopram			
ORAL		Blood Glucose Decreased		(Citalopram)	SS		ORAL
ORAL							
ORAL		Blood Ph Decreased		Clonazepam			
ORAL		Coma		(Clonazepam)	SS		ORAL
ORAL							
ORAL		Meningitis		Buspirone			
ORAL		Paralysis		(Buspirone)	SS		ORAL
ORAL							
ORAL		Pco2 Decreased		Fexofenadine			
ORAL		Pco2 Increased		(Fexofenadine)	SS		ORAL
ORAL							
ORAL		Sepsis		Gabapentin			
ORAL		Toxicologic Test Abnormal		(Gabapentin)	SS		ORAL
ORAL							
ORAL		Transaminases Increased		Dicyclomine/Phenobar			
ORAL		White Blood Cell Count		bital (Merbentyl			
ORAL		Decreased		With Phenobarbitone)	SS		ORAL
ORAL							
ORAL				Olanzapine			
ORAL				(Olanzapine)	SS		ORAL
ORAL							
ORAL				Topiramate			
ORAL				(Topiramate)	SS		ORAL

Date:09/23/04ISR Number: 4479088-8Report Type:Periodic

Company Report #208644

Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Diplopia	Health	Avastin			
		Myoclonus	Professional	(Bevacizumab) Pwdr &			
				Solvent, Infusion			
				Soln, 100mg	PS		
INTRAVENOUS	10 MG/KG,						

Q2W,

INTRAVENOUS

ORAL

Neurontin (Gabapentin)	SS	ORAL
Lithium (Lithium Nos)	C	
Oxycodone (Oxycodone)	C	
Vioxx (Rofecoxib)	C	

Date:09/24/04ISR Number: 4462239-9Report Type:Expedited (15-DaCompany Report #2004051197
 Age:80 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 3600 MG (3 IN 1 D), ORAL	Cardiac Disorder Creatinine Renal Clearance Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL

Lisinopril (Lisinopril)	C
Atorvastatin (Atorvastatin)	C
Fenofibrate (Fenofibrate)	C
Ranitidine Hydrochloride (Ranitidine Hydrochloride)	C

Freedom Of Information (FOI) Report

Hydromorphone Hydrochloride (Hydromorphone Hydrochloride)	C
Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotinamide, Metoprolol Tartrate (Metoprolol Tartrate)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C
Sertraline Hydrochloride (Sertraline Hydrochloride)	C
Amlodipine Besilate (Amlodipine Besilate)	C
Hydrochlorothiazide (Hydrochlorothiazide)	C

Date:09/24/04ISR Number: 4462242-9Report Type:Expedited (15-DaCompany Report #2004042915
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin			
Other		Arthropathy	Health	(Gabapentin)	PS		
		Back Disorder	Professional	All Other			
		Balance Disorder		Therapeutic Products			
		Bone Disorder		(All Other			
		Drug Ineffective		Therapeutic			
				Products)	C		

Date:09/24/04ISR Number: 4462247-8Report Type:Expedited (15-DaCompany Report #2004065605
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional	Neurontin (Gabapentin)	PS		
Date:09/24/04ISR Number: 4462249-1Report Type:Expedited (15-DaCompany Report #2004065631 Age:48 YR Gender:Female I/FU:I							
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Optic Neuritis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 2 IN 1 D), ORAL				Sertraline Hydrochloride (Sertraline Hydrochloride) Tizanidine	C		
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Freedom Of Information (FOI) Report

(Tizanidine) C
 Levothyroxine Sodium
 (Levothyroxine Sodium) C

Date:09/24/04ISR Number: 4462266-1Report Type:Expedited (15-DaCompany Report #2004066809

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (100 MG, 3 IN 1 D), ORAL		Condition Aggravated Convulsion Petit Mal Epilepsy	Consumer	Neurontin (Gabapentin)	PS		ORAL

Valproate Semisodium
 (Valproate Semisodium) C
 Citalopram
 Hydrobromide
 (Citalopram Hydrobromide) C
 Estrogens Conjugated
 (Estrogens Conjugated) C

Date:09/24/04ISR Number: 4462268-5Report Type:Expedited (15-DaCompany Report #2004066647

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional Company Representative	Neurontin (Gabapentin) Vicodin (Hydrocodone Bitartrate, Paracetamol)	PS C		

Date:09/24/04ISR Number: 4462285-5Report Type:Expedited (15-DaCompany Report #2004052217

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Activities Of Daily Living Impaired	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG (300 MG, 1 IN 1 D), ORAL		Blood Pressure Increased					
AS NEEDED (5 MG), ORAL		Condition Aggravated					
		Dizziness		Diazepam (Diazepam)	SS		ORAL
		Drug Interaction					
		Fatigue		Levothyroxine Sodium (Levothyroxine Sodium)	C		
		Feeling Abnormal		Estrogens Conjugated (Estrogens Conjugated)	C		
		Gait Disturbance		Potassium Chloride (Potassium Chloride)	C		
		Hangover		Furosemide (Furosemide)	C		
		Insomnia		Hyzaar (Hydrochlorothiazide , Losartan Potassium)	C		
		Knee Arthroplasty		Vicodin (Hydrocodone			
		Lumbar Puncture Abnormal					
		Multiple Sclerosis					
		Pain					
		Renal Disorder					
		Somnolence					
		Speech Disorder					
		Thyroid Disorder					
		Vision Blurred					

Freedom Of Information (FOI) Report

Bitartrate,
Paracetamol) C

Date:09/24/04ISR Number: 4462287-9Report Type:Expedited (15-DaCompany Report #2003034280
Age:39 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Convulsion	Consumer Health Professional	Neurontin (Gabapentin) Diazepam (Diazepam) Oxycocet (Paracetamol, Oxycodone Hydrochloride) Ibuprofen (Ibuprofen)	PS C C C		

Date:09/24/04ISR Number: 4462288-0Report Type:Expedited (15-DaCompany Report #2004049595
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1200 MG (600 MG, 2 IN 1 D), ORAL		Cardiac Disorder Coronary Artery Occlusion Diabetes Mellitus Inadequate Control Drug Ineffective Exposure To Toxic Agent Lymphoma Movement Disorder Myocardial Infarction Neuropathy	Consumer Health Professional	Neurontin (Tablets) (Gabapentin) Atenolol (Atenolol) Isosorbide (Isosorbide) Acetylsalicylic Acid (Acetylsalicylic Acid) Tocopherol (Tocopherol) Pyridoxine Hydrochloride (Pyridoxine Hydrochloride) Cyanocobalamin (Cyanocobalamin)	PS C C C C C C C		ORAL

Date:09/24/04ISR Number: 4462351-4Report Type:Expedited (15-DaCompany Report #2004065552

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Life-Threatening	Deformity	Consumer	Neurontin			
Disability	Drug Ineffective		(Gabapentin)	PS		
Other	Injury					
	Pain					
	Performance Status					
	Decreased					
	Suicide Attempt					

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Date:09/24/04ISR Number: 4462352-6Report Type:Expedited (15-DaCompany Report #2004065315

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure During Pregnancy Drug Withdrawal Syndrome Neonatal Premature Baby	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:09/24/04ISR Number: 4462436-2Report Type:Expedited (15-DaCompany Report #2004042649

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 2700 MG (900 Other MG, 3 IN 1 D), ORAL		Amnesia Balance Disorder Disorientation Overdose Tremor Vision Blurred	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Visual Disturbance		Carbamazepine (Carbamazepine)	SS		ORAL
				Potassium Chloride (Potassium Chloride)	C		
				Simvastatin (Simvastatin)	C		
				Furosemide (Furosemide)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Baclofen (Baclofen)	C		

Date:09/24/04ISR Number: 4462437-4Report Type:Expedited (15-DaCompany Report #2004063520

Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia	Consumer	Neurontin			

600 MG (300	Anxiety	(Gabapentin)	PS	ORAL
MG, 2 IN 1	Chest Discomfort			
D), ORAL	Depression			
1 IN 1 D,	Drug Effect Decreased	Zoloft (Sertraline)	SS	ORAL
ORAL	Drug Interaction			
	Headache	Escitalopram		
	Heart Rate Increased	(Escitalopram)	SS	
	Myocardial Infarction			
	Neck Pain			
	Refusal Of Treatment By			
	Patient			

Date:09/24/04ISR Number: 4462461-1Report Type:Expedited (15-DaCompany Report #2004058684
Age:54 YR Gender:Female I/FU:F

Outcome	PT
Other	Arrhythmia
	Asthma
	Crying
	Depression
	Diarrhoea
	Disorientation
	Dizziness

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG (1 IN 1 D), ORAL	(1 IN 2 D), ORAL	Fall Feeling Abnormal Feeling Drunk Homicidal Ideation Hypertension Hyperventilation Kidney Infection Multiple Allergies Multiple Sclerosis Nausea Neuropathy Pain Pain In Extremity Respiratory Rate Increased Suicidal Ideation Thinking Abnormal Urine Abnormality Visual Acuity Reduced Vomiting Projectile	Consumer Health Professional	Zoloft (Sertraline) Neurontin (Gabapentin) Potassium Chloride (Potassium Chloride) Losartan Potassium (Losartan Potassium) Lansoprazole (Lansoprazole) Metoclopramide (Metoclopramide) Salbutamol (Salbutamol)	PS SS C C C C C		ORAL ORAL

Date:09/27/04ISR Number: 4462696-8Report Type:Expedited (15-DaCompany Report #2004051397
Age:77 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Anticonvulsant Drug Level Decreased Balance Disorder Convulsion Fatigue Memory Impairment Osteoporosis Pharmaceutical Product Complaint Somnolence Speech Disorder	Health Professional	Dilantin Suspension (Phenytoin Sodium) Neurontin (Gabapentin) Phenytoin (Phenytoin)	PS SS SS		

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Systemic Lupus Erythematosus	Consumer	Neurontin (Gabapentin)	PS		
				Fentanyl (Fentanyl)	C		
				Levobunolol Hydrochloride (Levobunolol Hydrochloride)	C		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		

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Date:09/27/04ISR Number: 4462777-9Report Type:Expedited (15-DaCompany Report #2004066759

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

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
80 MG (80 MG , 1 IN 1 D)		Cardiac Disorder Diabetes Mellitus Myocardial Infarction	Consumer	Lipitor (Atorvastatin)	PS		
900 MG (300 MG, 3 IN 1 D)		Nervous System Disorder Tremor		Neurontin (Gabapentin)	SS		
20 MG (10 MG, 2 IN 1D)				Glipizide (Glipizide)	SS		
25 MG (25 MG, 1 IN 1 D)				Atenolol (Atenolol)	SS		
60 MG (30 MG, 2 IN 1 D)				Lansoprazole (Lansoprazole)	SS		
2.5 MG (2.5 MG, 1 IN 1 D)				Ramipril (Ramipril)	SS		
				Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
				Glyceryl Trinitrate (Glyceryl Trinitrate)	C		

Date:09/27/04ISR Number: 4462779-2Report Type:Expedited (15-DaCompany Report #2004065930

Age:56 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Death	Abnormal Behaviour	Literature	Neurontin	
	Completed Suicide	Consumer	(Gabapentin)	PS
	Memory Impairment			
	Social Avoidant Behaviour			

Date:09/27/04ISR Number: 4462796-2Report Type:Expedited (15-DaCompany Report #2004051408

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bladder Disorder	Consumer	Neurontin			
		Constipation		(Gabapentin)	PS		ORAL
3000 MG (300		Faecaloma					
MG), ORAL		Neoplasm		Clonazepam			
		Renal Cyst		(Clonazepam)	SS		ORAL
3 MG, ORAL		Urinary Retention		Oxycodone			
		Urinary Tract Infection		Hydrochloride			
		X-Ray Abnormal		(Oxycodone			
60 MG, ORAL				Hydrochloride)	SS		ORAL
				Venlafaxine			
				Hydrochloride			
				(Venlafaxine			
375 MG, ORAL				Hydrochloride)	SS		ORAL
				Quetiapine Fumarate			
				(Quetiapine			
				Fumarate)	C		
				Valsartan			
				(Valsartan)	C		

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Date:09/27/04ISR Number: 4462799-8Report Type:Expedited (15-DaCompany Report #2004066878
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Depression	Consumer	Neurontin			
Other		Heart Valve Insufficiency		(Gabapentin)	PS		ORAL
900 MG (300							
MG, 3 IN 1		Myocardial Infarction					
D), ORAL				Atenolol (Atenolol)	C		
				Simvastatin			
				(Simvastatin)	C		
				Fluoxetine			
				(Fluoxetine)	C		

Date:09/27/04ISR Number: 4462807-4Report Type:Expedited (15-DaCompany Report #2004066805
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Ineffective	Consumer	Neurontin			
300 MG (100		Epistaxis		(Gabapentin)	PS		ORAL
MG, 3 IN 1		Feeling Abnormal					
D), ORAL		Hypersensitivity					
		Vision Blurred		Promethazine			
		White Blood Cell Count		Hydrochloride			
		Increased		(Promethazine			
ORAL				Hydrochloride)	SS		ORAL
				Vicodin (Hydrocodone			
				Bitartrate,			
				Paracetamol)	C		
				Cyclobenzaprine			
				Hydrochloride			
				(Cyclobenzaprine			
				Hydrochloride)	C		

Date:09/27/04ISR Number: 4462930-4Report Type:Expedited (15-DaCompany Report #2004053846
Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain	Foreign	Neurontin			
Other		Difficulty In Walking	Consumer	(Gabapentin)	PS		
		Headache		Nortriptyline			
		Intervertebral Disc		(Nortriptyline)	C		
		Protrusion		Clonazepam			
		Pain In Extremity		(Clonazepam)	C		
				All Other			
				Contraceptive Device			
				(Intrauterine			
				Contraceptive			
				Uterine)	C		

Date:09/28/04ISR Number: 4462510-0Report Type:Expedited (15-DaCompany Report #2004065277
Age:73 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Cholestasis
Initial or Prolonged	Chronic Hepatitis
Other	Hepatic Steatosis
	Hepatomegaly
	Intra-Abdominal

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Haemangioma
Myalgia
Pyrexia

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
ORAL		Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:09/28/04ISR Number: 4462604-XReport Type:Expedited (15-DaCompany Report #2004034256
Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	1200 MG (400 MG, 3 IN 1 D), ORAL	Aplastic Anaemia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:09/28/04ISR Number: 4462615-4Report Type:Expedited (15-DaCompany Report #2004066974
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	ORAL	Hyperpyrexia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:09/28/04ISR Number: 4463325-XReport Type:Expedited (15-DaCompany Report #2004068340
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Life-Threatening	Condition Aggravated Headache Overdose	Consumer	Neurontin (Gabapentin) All Other	PS		

Panic Attack
Suicide Attempt

Therapeutic Products
(All Other
Therapeutic
Products) SS

Date:09/28/04ISR Number: 4463334-0Report Type:Expedited (15-DaCompany Report #HQWYE119220SEP04
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction Panic Reaction Suicidal Ideation Thinking Abnormal	Consumer	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		
SEE IMAGE							
				Neurontin (Gabapentin,)	SS		
SEE IMAGE							

Date:09/28/04ISR Number: 4463337-6Report Type:Expedited (15-DaCompany Report #2004068235
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Surgery Unevaluable Event	Consumer	Neurontin (Gabapentin)	PS		

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Date:09/28/04ISR Number: 4465134-4Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 228219

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - SEE IMAGE	Depression		Neurontin 300 Mg	PS		ORAL
Initial or Prolonged	Somnolence		.	C		
			Asa	C		
			Clonazepam	C		
			Clopidogrel	C		
			Donepezil	C		
			Ferrous Sulfate	C		
			Ipratropium	C		
			Lisinopril	C		
			Metoprolol	C		
			Omeprazole	C		

Date:09/29/04ISR Number: 4463846-XReport Type:Expedited (15-DaCompany Report #2004060890
 Age:83 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 1200 MG (400 Initial or Prolonged MG, 3 IN 1	Confusional State Narcolepsy Oxygen Saturation Decreased	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
D), ORAL			Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		ORAL
20 MG (20 MG, 1 IN 1 D), ORAL						

Date:09/29/04ISR Number: 4463875-6Report Type:Expedited (15-DaCompany Report #2004068343
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Confusional State	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Neuropathy Pain Pyrexia		Atrovastatin Calcium (Atorvastatin Calcium) Bromazepam (Bromazepam)	C C		

Date:09/29/04ISR Number: 4464543-7Report Type:Expedited (15-DaCompany Report #2004234715US
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Celebrex (Celecoxib)	PS		ORAL
200 MG, QD,		Bone Graft					
ORAL		Cognitive Deterioration Confusional State		Neurontin (Gabapentin)	SS		ORAL
600 MG, TID,		Dental Caries					
ORAL; 600		Facial Bones Fracture					
MG, BID,		Gingival Hypoplasia Impaired Healing Loss Of Consciousness Oral Infection Tooth Fracture Vision Blurred		Wellbutrin (Amfebutamone Hydrochloride) Seroquel (Quetiapine) Elavil	C C		

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Freedom Of Information (FOI) Report

(Amitriptyline
Hydrochloride) C
Synthroid(Levothyrox
ine Sodium) C
Tricor C
Pravachol
(Pravastatin Sodium) C

Date:09/29/04ISR Number: 4465317-3Report Type:Expedited (15-DaCompany Report #2004066805
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Choking Cough	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG, 3 IN 1 D), ORAL		Dizziness Drug Ineffective					
ORAL		Dry Mouth Dysphagia Dyspnoea Epistaxis		Promethazine Hydrochloride (Promethazine Hydrochloride)	SS		ORAL
		Fatigue Feeling Abnormal Hypersensitivity Oropharyngeal Swelling Pyrexia Somnolence Tongue Exfoliation Vision Blurred Vomiting White Blood Cell Count Increased		Vicodin (Hydrocodone Bitartrate, Paracetamol) Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride)	C C		

Date:09/29/04ISR Number: 4465340-9Report Type:Expedited (15-DaCompany Report #2004068315
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Disorder	Consumer	Neurontin			

1600 MG (800
 MG, 2 IN 1
 D), UNKNOWN

Drug Ineffective (Gabapentin) PS
 Economic Problem
 Treatment Noncompliance

Insulin (Insulin) C

Date:09/30/04ISR Number: 4465944-3Report Type:Direct Company Report #CTU 228423
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 7200 MG DAILY Initial or Prolonged ORAL		Coordination Abnormal Dysarthria Medication Error		Gabapentin 800 Mg Trazodone Albuterol Asa Felodipine Ferrous Sulfate Furosemide Glyburide Ipratropium Rabeprazole Lisinopril Isosorbide	PS C C C C C C C C C C		ORAL

Freedom Of Information (FOI) Report

Mononitrate	C
Potassium Chloride	C
Multivitamin	C
Toprol Xl	C
Ibuprofen	C
Nortriptyline	C

Date:09/30/04ISR Number: 4466065-6Report Type:Expedited (15-DaCompany Report #2002069039
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Foreign	Gabapentin			
		Clonic Convulsion	Health	(Gabapentin)	PS		ORAL
1800 MG							
(DAILY), ORAL		Condition Aggravated	Professional				
		Grand Mal Convulsion		Clozapine			
		Loss Of Consciousness		(Clozapine)	SS		ORAL
ORAL		Myoclonic Epilepsy		Venlafaxine	C		
		Myoclonus		Panadeine Co			
		Neuralgia		(Codeine Phosphate,			
		Petit Mal Epilepsy		Paracetamol)	C		
		Urinary Incontinence					

Date:09/30/04ISR Number: 4466069-3Report Type:Direct Company Report #CTU 228379
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aggression		Neurontin 100 Mg			
		Amnesia		Pfizer	PS	Pfizer	ORAL
1 CAPSUL A							
NIGHT ORAL		Anxiety					
		Crying					
		Discomfort					
		Emotional Disorder					
		Family Stress					
		Hallucination					
		Hostility					
		Hypersomnia					
		Impaired Work Ability					

Insomnia
Mood Altered
Paranoia
Suspiciousness

Date:09/30/04ISR Number: 4466385-5Report Type:Expedited (15-DaCompany Report #2004067531
Age:42 YR Gender:Male I/FU:F

Outcome	PT
Death	Accident
Hospitalization -	Alcohol Use
Initial or Prolonged	Anxiety
Other	Arthralgia
	Atherosclerosis
	Brain Oedema
	Burning Sensation
	Cardio-Respiratory Arrest
	Cardiomegaly
	Coma
	Coronary Artery
	Atherosclerosis
	Disease Recurrence

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Drug Abuser Drug Interaction Drug Screen Positive	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, 2 IN 1 D), ORAL		Excoriation Feeling Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Haematemesis					
		Hepatic Steatosis					
ORAL		Hypertensive Heart Disease		Benadryl (Diphenhydramine)	SS		ORAL
1-3 TABLETS Q12H, ORAL		Hypoaesthesia Ill-Defined Disorder Insomnia Intentional Self-Injury		Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL
		Laceration					
1-2 EVERY 4-6 HOURS (10 MG), ORAL		Major Depression Multi-Organ Disorder		Hydrocodone (Hydrocodone)	SS		ORAL
		Muscle Spasms					
		Muscle Spasticity					
10 MG (10 MG, 1 IN 1 D), ORAL		Pain Pulmonary Oedema		Zolpidem Tartrate (Zolpidem Tartrate)	SS		ORAL
		Radiculopathy					
		Renal Cyst					
80 MG (80 MG, 1 IN 1 D), ORAL		Suicide Attempt		Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		ORAL
				Amitriptyline Hydrochloride (Amitriptyline			

Hydrochloride)	C
Fentanyl (Fentanyl)	C
Cyclobenzaprine	
Hydrochloride	
(Cyclobenzaprine	
Hydrochloride)	C
Bupropion	
Hydrochloride	
(Bupropion	
Hydrochloride)	C
Quinine Sulfate	
(Quinine Sulfate)	C
Sertraline	
Hydrochloride	
(Sertraline	
Hydrochloride)	C
Trazodone	
(Trazodone)	C
Hydroxyzine Embonate	
(Hydroxyzine	
Embonate)	C

Date:09/30/04ISR Number: 4466386-7Report Type:Expedited (15-DaCompany Report #2004068966

Age: Gender:Unknown I/FU:I

Outcome	PT
Other	Arthralgia
	Pain In Extremity
	Sleep Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Surgery

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Consumer	Neurontin (Gabapentin)	PS		
			Hydrocortisone (Hydrocortisone)	SS		

Date:09/30/04ISR Number: 4466393-4Report Type:Expedited (15-DaCompany Report #2004039295
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Neurontin (Gabapentin)	PS		ORAL
Other		Leukopenia Neutropenia	Professional	Nortriptyline Hydrochloride (Nortriptyline Hydrochloride)	C		
3600 MG, ORAL							

Date:09/30/04ISR Number: 4466394-6Report Type:Expedited (15-DaCompany Report #2004006430
Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Health	Neurontin (Gabapentin)	PS		ORAL
Other		Convulsion Pharmaceutical Product	Professional				
1200 MG (2 IN		Complaint					
1 D), ORAL		Therapeutic Response Decreased Unevaluable Event		Dilantin Kapseals (Phenytoin Sodium) (Phenytoin Sodium)	SS		ORAL
ORAL		Vitamin B Complex Deficiency		Phenytoin (Phenytoin)	SS		

Date:09/30/04ISR Number: 4466396-XReport Type:Expedited (15-DaCompany Report #2004068968
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin			
2400 MG		Migraine		(Gabapentin)	PS		
		Osteoarthritis		Muscle Relaxant			
		Road Traffic Accident		(Muscle Relaxant)	SS		
				Vicodin (Hydrocodone			
				Bitartrate,			
				Paracetamol)	SS		
				Ibuprofen			
				(Ibuprofen)	SS		
				Paracetamol			
				(Paracetamol)	SS		
				Exedrin P.M.			
				(Diphenhydramine			
				Citrate,			
				Paracetamol)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:09/30/04ISR Number: 4466733-6Report Type:Expedited (15-DaCompany Report #2004051397
Age:77 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder	Health	Dilantin Suspension			
		Convulsion	Professional	(Phenytoin Sodium)	PS		
		Drug Effect Decreased		Neurontin			
		Fatigue		(Gabapentin)	SS		
		Memory Impairment		Phenytoin			
		Osteoporosis		(Phenytoin)	SS		
		Somnolence					
		Speech Disorder					

Date:09/30/04ISR Number: 4466778-6Report Type:Expedited (15-DaCompany Report #2004060932
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Appendicectomy	Consumer	Neurontin			
		Bladder Repair		(Gabapentin)	PS		ORAL
ORAL							
200 MG (200		Hysterectomy		Celebrex (Celecoxib)	SS		ORAL
MG, 1 IN 1		Inflammation					
D), ORAL		Muscle Twitching					
		Post Procedural		Aciclovir			
		Complication		(Aciclovir)	C		
		Post Procedural Pain		Pantoprazole			
		Treatment Noncompliance		(Pantoprazole)	C		
				Diazepam (Diazepam)	C		
				Quetiapine Fumarate			
				(Quetiapine			
				Fumarate)	C		
				Fluoxetine			
				Hydrochloride			
				(Fluoxetine			
				Hydrochloride)	C		

Date:10/01/04ISR Number: 4467699-5Report Type:Expedited (15-DaCompany Report #2004054409
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability 300 MG		Back Pain Gait Disturbance	Consumer	Neurontin (Gabapentin)	PS		
		Neck Pain Pain In Extremity		Metamucil "Procter & Gamble" (Glucose Monohydrate, Ispaghula Husk) Herbal Nos/Vitamins Nos	C C		

Date:10/01/04ISR Number: 4467700-9Report Type:Expedited (15-DaCompany Report #2004068375
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1600 MG, ORAL		Alanine Aminotransferase Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Aspartate Aminotransferase Increased Hepatotoxicity					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/01/04ISR Number: 4470499-3Report Type:Expedited (15-DaCompany Report #2004AL000197
Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other Required Intervention to Prevent Permanent Impairment/Damage 40 MG; QD 900 MG; QD		Agitation Confusional State Drug Ineffective Dry Mouth Emotional Distress Haemoglobin Decreased Hallucination Hypotension Myoclonus Pain Paresis Somnolence Vomiting	Foreign Literature Consumer Other	Kadian(Morphine Sulfate Sustained Release) Capsules 100 Mg Methadone Hydrochloride (,Nul>) Gabapentin (<Null>) Bupivacaine Clonidine Ketamine Promazine Pilocarpine Dexamethasone Clonazepam	PS SS SS C C C C C C C	Alpharma	

Date:10/04/04ISR Number: 4467000-7Report Type:Expedited (15-DaCompany Report #2004068261
Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL 900 MG (300 MG, 3 IN 1 D), ORAL		Arthralgia Lethargy Myalgia Palpitations Somnolence Transient Ischaemic Attack Trigeminal Neuralgia	Consumer	Lipitor (Atorvastatin) Neurontin (Gabapentin) Ezetimibe (Ezetimibe)	PS SS C		ORAL ORAL

Date:10/04/04ISR Number: 4467093-7Report Type:Expedited (15-DaCompany Report #2004068964
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG, 3 IN 1 D)		Asthenia Balance Disorder Coordination Abnormal Depression Difficulty In Walking Dizziness Fall Fatigue Feeling Abnormal Gait Disturbance Headache Limb Injury Loss Of Consciousness Medication Error Mood Swings Relationship Breakdown Somnolence Thinking Abnormal Upper Limb Fracture	Consumer	Neurontin (Gabapentin) Sertraline Hydrochloride (Sertraline Hydrochloride) Lorazepam (Lorazepam)	PS C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/04/04ISR Number: 4467098-6Report Type:Expedited (15-DaCompany Report #2004070205

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Malaise	Consumer	Neurontin (Gabapentin)	PS		
		Spinal Cord Injury		Rofecoxib (Rofecoxib)	SS		

Date:10/04/04ISR Number: 4467372-3Report Type:Expedited (15-DaCompany Report #2004069049

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Cholelithiasis	Consumer	Neurontin (Gabapentin)	PS		ORAL
SEE IMAGE		Gastric Haemorrhage					
Other		Oesophageal Disorder		Clonazepam (Clonazepam)	C		
		Stomach Discomfort		Baclofen (Baclofen)	C		
				Oxycodone Hydrochloriide (Oxycodone Hydrochloride)	C		

Date:10/04/04ISR Number: 4467818-0Report Type:Expedited (15-DaCompany Report #2004065273

Age:76 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abasia	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
200 MG (100 MG, 2 IN 1 D), ORAL		Pleurothotonus	Professional				
				Rivastigmine (Rivastigmine)	SS		ORAL
6 MG (3 MG, 2 IN 1 D), ORAL							

Salmeterol Xinafoate
(Salmeterol
Xinafoate) C
Calcium (Calcium) C
Oxazepam (Oxazepam) C
Warfarin Sodium
(Warfarin Sodium) C
Fluticasone
Propionate
(Fluticasone
Propionate) C
Salbutamol
(Salbutamol) C
Pantoprazole
(Pantoprazole) C
Allopurinol
(Allopurinol) C
Ramipril C

Date:10/04/04ISR Number: 4468045-3Report Type:Expedited (15-DaCompany Report #2004069046
Age:80 YR Gender:Male I/FU:I

Outcome PT
Death Drug Ineffective
Erectile Dysfunction

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Freedom Of Information (FOI) Report

Myocardial Infarction

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		
			Nicardipine Hydrochloride (Nicardipine Hydrochloride)	C		
			Glibenclamide (Glibenclamide0)	C		
			Dipyridamole (Dipyridamole)	C		
			Allopurinol (Allopurinol)	C		
			Blopress Plus (Candesartan Cilexetil, Hydrochlorothiazide)	C		
			Vaseretic (Enalapril Maleate, Hydrochlorothiazide)	C		
			Pravastatin Sodium (Pravastatin Sodium)	C		
			Metformin Hydrochloride (Metformin Hydrochloride)	C		
			Acarbose (Acarbose)	C		
			Glyceryl Trinitrate (Glyceryl Trinitrate)	C		

Date:10/05/04ISR Number: 4469449-5Report Type:Expedited (15-DaCompany Report #2004046622
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Neutropenia White Blood Cell Count Decreased	Health Professional	Neurontin (Gabapentin)	PS		

Outcome	PT
Death	Anxiety
Hospitalization -	Atrial Fibrillation
Initial or Prolonged	Blood Creatinine Increased Blood Ph Increased Blood Potassium Decreased Blood Pressure Systolic Increased Blood Sodium Decreased Body Temperature Increased Cardio-Respiratory Arrest Cardiomegaly Diarrhoea

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Myocardial Infarction Neutrophil Count Decreased	Report Source	Product	Role	Manufacturer	Route
ORAL		Neutrophil Count Increased	Foreign Literature Health	Ms Contin Tablets (Morphine Sulfate) Cr Tablet	PS		ORAL
ORAL		Oxygen Saturation Decreased	Professional	Feldene (Piroxicam)	SS		ORAL
ORAL		Pulmonary Oedema	Other	Celebrex (Celecoxib)	SS		ORAL
ORAL		Respiratory Alkalosis Respiratory Rate		Neurontin (Gabapentin)	SS		ORAL
		Increased Urine Output Decreased Ventricular Fibrillation Vomiting White Blood Cell Count Decreased White Blood Cell Count Increased		Avapro (Irbesartan) Prozac (Fluoxetine Hydrochloride)	SS SS		

Date:10/05/04ISR Number: 4469491-4Report Type:Expedited (15-DaCompany Report #2004069182

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Asthenia Economic Problem	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (200 MG, 3 IN 1 D), ORAL		Gait Disturbance Musculoskeletal Stiffness		Methadone (Methadone)	C		
		Vomiting Weight Decreased Weight Fluctuation					

Date:10/05/04ISR Number: 4469509-9Report Type:Expedited (15-DaCompany Report #2004045415

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abortion Spontaneous Drug Exposure During Pregnancy	Health Professional	Neurontin (Gabapentin)	PS		ORAL
3600 MG (1200 MG, 3 IN 1 D), ORAL		Pregnancy	Company Representative	Topiramate (Topiramate)	C		

Date:10/05/04ISR Number: 4470831-0Report Type:Expedited (15-DaCompany Report #2004070771
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 40 MG (20 MG , 2 IN 1 D)		Abnormal Behaviour Aggression Agitation	Literature Health Professional	Ziprasidone (Caps) (Ziprasidone)	PS		
400 MG (200 MG, 2 IN 1 D)		Condition Aggravated Confusional State Disorientation Disturbance In Attention		Carbamazepine (Gabapentin) (Carbamazepine)	SS		
900 MG (300 MG IN 1 D)		Drug Ineffective Hallucinations, Mixed Incoherent Mood Swings Speech Disorder		Gabapentin (Escitalopram) (Gabapentin) Escitalopram	SS		

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Freedom Of Information (FOI) Report

(Escitalopram) SS

10 MG (10 MG,
1 IN 1 D)

Date:10/06/04ISR Number: 4467393-0Report Type:Expedited (15-DaCompany Report #PHNU2004DE00766

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abnormal Behaviour
Initial or Prolonged	Alopecia
Disability	Anosmia
Other	Aphasia
	Basal Ganglion
	Degeneration
	Bradyphrenia
	Cerebellar Syndrome
	Chills
	Cogwheel Rigidity
	Condition Aggravated
	Depression
	Difficulty In Walking
	Disturbance In Attention
	Dysarthria
	Dyskinesia
	Electroencephalogram
	Abnormal
	Encephalopathy
	Erectile Dysfunction
	Fall
	Fatigue
	Gait Disturbance
	General Physical Health
	Deterioration
	Grand Mal Convulsion
	Hypokinesia
	Liver Function Test
	Abnormal
	Malaise
	Masked Facies
	Memory Impairment
	Mental Impairment
	Micrographia
	Mobility Decreased
	Motor Dysfunction
	Movement Disorder

Muscle Spasms
Muscle Twitching
Myalgia
Nerve Conduction Studies
Abnormal
Neuroleptic Malignant
Syndrome
Paralysis
Parkinson'S Disease
Performance Status
Decreased
Posture Abnormal
Regressive Behaviour
Salivary Hypersecretion
Simple Partial Seizures
Speech Disorder
Tongue Paralysis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tremor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
400 mg, QOD			Tegretal	PS	Novartis Sector: Pharma	ORAL
UNKNOWN			Ergenyl "Sanofi-Synthelabo"	SS		
UNKNOWN			Neurontin	SS		
60mg/day			Gabitril	SS		ORAL
100mg/day	549 DAY		Topamax	SS		ORAL
400 mg, QID			Carbabeta	SS		ORAL
400 mg, QID			Carbabeta	SS		ORAL
1DF/day			Movergan	SS		ORAL
.5 mg, QID			Levodopa	SS		ORAL
1mg/day			Parkotil	SS		ORAL
0.20mg/day			Parkotil	SS		ORAL
200 mg, QID			Comtess	SS		ORAL
50 mg, QID			Amantadin	SS		ORAL
62.5 mg, TID			Levodopa	SS		ORAL
400 mg, TID			Madopar	SS		ORAL
UNKNOWN	3DF/Day		Carbabeta	SS		ORAL
UNKNOWN	1000-1500mg/d		Zentropil	C		
ay			Keppra	C		
UNKNOWN			Nacom "Dupont Pharma"	C		

15mg/day	366 DAY	Tasmar	C	
		Remergil	C	ORAL
		L-Dopa	C	
gradually		Levodopa-Carbidopa	C	
increased		Pramipexole	C	
		Pergolide	C	
		Aman	C	
		Levetiracetam	C	
5mg/day		Selegiline	C	
		Phenhydan	C	
UNKNOWN	500 mg, BID	Keppra	C	
		Lamictal		
UNKNOWN		"Glaxosmithkline"	C	

Date:10/06/04ISR Number: 4471209-6Report Type:Expedited (15-DaCompany Report #2004061480
Age:85 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fall	Foreign	Neurontin			
		Rib Fracture	Health	(Gabapentin)	PS		ORAL
900 MG, ORAL		Vertigo	Professional	Amlodipine Besilate			
			Other	(Amlodipine			
				Besilate)	C		
				Paroxetine			
				Hydrochloride			
				(Paroxetine			
				Hydrochloride)	C		
				Rilmenidine			
				Dihydrogen Phosphate			
				(Rilmenidine			
				Dihydrogen			

Freedom Of Information (FOI) Report

Phosphate) C
 Dextropropoxyphene
 (Dextropropoxyphene) C

Date:10/06/04ISR Number: 4471466-6Report Type:Expedited (15-DaCompany Report #2004065345
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (3 IN 1 D), ORAL		Faecal Incontinence Somnolence	Foreign Health Professional Company Representative	Neurontin (Tablets) (Gabapentin) All Other Therapeutic Products (All Other Therapeutic Products) Perindopril (Perindopril) Insulin (Insulin) Panadeine Co (Codeine Phosphate, Paracetamol)	PS C C C C		ORAL

Date:10/06/04ISR Number: 4471808-1Report Type:Expedited (15-DaCompany Report #2004069999
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Confusional State Nervous System Disorder	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/06/04ISR Number: 4471810-XReport Type:Expedited (15-DaCompany Report #2004066809
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Condition Aggravated	Consumer	Neurontin			

300 MG (100 Convulsion (Gabapentin) PS ORAL
MG, 3 IN 1 Petit Mal Epilepsy
D), ORAL

Valproate Semisodium
(Valproate
Semisodium) C
Citalopram
Hydrobromide
(Citalopram
Hydrobromide) C
Estrogens Conjugated
(Estrogens
Conjugated) C

Date:10/06/04ISR Number: 4471812-3Report Type:Expedited (15-DaCompany Report #2004061448
Age: Gender:Male I/FU:F

Outcome PT
Other Breast Cyst
Nephrolithiasis
Neuralgia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Restless Legs Syndrome Scrotal Mass		Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Health Professional	Neurontin (Gabapentin)	PS		
600 MG (300 MG, 2 IN 1 D),		Company Representative				
			Pramipexole (Pramipexole)	C		
			Nortriptyline (Nortriptyline)	C		
			Fexofenadine Hydrochloride (Fexofenadine Hydrochloride)	C		
			Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		

Date:10/06/04ISR Number: 4471916-5Report Type:Expedited (15-DaCompany Report #2004056753
Age:65 YR Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Bone Marrow Depression Leukopenia	Health Professional	Neurontin (Gabapentin)	PS		
Other		Polycythaemia Vera	Company Representative				
1800 MG (600 MG, 3 IN 1 D)							

Date:10/06/04ISR Number: 4471918-9Report Type:Expedited (15-DaCompany Report #2004013492
Age: Gender:Male I/FU:F

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration	Bipolar Disorder Brain Damage Closed Head Injury	Consumer	Neurontin (Gabapentin)	PS		
Other							

Depressed Level Of
Consciousness
Drug Dependence
Drug Withdrawal Syndrome
Fear
Hallucination, Auditory
Hallucination, Visual
Intentional Self-Injury
Malaise
Off Label Use
Pain
Psychotic Disorder
Sensory Disturbance
Stupor
Suicide Attempt

Date:10/07/04ISR Number: 4471949-9Report Type:Expedited (15-DaCompany Report #2004071853
Age:46 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Brain Scan Abnormal
Initial or Prolonged	Depressed Level Of Consciousness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Epilepsy Speech Disorder Stereotypy	Report Source	Product	Role	Manufacturer	Route
			Foreign Health Professional	Neurontin (Gabapentin)	PS		
				Phenytoin Suspension (Phenytoin Suspension)	SS		
				Lamotrigine (Lamotrigine)	SS		
				Topiramate (Topiramate)	SS		
				Valproic Acid (Valproic Acid)	SS		
				Carbamazepine (Carbamazepine)	C		
				Levetiracetam (Levetiracetam)	C		
				Levodopa (Levodopa)	C		
				Entacapone (Entacapone)	C		
				Pergolide (Pergolide)	C		
				Amantadine (Amantadine)	C		

Date:10/07/04ISR Number: 4472077-9Report Type:Expedited (15-DaCompany Report #2004062069

Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cytolytic Hepatitis	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other		Liver Disorder					
1800 MG (600 MG, 3 IN 1 D), ORAL		Metastases To Liver	Company Representative	Omeprazole (Omeprazole)	C		
				Prednisolone Sodium Sulfobenzoate (Prednisolone Sodium Sulfobenzoate)	C		

Tianeptine
(Tianeptine) C
Insulin Human
Injection, Isophane
(Insulin Human
Injection, Isophane) C

Date:10/07/04ISR Number: 4472293-6Report Type:Expedited (15-DaCompany Report #2004062829

Age:61 YR Gender:Female I/FU:F

Outcome	PT
Disability	Arachnoiditis
Other	Back Pain
	Economic Problem
	Fracture Nonunion
	Humerus Fracture
	Migraine
	Post Procedural
	Complication

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Stress Treatment Noncompliance	Report Source	Product	Role	Manufacturer	Route
1600 MG (800 M G, 2 IN 1 D), ORAL			Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Professional	Tizanidine (Tizanidine)	C		
				Carisoprodol (Carisoprodol)	C		
				Temazepam (Temazepam)	C		
				Methadone (Methadone)	C		
				Valdecoxib (Valdecoxib)	C		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:10/07/04ISR Number: 4472322-XReport Type:Expedited (15-DaCompany Report #2004042647
Age:47 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG (600 MG, 3 IN 1 D), ORAL		Condition Aggravated Depression	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
		Insomnia	Professional				
		Neuropathic Pain					
		Somnolence Suicidal Ideation Vision Blurred		Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Valsartan (Valsartan)	C		

Date:10/07/04ISR Number: 4472356-5Report Type:Expedited (15-DaCompany Report #2004070397
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Deformity	Consumer	Neurontin			
Other		Drug Ineffective Pain Polytraumatism Suicide Attempt		(Gabapentin)	PS		

Date:10/08/04ISR Number: 4472088-3Report Type:Expedited (15-DaCompany Report #2004051151
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crohn'S Disease Drug Ineffective	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG (600		Neuropathic Pain	Professional				
MG, 3 IN 1		Neuropathy					
D), ORAL		Osteosclerosis		Celecoxib (Celecoxib) Rofecoxib	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Rofecoxib) SS

Date:10/08/04ISR Number: 4472092-5Report Type:Expedited (15-DaCompany Report #2004062323

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain	Consumer	Neurontin			
		Drug Withdrawal Syndrome	Health	(Gabapentin)	PS		ORAL
6400 MG, ORAL		Muscle Spasms	Professional	Fentanyl (Fentanyl)	C		
		Treatment Noncompliance		Benzocaine	C		
				(Benzocaine)	C		
				Clonidine	C		
				(Clonidine)	C		
				Clopidogrel Sulfate			
				(Clopidogrel			
				Sulfate)	C		
				Temazepam			
				(Temazepam)	C		
				Trazodone			
				(Trazodone)	C		
				Imipramine			
				(Imipramine)	C		
				Levothyroxine Sodium			
				(Levothyroxine			
				Sodium)	C		
				Risperidone			
				(Risperidone)	C		

Date:10/08/04ISR Number: 4472098-6Report Type:Expedited (15-DaCompany Report #2004071938

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Brain Oedema	Consumer	Neurontin (Tablets)			
Initial or Prolonged		Central Nervous System		(Gabapentin)	PS		ORAL
3200 MG (400							
Other		Infection					
MG, 8 IN 1							
		Head Deformity					
		Nervous System Disorder					
D), ORAL		Post Procedural					

Complication

Date:10/08/04ISR Number: 4472122-0Report Type:Expedited (15-DaCompany Report #2004067531
Age:42 YR Gender:Male I/FU:F

Outcome	PT
Death	Accident
Hospitalization -	Alcohol Use
Initial or Prolonged	Anxiety
Other	Atherosclerosis
	Back Pain
	Brain Oedema
	Burning Sensation
	Cardio-Respiratory Arrest
	Cardiomegaly
	Coma
	Coronary Artery
	Atherosclerosis
	Drug Abuser
	Drug Interaction
	Drug Level Above

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Therapeutic Drug Screen Positive Excoriation Feeling Abnormal	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, 2 IN 1 D), ORAL		Haematemesis Hepatic Steatosis Hypertensive Heart Disease	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Hypoaesthesia Intentional Self-Injury		Benadryl (Diphenhydramine)	SS		ORAL
1-3 TABLETS Q12H, ORAL		Laceration Major Depression Muscle Spasms Pain		Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL
1-2 EVERY 4-6 HOURS (10 MG), ORAL		Pulmonary Oedema Renal Cyst Sleep Disorder Due To General Medical Condition, Insomnia Type		Hydrocodone (Hydrocodone)	SS		ORAL
10 MG (10 MG , 1 IN 1 D), ORAL		Snoring Suicidal Ideation Therapy Non-Responder		Zolpidem Tartrate (Zolpidem Tartrate)	SS		ORAL
80 MG (80 MG, 1 IN 1 D), ORAL				Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		ORAL
				Amitriptyline Hydrochloride(Amitri ptyline			

Hydrochloride)	C
Fentanyl (Fentanyl)	C
Cyclobenzaprine	
Hydrochloride	
(Cyclobenzaprine	
Hydrochloride)	C
Bupropion	
Hydrochloride	
(Bupropion	
Hydrochloride)	C
Quinine Sulfate	
(Quinine Sulfate)	C
Sertraline	
Hydrochloride	
(Sertraline	
Hydrochloride)	C
Trazodone	
(Trazodone)	C
Hydroxyzine Embonate	
(Hydroxyzine	
Embonate)	C

Date:10/08/04ISR Number: 4472853-2Report Type:Expedited (15-DaCompany Report #2004UW20198
Age:34 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
25 MG DIALY		Dysphemia	Foreign	Seroquel	PS		ORAL
PO	2 MON	Speech Disorder	Health				
300 MG DAILY	2 MON		Professional	Gabapentin	SS		
			Other				

Date:10/08/04ISR Number: 4473826-6Report Type:Direct Company Report #CTU 229160
 Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 300 MG 1 X		Asthenia		Neurontin	PS		
Initial or Prolonged DAY		Chest Pain					
Other		Dizziness Dyspnoea Syncope Tremor					

Date:10/11/04ISR Number: 4471109-1Report Type:Expedited (15-DaCompany Report #PHRM2004FR02832
 Age:24 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cytolytic Hepatitis Drug Ineffective		Trileptal	PS	Novartis Sector: Pharma	ORAL
900 mg daily	185 DAY	Gamma-Glutamyltransferase		Depakine	SS		ORAL
750 mg, BID		Increased		Neurontin	SS		ORAL
600 mg, TID		Hepatitis Transaminases Increased					

Date:10/12/04ISR Number: 4472663-6Report Type:Direct
Age:59 YR Gender:Female I/FU:I

Company Report #CTU 229219

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Neurontin	PS		
3X DAY		Abnormal Dreams		Topomax (Samples Fr			
		Abnormal Sensation In Eye		Doctor)	SS		
3 X 11 -		Back Pain					
REDUCED TO 1		Confusional State					
		Coordination Abnormal					
		Dry Mouth					
		Ocular Hyperaemia					
		Oedema Peripheral					
		Rash					
		Speech Disorder					
		Thinking Abnormal					
		Vision Blurred					
		Visual Disturbance					
		Weight Increased					

Date:10/12/04ISR Number: 4472849-0Report Type:Direct
Age:48 YR Gender:Female I/FU:I

Company Report #CTU 229393

Outcome	PT
Other	Chest Pain
	Dyspnoea
	Fatigue

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Heart Rate Increased Hypersomnia	Report Source	Product	Role	Manufacturer	Route
300MG	TID			Neurontin 300mg Park-Davis	PS	Park-Davis	ORAL
ORAL							
100MG	TID			Neurontin 100mg Tid Park-Davis	SS	Park-Davis	ORAL
ORAL							

Date:10/12/04ISR Number: 4475241-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
Age:44 YR Gender:Male I/FU:F

Outcome
Hospitalization - PT
Initial or Prolonged Abdominal Pain Upper
Disability Abdominal Tenderness
Other Acrochordon
Actinic Keratosis
Acute Sinusitis
Albumin Globulin Ratio
Decreased
Alcoholism
Angina Unstable
Anxiety
Back Injury
Bipolar Disorder
Bladder Disorder
Blood Albumin Decreased
Blood Alkaline
Phosphatase Increased
Blood Calcium Decreased
Blood Triglycerides
Increased
Blood Urea
Nitrogen/Creatinine Ratio
Increased
Bone Cyst
Brain Neoplasm
Bronchitis Acute
Cardiac Arrest

Carotid Artery Disease
Cerebral Perfusion
Pressure Decreased
Chest Wall Mass
Cholecystitis
Cholelithiasis
Cognitive Disorder
Computerised Tomogram
Abnormal
Condyloma Acuminatum
Conversion Disorder
Convulsion
Coordination Abnormal
Cyst
Deafness Unilateral
Depression
Diabetes Mellitus
Difficulty In Walking
Diplopia
Dissociative Disorder

Freedom Of Information (FOI) Report

Disturbance In Attention
Dysphemia
Dysstasia
Ear Discomfort
Electrocardiogram P Wave
Abnormal
Emotional Disorder
Ependymoma
Exostosis
Exposure To Toxic Agent
Fall
Fibromyalgia
Folliculitis
Gait Disturbance
Gallbladder Disorder
Gamma-Glutamyltransferase
Increased
Gastritis
Gastrointestinal Disorder
Gastrooesophageal Reflux
Disease
Glioma
Gynaecomastia
Haematocrit Decreased
Hallucination
Headache
Hearing Impaired
Hepatic Enzyme Increased
Hepatic Steatosis
High Density Lipoprotein
Decreased
Hyperaemia
Hyperhidrosis
Impaired Driving Ability
Insomnia
Intervertebral Disc
Displacement
Intervertebral Disc
Protrusion
Joint Injury
Lower Gastrointestinal
Haemorrhage
Lumbar Spinal Stenosis
Marital Problem
Memory Impairment
Meniscus Lesion
Monocyte Count Decreased

Musculoskeletal Stiffness
Myocardial Infarction
Nervous System Disorder
Neurodegenerative
Disorder
Oedema Peripheral
Osteoarthritis
Pain
Paraesthesia
Paraneoplastic Syndrome
Patellofemoral Pain
Syndrome
Peptic Ulcer
Petit Mal Epilepsy

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
900 MG (1D), ORAL		Pharyngolaryngeal Pain Pigmented Naevus Plantar Fasciitis Post Procedural Complication Reading Disorder Road Traffic Accident Scar Seborrhoeic Keratosis Sick Sinus Syndrome Single Photon Emission Computerised Tomogram Abnormal Speech Disorder Stereotypy Strabismus Suicidal Ideation Synovitis Tension Tinnitus Toe Deformity Vision Blurred Visual Disturbance Weight Decreased	Consumer Health Professional Company Representative	Neurontin (Gabapentin) Valproate Sodium (Valproate Sodium) Buspirone Hydrochloride (Buspirone Hydrochloride) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Omeprazole (Omeprazole) Capsaicin (Capsaicin) Celecoxib (Celecoxib) Lansoprazole (Lansoprazole)	PS SS C C C C		ORAL

Date:10/12/04ISR Number: 4475312-6Report Type:Expedited (15-DaCompany Report #2004070920

Age: Gender:Female I/FU:I

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
1500 MG (300 MG), ORAL; 1500 MG (600 MG, QID INTERVAL:		PT Arthritis Burning Sensation Headache Transient Ischaemic Attack Weight Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL

50 MG (50 MG,

1 IN 1 D),

ORAL

Zoloft (Sertraline) SS

ORAL

Pravastatin Sodium (Pravastatin Sodium)	C
Telmisartin (Temisartan)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C
Omeprazole (Omeprazole)	C
Iron (Iron)	C
All Other Therapeutic Products (All Other Therapeutic Products)	C
Quinine (Quinine)	C
Temazepam (Temazepam)	C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/12/04ISR Number: 4475314-XReport Type:Expedited (15-DaCompany Report #2004070411

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Asthenia Benign Prostatic	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other 2 MG (2 MG, 1 IN 1 D), ORAL		Hyperplasia Chest Discomfort Chills Condition Aggravated Constipation Diarrhoea Difficulty In Walking Dizziness Dysuria Heart Rate Increased Malaise Medication Error Neuropathy Peripheral Somnolence Urinary Retention		Cardura (Doxazosin Mesilate) Triobe (Cyanocobalamin, Folic Acid, Pyridoxine) Acetylsalicylic Acid (Acetylsalicylic Acid)	SS C C		ORAL

Date:10/12/04ISR Number: 4475868-3Report Type:Expedited (15-DaCompany Report #2004070512

Age:92 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Medication Error Multiple Drug Overdose	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Clozapine (Clozapine)	SS		ORAL

Date:10/12/04ISR Number: 4475872-5Report Type:Expedited (15-DaCompany Report #2004070289

Age:38 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Adverse Event	Literature	Gabapentin	PS	ORAL
ORAL	Cardiac Arrest	Health	(Gabapentin)		
ORAL	Multiple Drug Overdose	Professional	Methadone	SS	ORAL
ORAL	Respiratory Arrest		(Methadone)		
ORAL			Fluoxetine	SS	ORAL
ORAL			(Fluoxetine)		
ORAL			All Other		
ORAL			Therapeutic Products		
ORAL			(All Other		
ORAL			Therapeutic	SS	ORAL
ORAL			Products)		

Date:10/12/04ISR Number: 4475938-XReport Type:Expedited (15-DaCompany Report #2004061440

Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Back Pain
Initial or Prolonged	Bone Neoplasm
Other	Burning Sensation
	Coma Hepatic
	Cyst
	Hallucination, Visual
	Neuralgia
	Neuropathy

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Professional				
1800 MG (600 MG, 3 IN 1 D), ORAL		Consumer Health	Neurontin (Gabapentin)	PS		ORAL
			Tiagabine Hydrochloride (Tiagabine Hydrochloride)	C		
			Cyclobenzaprine (Cyclobenzaprine)	C		
			Estrogens Conjugated (Estrogens Conjugated)	C		
			Zolpidem Tartrate (Zolpidem Tartrate)	C		
			Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
			Tizanidine (Tizanidine)	C		
			All Other Therapeutic Products	C		
			Seretide Mite (Fluticasone Propionate, Slameterol Xinafoate)	C		
			Salbutamol (Salbutamol)	C		
			Epinephrine (Epinephrine)	C		

Date:10/12/04ISR Number: 4475940-8Report Type:Expedited (15-DaCompany Report #2004058531
Age:69 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other	Convulsion	Health	Neurontin		
1600 MG (400	Hypoaesthesia	Professional	(Gabapentin)	PS	ORAL
MG, 4 IN 1	Intervertebral Disc				
D), ORAL	Protrusion				
100 MG, ORAL	Neuritis		Dilantin (Phenytoin		
	Pharmaceutical Product		Sodium)	SS	ORAL
	Complaint				

Date:10/12/04ISR Number: 4476014-2Report Type:Expedited (15-DaCompany Report #2004070292
Age:49 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Gabapentin			
ORAL		Intentional Misuse	Health	(Gabapentin)	PS		ORAL
ORAL			Professional	Methadone			
ORAL				(Methadone)	SS		ORAL
ORAL				Methocarbamol			
				(Methocarbamol)	SS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/12/04ISR Number: 4476034-8Report Type:Expedited (15-DaCompany Report #2004071970
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety Completed Suicide Drug Ineffective Pain Polytraumatism Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		

Date:10/12/04ISR Number: 4476066-XReport Type:Expedited (15-DaCompany Report #2004068235
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Surgery	Consumer	Neurontin (Gabapentin)	PS		

Date:10/12/04ISR Number: 4476085-3Report Type:Expedited (15-DaCompany Report #2004070288
Age:45 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Multiple Drug Overdose	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		ORAL

Date:10/12/04ISR Number: 4476482-6Report Type:Expedited (15-DaCompany Report #2004070525
Age:31 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest Drug Abuser Respiratory Arrest	Literature Health Professional	Gabapentin (Gabapentin) Cocaine (Cocaine)	PS SS		

Ethanol (Ethanol) SS
All Otehr
Therapeutic
Products) SS

Date:10/12/04ISR Number: 4476484-XReport Type:Expedited (15-DaCompany Report #2004070485
Age:19 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Baclofen (Baclofen)	SS		ORAL
ORAL				Oxybutynin (Oxybutynin)	SS		ORAL

Date:10/12/04ISR Number: 4476485-1Report Type:Expedited (15-DaCompany Report #2004070295
Age:53 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Levothyroxine			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL				(Levothyroxine)	SS		ORAL
				All Other Therapeutic Products (All Other Therapeutic Products)	SS		ORAL
ORAL				Macrogol (Macrogol)	C		

Date:10/12/04ISR Number: 4476489-9Report Type:Expedited (15-DaCompany Report #2004070501
Age:50 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Multiple Drug Overdose	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Chlorpromazine (Chlorpromazine)	SS		ORAL
ORAL				Quetiapine (Quetiapine)	SS		ORAL
ORAL				All Other Therapeutic Products (All Other Therapeutic Products)	SS		ORAL

Date:10/12/04ISR Number: 4476491-7Report Type:Expedited (15-DaCompany Report #2004070297
Age:44 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Multiple Drug Overdose	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Valproic Acid (Valproic Acid)	SS		ORAL
ORAL				Levothyroxine			

ORAL				(Levothyroxine)	SS		ORAL
				All Other Therapeutic Products (All Other Therapeutic Products)	SS		ORAL

Date:10/12/04ISR Number: 4476495-4Report Type:Expedited (15-DaCompany Report #2004070286
 Age:23 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Intentional Misuse	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		ORAL

Date:10/12/04ISR Number: 4477559-1Report Type:Expedited (15-DaCompany Report #2004070936
 Age:20 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Abuser	Literature Health	Sertraline (Sertraline)	PS		
			Professional	Methadone (Methadone)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Gabapentin
 (Gabapentin) SS
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C

Date:10/12/04ISR Number: 4477563-3Report Type:Expedited (15-DaCompany Report #2004070293
 Age:35 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Tramadol (Tramadol)	SS		ORAL
ORAL				Citalopram (Citalopram)	SS		ORAL
ORAL				All Other Therapeutic Product	SS		ORAL

Date:10/13/04ISR Number: 4475313-8Report Type:Direct Company Report #CTU 229534
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abasia Drug Ineffective Pain In Extremity Pharmaceutical Product Complaint		Gabapentin	PS		

Date:10/13/04ISR Number: 4476533-9Report Type:Expedited (15-DaCompany Report #2004010780
 Age:69 YR Gender:Male I/FU:F

Outcome	PT
Disability	Actinic Keratosis

Other

Aldolase Increased
Anhedonia
Arthralgia
Asthenia
Blood Creatinine
Increased
Blood Glucose Increased
Blood Potassium Increased
Blood Sodium Increased
Blood Urine Present
Bundle Branch Block Left
Difficulty In Walking
Diverticulum
Drug Hypersensitivity
Emotional Disorder
Fatigue
Feeling Cold
Flushing
Head Injury
Hyperhidrosis
Insomnia
Joint Crepitation
Joint Range Of Motion

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Decreased Kyphosis Low Density Lipoprotein Increased					
10 MG (DAILY)		Mental Disorder Muscle Atrophy	Consumer	Lipitor (Atorvastatin)	PS		
(DAILY), ORAL		Muscle Spasms Muscular Weakness Myalgia Myopathy		Hyzaar (Hydrochlorothiazide , Losartan Potassium)	SS		ORAL
300 MG ,ORAL		Neck Injury Nervousness		Neurontin (Gabapentin)	SS		ORAL
		Neuralgia Neuropathy Pain Pain In Extremity Polymyositis Polytraumatism Prostatic Specific Antigen Increased Protein Urine Present Rectal Polyp Spinal Column Stenosis Tenderness Visual Acuity Reduced Weight Decreased		Influenza Vaccine (Influenza Vaccine) All Other Therapeutic Products (All Other Therapeutic Products) Vicodin (Hydrocodone Bitartrate, Paracetamol) Acetylsalicylic Acid (Acetylsalicylic Acid) Amlodipine Besilate (Amlodipine Besilate)	C C C C		

Date:10/13/04ISR Number: 4476535-2Report Type:Expedited (15-DaCompany Report #2004070411

Age: Gender:Male I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hospitalization - Initial or Prolonged ORAL	Consumer	Neurontin (Gabapentin)	PS		ORAL
2 MG (2 MG, 1 IN 1 D), ORAL		Benign Prostatic Hyperplasia Chest Discomfort		Cardura (Doxazosin Mesilate)	SS		ORAL

Chills
 Condition Aggravated
 Constipation
 Diarrhoea
 Dizziness
 Drug Ineffective
 Dysuria
 Heart Rate Increased
 Malaise
 Medication Error
 Neuropathy Peripheral
 Somnolence

Triobe
 (Cyanocobalamin,
 Folic Acid,
 Pyridoxine) C
 Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C

Date:10/13/04ISR Number: 4476539-XReport Type:Expedited (15-DaCompany Report #2004074577

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 Other MG, 2 IN 1 D)		Hepatic Encephalopathy	Health Professional	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/13/04ISR Number: 4476594-7Report Type:Expedited (15-DaCompany Report #HQWYE542005OCT04
Age:49 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Literature	Methocarbamol (Methocarbamol, Tablet, 500 Mg)	PS		ORAL
ORAL				Gabapentin (Gabapentin,)	SS		ORAL
ORAL				Methadone (Methadone,)	SS		ORAL

Date:10/14/04ISR Number: 4477630-4Report Type:Expedited (15-DaCompany Report #2004051682
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	900 MG (300 MG, 3 IN 1 D), ORAL	Difficulty In Walking Disease Progression Myelopathy Oedema Peripheral	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Vioxx (Rofecoxib)	SS		ORAL
				Fexofenadine Hydrochloride (Fexofenadine Hydrochloride)	C		
				Fluticasone Propionate (Fluticasone Propionate)	C		
				Atorvastatin (Atorvastatin)	C		
				Gezor (Hydrochlorothiazide , Quinapril Hydrochloride)	C		
				Propranolol			

Hydrochloride
(Propranolol Hydrochloride) C
Baclofen (Baclofen) C
Sertraline Hydrochloride
(Sertraline Hydrochloride) C
Alprazolam (Alprazolam) C
Potassium Chloride (Potassium Chloride) C
Furosemide (Furosemide) C

Date:10/14/04ISR Number: 4477634-1Report Type:Expedited (15-DaCompany Report #2004061440
Age:47 YR Gender:Female I/FU:F

Outcome	PT
Life-Threatening	Back Pain
Hospitalization -	Coma
Initial or Prolonged	Feeling Hot
Other	Hallucination, Visual
	Hepatic Function Abnormal

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
	Intraneural Cyst Mental Status Changes Neoplasm					
1800 MG (600 MG, 3 IN 1 1 D), ORAL	Neuralgia Night Sweats Persecutory Delusion Post Procedural Pain Suicidal Ideation	Consumer Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Tiagabine Hydrochloride (Tiagabine Hydrochloride)	C		
			Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride)	C		
			Estrogens Conjugated (Estrogens Conjugated)	C		
			Zolpidem Tartrate (Zolpidem Tartrate)	C		
			Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
			Tizanidine (Tizanidine)	C		
			Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)	C		
			Salbutamol (Salbutamol)	C		
			Epinephrine (Epinephrine)	C		
			Prednisone (Prednisone)	C		
			Montelukast Sodium (Montelukast Sodium)	C		
			Lorazepam (Lorazepam)	C		

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Consumer	Neurontin			
(600 MG)		Knee Arthroplasty		(Gabapentin)	PS		
		Osteoporosis		Fentanyl (Fentanyl)	C		
				Ibuprofen			
				(Ibuprofen)	C		
				Rofecoxib			
				(Rofecoxib)	C		
				Helianthus Tuberosus			
				(Helianthus			
				Tuberosus	C		
				All Othr Tehrapeutic			
				Products (All Other			
				Therapeutic Product)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/14/04ISR Number: 4477648-1Report Type:Expedited (15-DaCompany Report #2004075239

Age:71 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Chest Pain	Consumer	Neurontin			
Initial or Prolonged	Coma		(Gabapentin)	PS		ORAL
ORAL						
Other	Malaise		Pantoprazole			
	Nausea		(Pantoprazole)	C		
	Renal Failure		Magnesium			
			(Magnesium)	C		
			Oxycocet (Oxycodone			
			Hydrochloride,			
			Paracetamol)	C		
			Clopidogrel Sulfate			
			(Clopidogrel			
			Sulfate)	C		
			Losartan Potassium			
			(Losartan Potassium)	C		
			Fursoemide			
			(Furosemide)	C		
			Isosorbide			
			(Isorbide)	C		
			Potassium			
			(Potassium)	C		
			Simvastatin			
			(Simvastatin)	C		
			Escitalopram			
			(Escitalopram)	C		
			Acetyl Salicylic			
			Acid			
			(Acetylsalicylic			
			Acid)	C		
			Diphenhydramine			
			Hydrochloride			
			(Diphenhydramine			
			Hydrochloride)	C		
			All Otehr			
			Therapeutic Products			
			(All Other			
			Therapeutic			
			Products)	C		
			Methylphenidate			
			Hydrochloride			
			(Methylphenidate			
			Hydrochloride)	C		

Seretide Mite
 (Fluticasone
 Propionate,
 Salmeterol
 Xinafoate) C
 Oxygen (Oxygen) C
 Fentanyl (Fentanyl) C

Date:10/14/04ISR Number: 4477770-XReport Type:Expedited (15-DaCompany Report #2004073296
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 160 MG		Cognitive Deterioration	Health	Geodon (Ziprasidone)	PS		
		Pain Paraesthesia Speech Disorder	Professional Company Representative	Neurontin (Gabapentin) Quetiapine Fumarate	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

100 MG

(Quetiapine
Fumarate) SS

Cannabis (Cannabis) SS

Date:10/14/04ISR Number: 4478881-5Report Type:Expedited (15-DaCompany Report #2004075359
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (400 MG, 3 IN 1 D)		Hepatotoxicity	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Paroxetine (Paroxetine) Temozolomide (Temozolomide)	PS SS C		

Date:10/15/04ISR Number: 4479300-5Report Type:Expedited (15-DaCompany Report #2004074993
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4504970-2Report Type:Periodic Company Report #2003125334
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1200 MG (600, BID), ORAL		Feeling Abnormal Suicidal Ideation	Consumer	Neurontin (Gabapentin) Mirtazapine (Mirtazapine) Trazodone (Trazodone)	PS C C		ORAL

Olanzapine
(Olanzapine) C

Date:10/15/04ISR Number: 4504971-4Report Type:Periodic Company Report #2003125900
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4504972-6Report Type:Periodic Company Report #2003125902
Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain Impaired Driving Ability	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (TID), ORAL		Visual Disturbance Weight Increased		Vicodin (Paracetamol, Hydrocodone Bitartrate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4504973-8Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2003125905

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intentional Misuse Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		
2400 MG (TID)		Vomiting		All Other Therapeutic Products (All Other Therapeutic Products)	SS		
				Trazodone (Trazodone)	C		
				Lansoprazole (Lansoprazole)	C		

Date:10/15/04ISR Number: 4504974-XReport Type:Periodic
Age:39 YR Gender:Male I/FU:I

Company Report #2003125937

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression Somnolence	Consumer	Neurontin (Gabapentin)	PS		
900 MG		Suicidal Ideation		Modafinil (Modafinil)	C		
				Vitamins (Vitamins)	C		

Date:10/15/04ISR Number: 4504975-1Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2003125941

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Sensation In Eye Blood Glucose Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
(TID), ORAL		Blood Pressure Increased		Prednisone (Prednisone)	SS		
		Convulsion		Lipitor (Atorvastatin)	C		
		Musculoskeletal Stiffness Pain		Antivirals For			
		Tachycardia					

Tremor

Systemic Use (Antivirals For Systemic Use)	C
Metformin Hydrochloride (Metformin Hydrochloride)	C
Vicodin (Paracetamol, Hydrocodone Bitartrate)	C
Valsartan (Valsartan)	C
Paracetamol (Paracetamol)	C

Date:10/15/04ISR Number: 4504977-5Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2003126925

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
		Depression Mental Impairment Personality Change Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4504978-7Report Type:Periodic Company Report #2004000179
Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Face Oedema	Health Professional	Neurontin (Gabapentin)	PS		ORAL

900 MG (TID)

ORAL

Antihypertensives
(Antihypertensives) C

Date:10/15/04ISR Number: 4504979-9Report Type:Periodic Company Report #2004000879
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Crying Depression	Consumer Health Professional	Neurontin (Gabapentin)	PS		
Other			Drug Ineffective Nerve Injury Suicide Attempt		Hydrocodone (Hydrocodone) Carisoprodol (Carisoprodol)	C C		

Date:10/15/04ISR Number: 4504980-5Report Type:Periodic Company Report #2004001003
Age:40 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Abdominal Pain	Consumer	Neurontin			

600 MG (BID)	Convulsion	(Gabapentin)	PS	ORAL
ORAL	Deafness			
	Dizziness	Clonazepam		
	Dyspepsia	(Clonazepam)	C	
	Ear Disorder			
	Intentional Misuse			
	Medication Error			
	Neuralgia			
	Sedation			
	Vestibular Disorder			

Date:10/15/04ISR Number: 4504981-7Report Type:Periodic Company Report #2003118176
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Facial Palsy	Consumer	Neurontin			
		Feeling Abnormal		(Gabapentin)	PS		
		Pain					
		Stomach Discomfort					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4504982-9Report Type:Periodic
 Age:65 YR Gender:Female I/FU:I

Company Report #2003118178

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Depression Intentional Self-Injury	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1800 MG (TID)							

ORAL

Dilantin (Phenytoin Sodium)	C
Zoloft (Sertraline)	C
Atenolol	C
Ranitidine Hydrochloride	C
Valerian Root (Valeriana Officinalis Root)	C
Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Metamucil "Procter & Gamble" (Glucose Monohydrate, Ispaghula Husk)	C

Date:10/15/04ISR Number: 4504983-0Report Type:Periodic
 Age:50 YR Gender:Female I/FU:I

Company Report #2003118686

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Loss Of Consciousness	Consumer Company	Neurontin (Gabapentin)	PS		ORAL

ORAL

Representative

Date:10/15/04ISR Number: 4504984-2Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #2003118691

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Loss Of Consciousness	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1800 MG TID			Company Representative				

Date:10/15/04ISR Number: 4504985-4Report Type:Periodic Company Report #2003119168
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Intentional Misuse Medication Error	Health Professional	Neurontin (Gabapentin)	PS		ORAL
3900 MG DAILY		Non-Hodgkin'S Lymphoma	Company Representative				

Date:10/15/04ISR Number: 4504986-6Report Type:Periodic Company Report #2003120555
Age:74 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Abasia Abnormal Behaviour

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Asthenia Atrial Fibrillation Confusional State	Report Source				
600 MG (BID)		Delusion Dysarthria	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Insomnia					
		Intentional Misuse		Zoloft (Sertraline)	C		
		Irritability		Celebrex (Celecoxib)	C		
		Lethargy		Atenolol	C		
		Medication Error		Amiodarone			
		Nausea		Hydrochloride	C		
		Oedema Peripheral		Amitriptyline			
		Oral Intake Reduced		Hydrochloride	C		
		Parosmia		Benazepril			
		Somnolence		Hydrochloride	C		
		Tremor		Fentanyl	C		
		Vomiting		Lidocaine	C		
				Fexofenadine			
				Hydrochloride	C		
				Magnesium	C		
				Montelukast Sodium	C		
				Esomeprazole	C		
				Raloxifene			
				Hydrochloride	C		
				Warfarin Sodium	C		
				Calcium	C		
				Tocopherol	C		
				Estradiol	C		
				Hydrochlorothiazide	C		
				Meclozine	C		
				Simvastatin	C		
				Centrum (Vitamins Nos, Minerals Nos)	C		
				Morphine	C		

Date:10/15/04ISR Number: 4504987-8Report Type:Periodic
Age:40 YR Gender:Female I/FU:I

Company Report #2003121503

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion	Health	Neurontin			

400 MG (200 Drug Level Decreased Professional (Gabapentin) PS ORAL
BID) ORAL

Date:10/15/04ISR Number: 4504988-XReport Type:Periodic Company Report #2003123205
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4504990-8Report Type:Periodic Company Report #2003124351
Age:12 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Stevens-Johnson Syndrome	Health Professional	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4504991-XReport Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2003124522

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG TID		Aggression Agitation	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other ORAL		Amnesia	Professional				
		Depression Insomnia Intentional Misuse Suicide Attempt		Paroxetine Hydrochloride Venlafaxine Hydrochloride	C C		

Date:10/15/04ISR Number: 4504992-1Report Type:Periodic
 Age:57 YR Gender:Female I/FU:I

Company Report #2003114281

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blindness	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4504994-5Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #2003114318

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 3200 OR 3800 Initial or Prolonged MG TID		Intentional Misuse Medication Error	Consumer	Neurontin (Gabapentin)	PS		
Other		Negative Thoughts Suicidal Ideation Suicide Attempt					

Date:10/15/04ISR Number: 4504996-9Report Type:Periodic
 Age:66 YR Gender:Male I/FU:I

Company Report #2003115407

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Asthenia	Consumer	Neurontin		
	Fatigue		(Gabapentin)	PS	ORAL
900 MG TID					
	Influenza Like Illness				
ORAL					
	Loss Of Consciousness		Phenytoin Sodium	C	
	Visual Disturbance		Losartan Potassium	C	
			Esomeprazole	C	
			Acetylsalicylic Acid	C	
			Finasteride	C	
			Cyclobenzaprine	C	

Date:10/15/04ISR Number: 4504997-0Report Type:Periodic Company Report #2003115804
Age:52 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Suicide Attempt	Health	Neurontin			
Initial or Prolonged		Professional	(Gabapentin)	PS		
900 MG TID						
Other			Escitalopram			
			(Escitalopram)	SS		
10 MG DAILY						
			Oxazepam (Oxazepam)	SS		
30 MG TID						
			Ethanol (Ethanol)	SS		
			Nadolol	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4504998-2Report Type:Periodic
Age:56 YR Gender:Male I/FU:I

Company Report #2003116123

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams	Health	Neurontin			
		Chest Pain	Professional	(Gabapentin)	PS		ORAL
		Nightmare					
		Suicidal Ideation		Fentanyl	C		
				Atenolol	C		
				Diltiazem	C		

Date:10/15/04ISR Number: 4505000-9Report Type:Periodic
Age:26 YR Gender:Male I/FU:I

Company Report #2003116318

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain	Consumer	Neurontin			
		Abdominal Pain Upper	Health	(Gabapentin)	PS		ORAL
		Adverse Drug Reaction	Professional				
		Blood Pressure Increased		Clonazepam	C		
		Family Stress					
		Fluid Retention					
		Haemorrhoids					
		Mood Swings					
		Night Sweats					
		Pain					
		Personality Change					
		Suicidal Ideation					

Date:10/15/04ISR Number: 4505003-4Report Type:Periodic
Age:45 YR Gender:Female I/FU:I

Company Report #2003116566

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
		Headache		(Gabapentin)	PS		ORAL
		Loss Of Consciousness		Citalopram			

Hydrobromide

C

Date:10/15/04ISR Number: 4505004-6Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2003116657

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetic Neuropathy	Consumer	Neurontin			
		Diarrhoea	Health	(Gabapentin)	PS		ORAL
900 MG TID		Dizziness	Professional				
ORAL		Nausea		Glipizide	C		
		Stool Analysis Abnormal		Levothyroxine Sodium	C		
				Pravastatin Sodium	C		
				Atenolol	C		
				Enalapril	C		
				Hydrochlorothiazide	C		
				Metformin			
				Hydrochloride	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505005-8Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2003117026

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Overdose	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:10/15/04ISR Number: 4505007-1Report Type:Periodic
 Age:30 YR Gender:Female I/FU:I

Company Report #2003117626

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 100 MG DAILY		Abdominal Pain Asthenia	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Blood Bilirubin Decreased					
ORAL		Blood Thyroid Stimulating Hormone Increased Constipation Diarrhoea Dyspepsia Hepatic Enzyme Increased Lethargy Tongue Discolouration		Escitalopram Levothyroxine Sodium Prednisone	C C C		

Date:10/15/04ISR Number: 4505009-5Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2003041594

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatine Phosphokinase Increased	Health Professional	Neurontin (Gabapentin)	PS		
1800 MG (TID)		Convulsion Intentional Misuse Medication Error Myalgia					

Date:10/15/04ISR Number: 4505010-1Report Type:Periodic Company Report #2003041702
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cataract	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505011-3Report Type:Periodic Company Report #2003110010
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Suicidal Ideation	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1800 MG							
(TID), ORAL							

Date:10/15/04ISR Number: 4505012-5Report Type:Periodic Company Report #2003111954
Age:15 YR Gender:Female I/FU:I

Outcome Other	PT Convulsion Drug Withdrawal Syndrome
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Eating Disorder Eye Pain Feeling Abnormal	3600 MG (DAILY)			Consumer	Neurontin (Gabapentin)	PS		
Headache Intentional Misuse Medication Error								
Nausea Sinusitis Weight Decreased					Topiramate (Topiramate) Clorazepate Dipotassium (Clorazepate Dipotassium)	SS C		

Date:10/15/04ISR Number: 4505014-9Report Type:Periodic Company Report #2003112721
Age: Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Blood Pressure Decreased Hypertension	600 MG (DAILY), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
					Hydrochlorothiazide (Hydrochlorothiazide) Nadolol (Nadolol) Atorvastatin (Atorvastatin)	C C C		

Date:10/15/04ISR Number: 4505015-0Report Type:Periodic Company Report #2003112984
Age:39 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Abdominal Pain Upper Depression	ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
Dysphemia Gastroesophageal Reflux					Methocarbamol (Methocarbamol)	SS		

Disease
 Hypersomnia
 Mental Disorder
 Muscle Spasms
 Speech Disorder
 Suicide Attempt
 Tic

Date:10/15/04ISR Number: 4505017-4Report Type:Periodic
 Age:76 YR Gender:Male I/FU:I

Company Report #2003113288

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Buttock Pain	Health	Neurontin			
Other		Dizziness	Professional	(Gabapentin)	PS		ORAL
300 MG		Ejaculation Disorder					
(DAILY), ORAL		Hypoaesthesia		Trimipramine			
		Intentional Misuse		(Trimipramine)	C		
		Medication Error		Acetylsalicylic Acid			
		Myocardial Infarction		(Acetylsalicylic			
		Somnolence		Acid)	C		
		Urinary Incontinence		All Other			
		Visual Disturbance		Therapeutic Products			
				(All Other			

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Freedom Of Information (FOI) Report

Therapeutic Products)	C
Furosemide (Furosemide)	C
Carvedilol (Carvedilol)	C
Captopril (Captopril)	C
All Other Therapeutic Products (All Other Therapeutic Products)	C
Ibuprofen (Ibuprofen)	C
Paracetamol (Paracetamol)	C
Arthrotec (Diclofenac Sodium, Misoprostol)	C
Dextropropoxyphene Hydrochloride (Dextropropoxyphene Hydrochloride)	C
Theragran (Vitamins Nos)	C
Thiamine Hydrochloride (Thiamine Hydrochloride)	C

Date:10/15/04ISR Number: 4505018-6Report Type:Periodic Company Report #2003113617
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505020-4Report Type:Periodic Company Report #2003113623
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -	Loss Of Consciousness	Consumer	Neurontin	
Initial or Prolonged	Urinary Incontinence		(Gabapentin)	PS
Other	Vomiting			

Date:10/15/04ISR Number: 4505021-6Report Type:Periodic Company Report #2003113767
 Age:58 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Amnesia	Consumer	Neurontin			
Hospitalization -	Suicide Attempt		(Gabapentin)	PS		ORAL
ORAL						
Initial or Prolonged			Antidepressants			
Other			(Antidepressants)	C		
			All Other			
			Therapeutic Products			
			(All Other			
			Therapeutic			
			Products)	C		

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Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505022-8Report Type:Periodic Company Report #2003035281
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cataract	Health	Neurontin			
		Depression	Professional	(Gabapentin)	PS		ORAL
		Vision Blurred					
				Escitalopram			
				(Escitalopram)	SS		
				Methylphenidate			
				Hydrochloride	C		
				Oral Contraceptives			
				Nos	C		

Date:10/15/04ISR Number: 4505023-XReport Type:Periodic Company Report #2003035548
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pulmonary Oedema	Health	Neurontin			
			Professional	(Gabapentin)	PS		
			Company				
			Representative				

Date:10/15/04ISR Number: 4505026-5Report Type:Periodic Company Report #2003035549
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pulmonary Oedema	Health	Neurontin			
			Professional	(Gabapentin)	PS		
			Company				
			Representative				

Date:10/15/04ISR Number: 4505028-9Report Type:Periodic Company Report #2003036398
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cardiac Failure Congestive	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG (100MG TID) ORAL		Difficulty In Walking					
				Spironolactone	C		
				Alendronate Sodium	C		
				Calcium	C		
				Paroxetine			
				Hydrochloride	C		
				Mirtazapine	C		
				Tocopherol	C		
				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrochloride, Folic Acid	C C		
				Vicodin (Paracetamol, Hydrocodone Bitartrate)	C		
				Celecoxib	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505029-0Report Type:Periodic
Age: Gender: I/FU:I

Company Report #2003037649

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intentional Misuse Medication Error	Health Professional	Neurontin (Gabapentin)	PS		
3600 MG QID		Psychotic Disorder		All Other Therapeutic Products (All Other Therapeutic Products)	SS		

Date:10/15/04ISR Number: 4505031-9Report Type:Periodic
Age: Gender: I/FU:I

Company Report #2003037886

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Psychotic Disorder	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505032-0Report Type:Periodic
Age:51 YR Gender:Female I/FU:I

Company Report #2003039268

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain Amnesia	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300MG (TWICE DAILY) ORAL		Coordination Abnormal		Nortriptyline Vicodin (Paracetamol, Hydrocodone Bitartrate)	C C		
		Diarrhoea Dizziness Dysarthria Dysphagia Eructation Flatulence Gastritis Haematochezia Nausea Neuropathic Pain Rectal Haemorrhage					

Somnolence
Vertigo

Date:10/15/04ISR Number: 4505033-2Report Type:Periodic Company Report #2003039651
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysphagia Nausea	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG TID OR		Oropharyngeal Swelling					
QID ORAL		Pain Swelling Face Vomiting		Antihypertensives	C		

Date:10/15/04ISR Number: 4505034-4Report Type:Periodic Company Report #2003040738
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Depression Mania	Consumer	Neurontin (Gabapentin) Aripiprazole	PS		

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Freedom Of Information (FOI) Report

(Aripiprazole) SS

Date:10/15/04ISR Number: 4505035-6Report Type:Periodic Company Report #2003041088
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Pericardial Effusion Pericarditis	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505038-1Report Type:Periodic Company Report #2003021325
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG (TID) ORAL		Asthenia Convulsion Fall	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Feeling Abnormal Head Discomfort Headache Hypersomnia Musculoskeletal Stiffness Nausea Nervousness Pain Paraesthesia Tremor		Propacet (Paracetamol, Dextropropoxyphene) Lorazepam Doxepin	C C C		

Date:10/15/04ISR Number: 4505040-XReport Type:Periodic Company Report #2003022165
 Age:54 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 200 MG DAILY		Abdominal Pain Upper Blood Pressure Decreased Chest Pain	Health Professional	Neurontin (Gabapentin) Topiramate	PS		

Dysstasia	(Topiramate)	SS
Ear Disorder	Nortriptyline	
Euphoric Mood	(Nortriptyline)	SS
Fatigue	Oxcarbazepine	
Feeling Abnormal	(Oxcarbazepine)	SS
Hepatic Enzyme Increased	Tiagabine	
Hepatitis	Hydrochloride	
Hepatomegaly	(Tiagabine	
Infectious Mononucleosis	Hydrochloride)	SS
Logorrhoea	Carbamazepine	
Sleep Disorder	(Carbamazepine)	SS
Syncope	Ascorbic Acid	C
Trigeminal Neuralgia	Tocopherol	C
Vertigo	Multivitamin "Lappe"	
	(Vitamins Nos)	C
	Glucosamine	
	W/Chondroitin	
	Sulfates (Ascorbic	
	Acid Manganese,	
	Chondroitin Sulfte,	C
	All Other	
	Therapeutic Products	C

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Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505041-1Report Type:Periodic
Age: Gender:Male I/FU:F

Company Report #2003026319

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 5400 (TID) Other		Drug Level Decreased Drug Withdrawal Syndrome Medication Error Overdose Tremor	Health Professional Company Representative	Neurontin (Gabapentin) Methadone	PS C		

Date:10/15/04ISR Number: 4505042-3Report Type:Periodic
Age:73 YR Gender:Male I/FU:F

Company Report #2003032864

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (300 MD) TID ORAL		Constipation Coordination Abnormal Cough Deafness Unilateral Disturbance In Attention Dizziness Dry Mouth Fatigue Feeling Abnormal Headache Lacrimation Increased Listless Mood Altered Pain Pharyngolaryngeal Pain Pyrexia Somnolence Vision Blurred Visual Disturbance	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:10/15/04ISR Number: 4505044-7Report Type:Periodic
Age:66 YR Gender:Male I/FU:F

Company Report #2003033496

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation	Consumer	Neurontin			
		Asthenia		(Gabapentin)	PS		ORAL
ORAL		Balance Disorder		Antihypertensives	C		
		Disorientation		Cholesterol And			
		Disturbance In Attention		Triglyceride			
		Dizziness		Reducers	C		
		Dysphagia		Alprazolam	C		
		Dyspnoea		Analgesics	C		
		Illogical Thinking		Atenolol	C		
		Insomnia		Captopril	C		
		Irritability		Metformin			
		Mood Altered		Hydrochloride	C		
		Rash					
		Restlessness					
		Somnolence					
		Speech Disorder					
		Vision Blurred					

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Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505050-2Report Type:Periodic
Age:61 YR Gender:Male I/FU:I

Company Report #2004052225

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Memory Impairment					
		Overweight		Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		
		Repetitive Speech		Co-Diovan (Hydrochlorothiazide , Valsartan)	C		
		Vomiting		Vitamins	C		

Date:10/15/04ISR Number: 4505053-8Report Type:Periodic
Age: Gender: I/FU:I

Company Report #2004052385

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicidal Ideation	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505056-3Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2004052393

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident	Consumer	Neurontin (Gabapentin)	PS		
		Diabetes Mellitus		Morphine (Morphine)	SS		
				Clonazepam (Clonazepam)	SS		
				Oxycocet (Oxycodone Hydrochloride, Paracetamol)	SS		
				All Other Therapeutic Products (All Other Therapeutic			

Date:10/15/04ISR Number: 4505059-9Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2004052502

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder	Consumer	Neurontin			
		Body Height Decreased		(Gabapentin)	PS		ORAL
300 MG (300		Breast Cancer					
MG 1 IN 1 D)		Dizziness					
ORAL		Dyskinesia		Oxcarbazepine			
		Gastric Disorder		(Oxcarazepine)	SS		
		Pain		Acetylsalicylic Acid	C		
		Weight Decreased		Lorazepam	C		

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Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505061-7Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #2004054070

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State Dysarthria	Consumer	Neurontin (Gabapentin)	PS		
	1800 MG (1 D)	Hepatitis Hypervigilance Speech Disorder Unevaluable Event		Levetiracetam	C		

Date:10/15/04ISR Number: 4505064-2Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #2004054424

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation Depression Difficulty In Walking Disturbance In Attention Dizziness Fatigue Joint Injury Limb Discomfort Mood Altered Pain Rash Pruritic Somnolence Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505065-4Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2004056047

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia Difficulty In Walking	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL				Glipizide	C		
				Atenolol	C		
				Simvastatin	C		
				Acetylsalicylic Acid	C		
				Loratadine	C		
				Metoprolol Tartrate	C		

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Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505070-8Report Type:Periodic
Age:65 YR Gender:Male I/FU:I

Company Report #2004059018

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Visual Field Defect	Consumer	Neurontin			
ORAL			Company	(Gabapentin)	PS		ORAL
			Representative	Antineoplastic			
				Agents	C		
				Vicodin (Hydrocodone			
				Bitartrate,			
				Paracetamol)	C		
				Lisinopril	C		
				Atorvastatin	C		

Date:10/15/04ISR Number: 4505201-XReport Type:Periodic
Age:64 YR Gender:Female I/FU:I

Company Report #2004046515

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nausea	Health	Neurontin			
300 MG (300		Suicidal Ideation	Professional	(Gabapentin)	PS		ORAL
MG, QD							
INTERVAL:							
EVERY DAY),							
ORAL							
				Amlodipine Beslate			
				(Amlodipine			
				Besilate)	C		
				Hydrochlorothiazide			
				(Hydrochlorothiazide			
)	C		
				Atenolol (Atenolol)	C		
				Trazodone			
				(Trazodone)	C		

Date:10/15/04ISR Number: 4505204-5Report Type:Periodic Company Report #2004046599
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diplopia Eye Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL		Vision Blurred		Warfarin Sodium (Warfarin Sodium)	C		

Date:10/15/04ISR Number: 4505206-9Report Type:Periodic Company Report #2004047166
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505207-0Report Type:Periodic Company Report #2004047490
Age:54 YR Gender:Female I/FU:I

Outcome	PT
Other	Arthritis Depression

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Myocardial Infarction
Somnolence

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
200 MG (100 MG, 1 IN 1 D), ORAL		Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
			All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:10/15/04ISR Number: 4505210-0Report Type:Periodic Company Report #2004047500
Age:18 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1 D), ORAL		Suicidal Ideation Tremor	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		
				Olanzapine (Olanzapine)	C		

Date:10/15/04ISR Number: 4505212-4Report Type:Periodic Company Report #2004047729
Age: Gender:Unknown I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Nephrolithiasis	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505215-XReport Type:Periodic Company Report #2004047833
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Failure Congestive	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505216-1Report Type:Periodic Company Report #2004048064
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation Suicidal Ideation	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505217-3Report Type:Periodic
Age:59 YR Gender:Female I/FU:I

Company Report #2004048448

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Constipation Diarrhoea	Consumer	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL		Faeces Discoloured Haematochezia					
ORAL		Somnolence		Diclofenac Sodium (Diclofenac Sodium)	SS		ORAL
ORAL				Cyanocobalamin (Cyanocobalamin)	SS		ORAL

Date:10/15/04ISR Number: 4505219-7Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2004051575

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505220-3Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2004039301

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Depression Muscle Twitching Suicidal Ideation	Health Professional	Neurontin (Gabapentin)	PS		
				Oxycodone (Oxycodone)	C		
				Atenolol (Atenolol)	C		
				Spiroinolactone (Spiroinolactone)	C		
				Temazepam (Temazepam)	C		
				Melatonin (Melatonin)	C		

Furosemide
(Furosemide) C
Fluoxetine
Hydrochloride
(Fluoxetine
Hydrochloride) C

Date:10/15/04ISR Number: 4505222-7Report Type:Periodic
Age:42 YR Gender:Female I/FU:I

Company Report #2004039616

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Feeling Abnormal Medication Error	Consumer	Neurontin (Gabapentin)	PS		ORAL
SEE IMAGE		Pain Paraesthesia Sedation Vision Blurred		Norethisterone (Norethisterone)	C		

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Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505224-0Report Type:Periodic
Age:79 YR Gender:Female I/FU:I

Company Report #2004040062

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Gingivitis	Health	Neurontin			
		Pain	Professional	(Gabapentin)	PS		ORAL
ORAL							

Date:10/15/04ISR Number: 4505227-6Report Type:Periodic
Age: Gender: I/FU:I

Company Report #2004040379

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident	Health	Neurontin			
			Professional	(Gabapentin)	PS		
			Company				
			Representative				

Date:10/15/04ISR Number: 4505228-8Report Type:Periodic
Age:16 YR Gender:Male I/FU:I

Company Report #2004041577

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Aggression	Consumer	Neurontin			
Initial or Prolonged		Agitation		(Gabapentin)	PS		

Date:10/15/04ISR Number: 4505231-8Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2004042631

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
				(Gabapentin)	PS		
				Celecoxib			
				(Celecoxib)	C		
				Vicodin (Hydrocodone			
				Bitartrate			
				Paracetamol)	C		
				Amitriptyline			
				(Amitriptyline			

(Amitriptyline) C
Cyclobenzaprine
(Cyclobenzaprine) C

Date:10/15/04ISR Number: 4505232-XReport Type:Periodic
Age:76 YR Gender:Male I/FU:I

Company Report #2004045199

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haemoptysis	Consumer	Neurontin			
ORAL		Head Discomfort		(Gabapentin)	PS		ORAL
		Headache		Bupropion			
		Neurological Symptom		Hydrochloride			
		Pain In Extremity		(Bupropion			
				Hydrochloride)	C		
				Tramadol (Tramadol)	C		
				Timolol Maleate			
				(Timolol Maleate)	C		
				Losartan Potassium			
				(Losartan Potassium)	C		
				Herbal Preparation			
				(Herbal Preparation)	C		
				Enzymes (Enzymes)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Centrym Silver
 (Ascorbic Acid,
 Calcium, Minerals
 Nos, Retinol,
 Tocopheryl Acetate, C
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C

Date:10/15/04ISR Number: 4505233-1Report Type:Periodic Company Report #2004045942
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Deafness	Health Professional	Neurontin (Gabapentin)	PS		ORAL
500 MG (2 IN 1 D), ORAL							

Date:10/15/04ISR Number: 4505236-7Report Type:Periodic Company Report #2004046202
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Renal Failure Urinary Retention	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505239-2Report Type:Periodic Company Report #2004046505
 Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Contusion Platelet Count Decreased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
EVERY DAY, ORAL							
			Company Representative	Acetylsalicylic Acid			

(Acetylsalicylic
Acid)

C

Date:10/15/04ISR Number: 4505241-0Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2004034323

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Failure Congestive Diabetes Mellitus Neuropathy Pneumonia Renal Failure	Consumer Health Professional	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505251-3Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2004034726

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 4800 MG		Intentional Misuse Medication Error	Health Professional	Neurontin (Gabapentin)	PS		
Other		Palpitations					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505268-9Report Type:Periodic
Age:43 YR Gender:Female I/FU:I

Company Report #2004035088

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Amnesia Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG 3 IN 1 D) ORAL		Corrective Lens User Hearing Impaired Suicide Attempt Weight Increased		Vicodon (Hydrocodone Bitartrate, Paracetamol)	C		

Date:10/15/04ISR Number: 4505270-7Report Type:Periodic
Age:69 YR Gender:Female I/FU:I

Company Report #2004035449

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Ageusia Anorexia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
1200 MG (300 MG, Q6H) INTERVAL: ONCE A DAY) ORAL		Dysgeusia Feeling Abnormal Nausea Pain	Professional	Methadone (Methadone)	SS		
10 MG (2.5 MG, 4 IN 1 D)				Estradiol Vitamins	C C		

Date:10/15/04ISR Number: 4505272-0Report Type:Periodic
Age:39 YR Gender:Female I/FU:I

Company Report #2004035950

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness Judgement Impaired	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Somnolence Weight Increased		Tizanidine Hydrochloride	C		
				Valdecoxib	C		
				Bupropion Hydrochloride	C		
				Propacet (Dextropropoxyphene Napsilate, Paracetamol)	C		
				Tramadol Hydrochloride	C		
				Amitriptyline	C		
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		
				Zolpidem Tartrate	C		

Date:10/15/04ISR Number: 4505275-6Report Type:Periodic Company Report #2004036485
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness	Health Professional Company Representative	Neurontin (Gabapentin) Ethanol (Ethanol)	PS SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505276-8Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2004036756

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intentional Self-Injury	Consumer	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505278-1Report Type:Periodic
 Age:62 YR Gender:Female I/FU:I

Company Report #2004038006

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Aphasia Cardiac Failure	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (300 Other M, 1 IN 1 D)		Cerebrovascular Accident					
ORAL		Feeling Abnormal					
		Myocardial Infarction Speech Disorder		All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:10/15/04ISR Number: 4505280-XReport Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #2004038267

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505281-1Report Type:Periodic
 Age:80 YR Gender:Male I/FU:I

Company Report #2004038469

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Renal Failure	Consumer	Neurontin (Gabapentin)	PS	ORAL
ORAL			Levothyroxine Sodium	C	
			Allopurinol	C	
			Folic Acid	C	

Date:10/15/04ISR Number: 4505306-3Report Type:Periodic Company Report #2004028049
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation Personality Change Suicidal Ideation	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505308-7Report Type:Periodic Company Report #2004031558
 Age:37 YR Gender:Male I/FU:I

Outcome	PT
Other	Dysphonia Hypersensitivity Local Swelling Pain

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pharyngolaryngeal Pain Swelling Face Upper Respiratory Tract Infection	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1 D), ORAL			Health Professional	Neurontin (Gabapentin)	PS		ORAL

Oxycocet (Oxycodone Hydrochloride, Paracetamol)	C
Vicodin (Hydrocodone Bitartrate, Paracetamol)	C
Rofecoxib (Rofecoxib)	C
Amitriptyline (Amitriptyline)	C
Cyclobenzaprine (Cyclobenzaprine)	C

Date:10/15/04ISR Number: 4505310-5Report Type:Periodic Company Report #2004032380
Age:44 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Aggression Asthenia Drug Ineffective Hypomania Unevaluable Event	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin)	PS		
				Topiramate (Topiramate)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		

Date:10/15/04ISR Number: 4505311-7Report Type:Periodic Company Report #2004032846
Age:81 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Bedridden	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin			

(100 MG,
UNKNOWN)
Cerebrovascular Accident
Dizziness
Muscle Spasms
Myalgia
(Gabapentin)
PS

Date:10/15/04ISR Number: 4505312-9Report Type:Periodic Company Report #2004033000
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Diplopia Dizziness	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
(300 MG, EVERYDAY), ORAL		Dysarthria Gait Disturbance Nausea Pruritus Unevaluable Event Vision Blurred Visual Disturbance	Professional				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505313-0Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2004033165

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1700 MG (TID) Other	Nephrolithiasis	Health Professional	Neurontin (Gabapentin)	PS		
			Sertraline Hydrochloride (Sertraline Hydrochloride)	C		

Date:10/15/04ISR Number: 4505314-2Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2004033214

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG (100 MG, DAILY), ORAL	Blood Pressure Fluctuation Blood Pressure Increased Dizziness Headache Insomnia Memory Impairment Muscle Twitching Nausea Tinnitus	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Warfarin (Warfarin)	C		
			Terazosin (Terazosin)	C		
			Hydrochlorothiazide (Hydrochlorothiazide)	C		
			Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
			Lisinopril (Lisinopril)	C		
			Atenolol (Atenolol)	C		
			Ranitidine (Ranitidine)	C		
			Combivent (Ipratropium Bromide, Salbutamol Sulfate)	C		
			Flunisolide (Flunisolide)	C		

Multivitamins
(Ascorbic Acid,
Ergocalciferol,
Folic Acid,
Nicotianamide, C
Lorazepam
(Lorazepam) C
Vicodin (Hydrocodone
Bitartrate,
Paracetamol) C
Escitalopram
(Escitalopram) C

Date:10/15/04ISR Number: 4505316-6Report Type:Periodic
Age:62 YR Gender:Female I/FU:I

Company Report #2004033865

Outcome PT
Disability Blindness
Other Eye Haemorrhage
Glaucoma
Impaired Healing

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pain Visual Disturbance Vomiting	Report Source	Product	Role	Manufacturer	Route
ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Glimepiride (Glimepiride)	C		

Date:10/15/04ISR Number: 4505317-8Report Type:Periodic Company Report #2004033866
 Age:26 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (600 Other MG, BID		Amnesia Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
INTERVAL: EVERY DAY),		Depression Fall Feeling Abnormal Loss Of Consciousness Road Traffic Accident		All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:10/15/04ISR Number: 4505320-8Report Type:Periodic Company Report #2004033872
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Liver Disorder Pyrexia Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Valproate Semisodium (Valproate Semisodium)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Diarrhoea Haematemesis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL		Haemoptysis					
250 MG (DAILY), ORAL		Nausea Vomiting		Gefitinib (Gefitinib)	SS		ORAL
				Propacet (Dextropropoxyphene Napsilate, Paracetamol)	C		
				Folic Acid (Folic Acid)	C		
				Calcium (Calcium)	C		
				Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotianamide,	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505322-1Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #2004021663

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Loss Of Consciousness	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505324-5Report Type:Periodic
Age:33 YR Gender:Male I/FU:I

Company Report #2004023142

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 1800 MG		Oedema Pneumonia Viral	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other (DAILY), ORAL		Sciatica Tooth Infection Toothache					

Date:10/15/04ISR Number: 4505325-7Report Type:Periodic
Age:60 YR Gender:Female I/FU:I

Company Report #2004024973

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		White Blood Cell Count Decreased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
1200 MG (300 MG, 4 IN 1 D), ORAL				Celecoxib (Celecoxib) Venlafaxine Hydrochloride (Venlafaxine Hydrochloride) Tramadol Hydrochloride	C		

(Tramadol
Hydrochloride) C

Date:10/15/04ISR Number: 4505326-9Report Type:Periodic Company Report #2004025941
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident Convulsion	Consumer	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505327-0Report Type:Periodic Company Report #2004026236
Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorexia Back Pain	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
300 MG (100		Difficulty In Walking	Professional				
MG, 3 IN 1		Pain In Extremity					
D), ORAL		Rectal Haemorrhage Weight Increased		Celecoxib (Celecoxib)	SS		ORAL
ORAL				Esomeprazole (Esomeprazole) Hemocyte-F Elixir	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Cyanocobalamin,
Folic Acid,
Polysaccharide-Iron
Complex) C
Paroxetine
Hydrochloride
(Paroxetine
Hydrochloride) C
Co-Diovan
(Hydrochlorothiazide
, Vaslartan) C
... C

Date:10/15/04ISR Number: 4505329-4Report Type:Periodic Company Report #2004026622
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Health Professional	Neurontin (Gabapentin)	PS		
1000 MG							
(DAILY),							

Date:10/15/04ISR Number: 4505331-2Report Type:Periodic Company Report #2004026711
Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agoraphobia Cataract	Consumer	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL							
25 MG		Chest Pain		Zoloft (Sertraline)	SS		ORAL
(DAILY), ORAL		Depression					
		Diarrhoea		Clonazepam (Clonazepam)	C		
		Dizziness					
		Drug Ineffective		Rabeprazole Sodium (Rabeprazole Sodium)	C		
		Feeling Abnormal					
		Nervousness		Lansoprazole (Lansoprazole)	C		
		Phobia					
		Vision Blurred		Maalox (Aluminium Hydroxide Gel,			
		Vomiting					

Date:10/15/04ISR Number: 4505332-4Report Type:Periodic
 Age:80 YR Gender:Male I/FU:I

Company Report #2004026960

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Oedema Peripheral Pulmonary Oedema	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG (100, TID), ORAL				Pravastatin Sodium (Pravastatin Sodium)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Valsartan (Valsartan)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505364-6Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2004027485

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation	Health	Neurontin			
ORAL		Dizziness	Professional	(Gabapentin)	PS		ORAL
		Dysstasia		Hydrocodone			
		Insomnia		(Hydrocodone)	C		
		Loss Of Consciousness		Carisoprodol			
		Pain		(Carisoprodol)	C		
		Rash Pruritic		Morphine (Morphine)	C		
				All Other			
				Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	C		
				Lidocaine			
				(Lidocaine)	C		

Date:10/15/04ISR Number: 4505366-XReport Type:Periodic
 Age:43 YR Gender:Female I/FU:I

Company Report #2004016724

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Neurontin			
900 MG DAILY		Dizziness	Health	(Gabapentin)	PS		ORAL
ORAL		Photophobia	Professional				
		Visual Disturbance		Metaxalone	C		

Date:10/15/04ISR Number: 4505368-3Report Type:Periodic
 Age:33 YR Gender:Female I/FU:I

Company Report #2004017465

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Adnexa Uteri Pain	Consumer	Neurontin			
900 MG TID		Blepharospasm		(Gabapentin)	PS		ORAL
ORAL		Eye Pain					

Menorrhagia
Metrorrhagia
Pelvic Pain
Phobia Of Driving
Reading Disorder
Vision Blurred

Vicodin
(Paracetamol,
Hydrocodone
Bitartrate)

C

Date:10/15/04ISR Number: 4505370-1Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2004017469

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Pulmonary Oedema Renal Disorder	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505371-3Report Type:Periodic
Age:63 YR Gender:Female I/FU:I

Company Report #2004018046

Outcome	PT
Other	Blister Fluid Retention Hip Swelling

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1 D)		Infection Musculoskeletal Stiffness	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Pioglitazone (Pioglitazone)	SS		
				Oxycocet (Paracetamol, Oxycodone Hydrochloride)	C		
				Digoxin	C		
				Metoprolol	C		

Date:10/15/04ISR Number: 4505374-9Report Type:Periodic Company Report #2004018157
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Cholesterol Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Blood Sodium Decreased Weight Increased		Oxcarbazepine (Oxcarbazepine)	SS		
				Clonazepam	C		
				Risperidone	C		
				Valproate Semisodium	C		
				Aripiprazole	C		

Date:10/15/04ISR Number: 4505376-2Report Type:Periodic Company Report #2004019211
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Stevens-Johnson Syndrome	Health Professional	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4505377-4Report Type:Periodic Company Report #2004019527
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea	Consumer	Neurontin			
400 MG ID		Muscle Spasms		(Gabapentin)	PS		ORAL
ORAL		Weight Increased					

Date:10/15/04ISR Number: 4505380-4Report Type:Periodic Company Report #2004019776
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Neurontin			
900 MG TID		Convulsion		(Gabapentin)	PS		ORAL
ORAL		Feeling Abnormal					
		Nausea		Oxycodone			
		Unevaluable Event		Hydrochloride			
				(Oxycodone			
				Hydrochloride)	SS		
				Methadone	C		
				Venlafaxine			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride C
 Vicodin
 (Paracetamol,
 Hydrocodone
 Bitartrate) C
 Trazodone C

Date:10/15/04ISR Number: 4505381-6Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2004020182

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain	Consumer	Neurontin			
		Atrial Fibrillation		(Gabapentin)	PS		
		Food Allergy		All Other			
		Hernia		Non-Therapeutic			
		Pain		Products (All Other			
		Unevaluable Event		Non-Therapeutic			
		Vision Blurred		Products)	SS		
				All Other			
				Therapeutic Products	C		

Date:10/15/04ISR Number: 4505382-8Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #2004021461

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Platelet Count Decreased	Consumer	Neurontin			
		Red Blood Cell Count		(Gabapentin)	PS		ORAL
900 MG TID		Increased					
ORAL		White Blood Cell Count		Erythropoietin	C		
		Decreased		Antihypertensive	C		
				Anti-Diabetics	C		
				Cardiovascular			
				System Drugs	C		

Date:10/15/04ISR Number: 4505384-1Report Type:Periodic
 Age:6 YR Gender:Female I/FU:I

Company Report #2004028505

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Complex Partial Seizures Intentional Misuse Medication Error	Health Professional	Neurontin (Solution) (Gabapentin)	PS		ORAL
3375 MG (1125 MG, 3 IN 1 D), ORAL				Valproate Semisodium (Valproate Semisodium)	C		

Date:10/15/04ISR Number: 4505387-7Report Type:Periodic Company Report #2004034117
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4505391-9Report Type:Periodic
Age:55 YR Gender:Female I/FU:I

Company Report #2004037845

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Abdominal Pain Upper	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL		Abnormal Faeces Faeces Discoloured Haematemesis Haematochezia Haemorrhoids Nausea	Professional	Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotinamide, Tocopherol (Tocopherol) Acetylsalicylic Acid (Acetylsalicylic Acid)	C C C		

Date:10/15/04ISR Number: 4505392-0Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2004040268

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Upper Atrial Fibrillation	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1500 MG (2 IN 1 D), ORAL		Blood Pressure Increased Erectile Dysfunction Hyperthyroidism		Clonazepam (Clonazepam)	SS		ORAL
3 MG (3 MG, 1 IN 1 D), ORAL		Insomnia Pollakiuria Pruritus Rash		Levothyroxine Sodium (Levothyroxine Sodium)	SS		
ORAL		Urinary Retention		Lovastatin (Lovastatin) Vitamins (Vitamins)	C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Neurontin (Tablets)			
(3 IN 1 D),		Drug Ineffective		(Gabapentin)	PS		ORAL
ORAL		Neuropathy Peripheral					
		Vision Blurred		Folic Acid (Folic Acid)	C		
				Furosemide (Furosemide)	C		
				Losartan Potassium (Losartan Potassium)	C		
				Spiroinolactone (Spiroinolactone)	C		
				Phenoxyethylpenicillin Potassium (Phenoxyethylpenicillin Potassium)	C		
				Potassium Chloride (Potassium Chloride)	C		
				Warfarin Sodium (Warfarin Sodium)	C		
				Digoxin (Digoxin)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ranitidine
Hydrochloride
(Ranitidine
Hydrochloride) C

Date:10/15/04ISR Number: 4507714-3Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2003035148

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG Other (TID), ORAL		Amnesia Body Height Decreased Cerebrovascular Accident Convulsion	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Levetiracetam (Levetiracetam)	C		
				Galantamine (Galantamine)	C		
				Famotidine (Famotidine)	C		
				Simvastatin (Simvastatin)	C		
				Folic Acid (Folic Acid)	C		
				Celecoxib (Celecoxib)	C		
				Multivitamins (Ergocalciferol, Ascorbic Acid, Folic Acid, Thiamine Hydrohchloride, Acetylsalicylic Acid (Acetylsalicylic Acid)	C		

Date:10/15/04ISR Number: 4507715-5Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2003037234

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Pressure Abnormal	Consumer	Neurontin (Tablets)			

900 MG (TID),	Contusion	(Gabapentin)	PS	ORAL
ORAL	Insomnia			
	Medication Error	Norvasc (Amlodipine		
	Pain	Besilate)		
MG (DAILY),	Platelet Count Decreased	(Amlodipine)	SS	ORAL
ORAL	Rash Macular			
		Simvastatin		
20 MG		(Simvastatin)	SS	ORAL
(DAILY), ORAL				
		Verapamil		
		(Verapamil)	C	
		Allopurinol		
		(Allopurinol)	C	
		Amiodarone		
		(Amiodarone)	C	
		Doxazosin Mesilate		
		(Doxazosin Mesilate)	C	
		Rofecoxib		
		(Rofecoxib)	C	
		All Other		
		Therapeutic Products		

Freedom Of Information (FOI) Report

(All Other
Therapeutic
Products) C

Date:10/15/04ISR Number: 4507720-9Report Type:Periodic
Age:76 YR Gender:Male I/FU:I

Company Report #2003040517

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (DAILY), ORAL		Herpes Zoster Medication Error Platelet Count Decreased Pruritus Somnolence Weight Decreased	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL

Date:10/15/04ISR Number: 4507722-2Report Type:Periodic
Age:28 YR Gender:Female I/FU:I

Company Report #2003116563

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 Muscle Twitching Suicide Attempt	Health Professional	Neurontin (Tablets) (Gabapentin) Bupropion Hydrochloride (Bupropion Hydrochloride) Baclofen (Baclofen) Vicodin (Paracetamol, Hydrocodone Bitartrate) Zolpidem Tartrate (Zolpidem Tartrate) Clonazepam (Clonazepam) Fluoxetine Hydrochloride (Fluoxetine)	PS SS C C C C		ORAL

Hydrochloride) C
Queitapine Fumarate
(Quetiapine
Fumarate) C

Date:10/15/04ISR Number: 4507723-4Report Type:Periodic Company Report #2003119381
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Depression	Consumer	Neurontin (Tablets)			
Other		Hallucination, Auditory	Health	(Gabapentin)	PS		ORAL
ORAL		Muscle Spasms	Professional				
		Pain					
		Suicidal Ideation					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4507724-6Report Type:Periodic
Age:40 YR Gender:Male I/FU:I

Company Report #2004004022

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2400 MG (800, TID), ORAL ORAL		Hallucination, Auditory Insomnia Medication Error Overdose Paranoia Psychomotor Hyperactivity Suicidal Ideation Weight Increased	Consumer	Neurontin (Tablets) (Gabapentin) Clonazepam (Clonazepam) Valproate Semisodium (Valproate Semisodium) Doxepin Hydrochloride (Doxepin Hydrochloride)	PS SS C C		ORAL ORAL

Date:10/15/04ISR Number: 4507728-3Report Type:Periodic
Age: Gender:Female I/FU:I

Company Report #2004008686

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Anxiety Blood Urine Calculus Urinary Delirium Difficulty In Walking Feeling Abnormal Gastroenteritis Viral Haematochezia Hunger Hyperphagia Insomnia Intentional Misuse Irritability Muscle Spasms Myalgia Nervousness Neuralgia Pain	Consumer	Neurontin (Tablets) (Gabapentin) Antidepressants (Antidepressants) All Other Therapeutic Products (All Other Therapeutic Products)	PS SS SS		ORAL

Paraesthesia
Sensation Of Pressure
Sinusitis
Somnolence
Urinary Tract Infection
Weight Decreased
Weight Increased

Date:10/15/04ISR Number: 4507729-5Report Type:Periodic
Age:72 YR Gender:Male I/FU:I

Company Report #2004024702

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG (BID), ORAL		Fall Insomnia Pain Patella Fracture	Consumer	Neurontin (Tablets) (Gabapentin) Lorazepam (Lorazepam) Bupropion Hydrochloride (Bupropion Hydrochloride)	PS C C		ORAL

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Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4507735-0Report Type:Periodic
Age:43 YR Gender:Female I/FU:I

Company Report #2004029814

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin (Tablets)			
600 MG		Migraine		(Gabapentin)	PS		ORAL
(DAILY), ORAL		Trigeminal Neuralgia					
				Levothyroxine Sodium			
				(Levothyroxine Sodium)	C		
				Pantoprazole			
				(Pantoprazole)	C		

Date:10/15/04ISR Number: 4507736-2Report Type:Periodic
Age: Gender: I/FU:I

Company Report #2004034116

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Neurontin (Tablets)			
				(Gabapentin)	PS		

Date:10/15/04ISR Number: 4511063-7Report Type:Periodic
Age:53 YR Gender:Female I/FU:I

Company Report #2004012890

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyslogia	Consumer	Neurontin			
900 MG (TID),		Muscle Twitching		(Gabapentin)	PS		ORAL
ORAL		Pain					
		Paraesthesia		All Other Therapeutic Products			
				(All Other Therapeutic Products)	C		

Date:10/15/04ISR Number: 4511064-9Report Type:Periodic Company Report #2004012911
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deafness	Consumer	Neurontin (Gabapentin)	PS		ORAL
1800 MG							
(DAILY), ORAL							

Rofecoxib (Rofecoxib)	C
Montelukast Sodium (Montelukast Sodium)	C
Prednisone (Prednisone)	C

Date:10/15/04ISR Number: 4511065-0Report Type:Periodic Company Report #2004013495
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Other		Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4511066-2Report Type:Periodic
Age:36 YR Gender:Female I/FU:I

Company Report #2004013500

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Back Pain Intentional Misuse	Consumer Health	Neurontin (Gabapentin)	PS		
TID Other ORAL	Medication Error Suicide Attempt	Professional	Clonazepam (Clonazepam)	SS		ORAL
			Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		
			Oxycocet (Oxycodone Hydrochloride, Paracetamol)	C		
			Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		

Date:10/15/04ISR Number: 4511067-4Report Type:Periodic
Age: Gender:Male I/FU:I

Company Report #2004013511

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Feeling Abnormal Suicidal Ideation Tinnitus	Consumer	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4511068-6Report Type:Periodic
Age: Gender:Unknown I/FU:I

Company Report #2004013669

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Cardiac Failure Congestive	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cataract Subcapsular Neuropathy	Consumer	Neurontin (Gabapentin)	PS		ORAL
500 MG (BID), ORAL				Atenolol (Atenolol) Hyzaar (Hydrochlorothiazide , Losartan Potassium) Acetylsalicylic Acid (Acetylsalicylic Acid) Antithrombotic Agents (Antithrombotic Agents) Torasemide (Torasemide) Cardiovascular System Drugs	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Cardiovascular System Drugs) C

Date:10/15/04ISR Number: 4511070-4Report Type:Periodic Company Report #2004014252
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Convulsion	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4511071-6Report Type:Periodic Company Report #2004015524
 Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Magnesium Decreased Blood Potassium Decreased	Consumer	Neurontin (Gabapentin)	PS		
600 MG (DAILY)		Intentional Misuse		Quetiapine Fumarate (Quetiapine Fumarate)	C		
		Medication Error Suicide Attempt		Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
				Valproate Semisodium (Valproate Semisodium)	C		

Date:10/15/04ISR Number: 4511072-8Report Type:Periodic Company Report #2004016555
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cardiac Failure Congestive	Health Professional Company	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4511073-XReport Type:Periodic
Age:79 YR Gender:Female I/FU:I

Company Report #2004008374

Outcome	PT
Disability	Anxiety
Other	Asthenopia
	Corneal Disorder
	Depressed Mood
	Difficulty In Walking
	Diplopia
	Disorientation
	Dizziness
	Eye Disorder
	Eye Irritation
	Eye Pain
	Fear Of Falling
	Headache
	Impaired Driving Ability
	Impaired Work Ability

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
		Ivth Nerve Paralysis Nuclear Magnetic Resonance Imaging					
1200 MG		Abnormal Pain	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
(TID), ORAL		Unevaluable Event	Professional				
		Visual Disturbance		Atorvastatin (Atorvastatin)	C		
				Esomeprazole (Esomeprazole)	C		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		
				Valsartan (Valsartan)	C		
				Amlodipine Besilate (Amlodipine Besilate)	C		
				Celecoxib (Celecoxib)	C		

Date:10/15/04ISR Number: 4511074-1Report Type:Periodic Company Report #2004009092
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Headache Intentional Misuse Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4511075-3Report Type:Periodic Company Report #2004009336
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Consumer	Neurontin (Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health	Neurontin			
600 MG (BID),		Nausea	Professional	(Gabapentin)	PS		ORAL
ORAL		Pain					
				All Other Therapeutic (All Other Therapeutic Products)	C		
				Alprazolam (Alprazolam)	C		
				Estrogens Conjugated (Estrogens Conjugated)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/15/04ISR Number: 4511077-7Report Type:Periodic
 Age: Gender:Female I/FU:I

Company Report #2004009795

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1200 MG (QID), ORAL		Depression Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Vicodin (Paracetamol, Hydrocodone Bitartrate)	C		
				Tizanidine Hydrochloride (Tizanidine Hydrochloride)	C		
				Trazodone (Trazodone)	C		
				Modafinil (Modafinil)	C		
				Prednisone (Prednisone)	C		

Date:10/15/04ISR Number: 4511078-9Report Type:Periodic
 Age:61 YR Gender:Male I/FU:I

Company Report #2004009820

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (DAILY), ORAL		Abnormal Dreams Arthritis	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Balance Disorder	Professional				
		Disorientation		Celebrex (Celecoxib)	SS		ORAL
		Headache					
		Loss Of Consciousness					

Date:10/15/04ISR Number: 4511079-0Report Type:Periodic
 Age: Gender:Male I/FU:I

Company Report #2004011342

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		

Date:10/15/04ISR Number: 4511080-7Report Type:Periodic Company Report #2004011602
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness Vision Blurred	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (BID)							
, ORAL							

Date:10/15/04ISR Number: 4511081-9Report Type:Periodic Company Report #2004011726
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							
Oxycodone Hydrochloride (Oxycodone Hydrochloride)							
SS							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lisinopril
(Lisinopril) SS

Date:10/15/04ISR Number: 4511082-0Report Type:Periodic Company Report #2004012512
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG, TID), ORAL		Asthenia Convulsion Hot Flush Panic Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
60 MG (60 MG, DAILY), ORAL				Phenobarbital (Phenobarbital)	SS		ORAL
				Quetiapine Fumarate (Quetiapine Fumarate)	SS		
				Oxcarbazepine (Oxcarbazepine)	SS		
				Escitalopram (Escitalopram)	C		
				Alprazolam (Alprazolam)	C		
				Clonazepam (Clonazepam)	C		

Date:10/15/04ISR Number: 4511083-2Report Type:Periodic Company Report #2004004295
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 6 YR		Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety Depression Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatinine Increased	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Renal Failure Acute					
				Valsartan (Valsartan)	C		
				Simvastatin (Simvastatin)	C		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Atenolol (Atenolol) C
 Lisinopril
 (Lisinopril) C

Date:10/15/04ISR Number: 4511086-8Report Type:Periodic Company Report #2004005857
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Chest Pain Renal Failure	Consumer	Gabapentin (Gabapentin)	PS		ORAL
ORAL							
Other		Somnolence					

Date:10/15/04ISR Number: 4511087-XReport Type:Periodic Company Report #2004005894
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Deafness	Health Professional	Neurontin (Gabapentin)	PS		
				Vfend (Voriconazole)	SS		
				Linezolid (Linezolid)	C		
				Lansoprazole (Lansoprazole)	C		

Date:10/15/04ISR Number: 4511088-1Report Type:Periodic Company Report #2004006516
 Age:9 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion Dehydration	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							
		Intentional Misuse Medication Error		Leuprorelin Acetate (Leuprorelin Acetate)	C		

Date:10/15/04ISR Number: 4511089-3Report Type:Periodic Company Report #2004007147
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Suicidal Ideation	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG, ORAL			Company Representative	Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:10/15/04ISR Number: 4511090-XReport Type:Periodic Company Report #2004007393
Age:89 YR Gender:Female I/FU:I

Outcome	PT
Other	Convulsion Defaecation Urgency

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Fatigue Muscle Contractions Involuntary	Report Source	Product	Role	Manufacturer	Route
900 MG (300, TID), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Donepezil Hydrochloride (Donepezil Hydrochloride)	C		

Date:10/15/04ISR Number: 4511091-1Report Type:Periodic Company Report #2004007889
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Accidental Overdose Diplopia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (TID), ORAL		Road Traffic Accident	Professional				
				Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
				Zolpidem Tartrate (Zolpidem Tartrate)	C		

Date:10/15/04ISR Number: 4511092-3Report Type:Periodic Company Report #2004008326
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Aggression Decreased Interest	Consumer	Neurontin (Gabapentin)	PS		ORAL
1600 MG (TID), ORAL		Depression Mood Swings Pain					

Suicidal Ideation
Unevaluable Event

Date:10/15/04ISR Number: 4511093-5Report Type:Periodic Company Report #2004001218
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anger	Health	Neurontin			
Other		Depression	Professional	(Gabapentin)	PS		
		Suicidal Ideation		Ritonavir			
				(Ritonavir)	C		
				Atazanavir			
				(Atazanavir)	C		
				Zidovudine			
				W/Lamivudine			
				(Zidovudine,			
				Lamivudine)	C		
				Cetirizine			
				Hydrochloride			
				(Cetirizine			
				Hydrochloride)	C		
				Fluticasone			
				(Propionate)			
				(Fluticasone			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Propionate) C
 Allopurinol
 (Allopurinol) C
 Aciclovir
 (Aciclovir) C
 Azithromycin
 (Azithromycin) C
 Bactrim
 (Sulfamethoxazole,
 Trimethoprim) C

Date:10/15/04ISR Number: 4511094-7Report Type:Periodic Company Report #2004001304
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Accidental Overdose	Consumer	Neurontin			
Other		Chest Pain		(Gabapentin)	PS		ORAL
2400 MG		Emotional Disorder					
(TID), ORAL		Suicidal Ideation		Zoloft (Sertraline)	C		
		Thinking Abnormal		Clonazepam			
				(Clonazepam)	C		
				Bupropion			
				Hydrochloride			
				(Bupropion			
				Hydrochloride)	C		
				Thyroid (Thyroid)	C		
				Methadone			
				(Methadone)	C		
				Oxycodone			
				(Oxycodone)	C		

Date:10/15/04ISR Number: 4511095-9Report Type:Periodic Company Report #2004001691
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Neurontin			
Other		Skeletal Injury		(Gabapentin)	PS		ORAL
300 MG		Suicide Attempt					
(DAILY), ORAL							

Effexor (Venlafaxine
Hydrochloride) C
Diazepam (Diazepam) C

Date:10/15/04ISR Number: 4511096-0Report Type:Periodic Company Report #2004001692
Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Renal Impairment	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Sleep Disorder					

Date:10/15/04ISR Number: 4511097-2Report Type:Periodic Company Report #2004001917
Age:25 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Abnormal Dreams
Initial or Prolonged	Balance Disorder
Other	Bipolar Disorder
	Depression
	Memory Impairment

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Freedom Of Information (FOI) Report

Dose	Duration	Mental Disorder Obsessive-Compulsive Disorder	Report Source	Product	Role	Manufacturer	Route
ORAL		Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Alprazolam (Alprazolam)	C		

Date:10/15/04ISR Number: 4511098-4Report Type:Periodic Company Report #2004002063
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anaemia Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other				Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		ORAL
				Naproxen (Naproxen)	C		
				Spironolactone (Spironolactone)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Furosemide (Furosemide)	C		
				Allopurinol (Allopurinol)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Bumetanide (Bumetanide)	C		
				Trandolapril (Trandolapril)	C		
				Combivent (Ipratropium Bromide, Salbutamol Sulfate)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Consumer	Neurontin			
900 MG		Fall		(Gabapentin)	PS		ORAL
(DAILY), ORAL		Humerus Fracture					
		Inappropriate Affect		Amitriptyline			
		Pain In Extremity		(Amitriptyline)	SS		
		Polyneuropathy		Carbamazepine			
ORAL				(Carbamazepine)	SS		ORAL
				Atorvastatin			
				(Atorvastatin)	C		
				Rosiglitazone			
				Maleate			
				(Rosiglitazone			
				Maleate)	C		
				Glimepiride			
				(Glimepiride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lisinopril
 (Lisinopril) C
 Metformin
 Hydrochloride
 (Metformin
 Hydrochloride) C

Date:10/15/04ISR Number: 4511100-XReport Type:Periodic Company Report #2004003996
 Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (APPROXIMATELY 900MG QD), ORAL		Amnesia Confusional State Delirium Drooling Hallucination Influenza Speech Disorder	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Zolpidem Tartrate (Zolpidem Tartrate) Hydromorphone Hydrochloride (Hydromorphone Hydrochloride)	C C		

Date:10/15/04ISR Number: 4511101-1Report Type:Periodic Company Report #2004004008
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300, Other BID), ORAL		Convulsion Micturition Disorder Pollakiuria Renal Failure Renal Pain	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:10/15/04ISR Number: 4511102-3Report Type:Periodic Company Report #2004004012
Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Renal Failure Acute	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/18/04ISR Number: 4480032-8Report Type:Expedited (15-DaCompany Report #001-0945-M0000697
Age:44 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain Upper
Initial or Prolonged	Abdominal Tenderness
Disability	Acrochordon
Other	Actinic Keratosis
	Acute Sinusitis
	Albumin Globulin Ratio
	Decreased
	Alcoholism
	Amnesia
	Angina Unstable
	Anticonvulsant Drug Level

Freedom Of Information (FOI) Report

Below Therapeutic
Anticonvulsant Drug Level
Increased
Anxiety
Aphasia
Asthenia
Back Injury
Back Pain
Bipolar Disorder
Bladder Disorder
Blood Alkaline
Phosphatase Increased
Blood Calcium Decreased
Blood Cholesterol
Increased
Blood Glucose Increased
Blood Triglycerides
Increased
Blood Urea
Nitrogen/Creatinine Ratio
Increased
Bone Cyst
Brain Neoplasm
Bronchitis Acute
Cardiac Arrest
Carotid Artery Disease
Chest Pain
Chest Wall Mass
Chest Wall Pain
Cholecystitis
Cholelithiasis
Chondropathy
Cognitive Disorder
Coma
Communication Disorder
Condyloma Acuminatum
Conversion Disorder
Convulsion
Coordination Abnormal
Deafness Unilateral
Depressed Level Of
Consciousness
Depression
Dermal Cyst
Dermatitis Contact
Diabetes Mellitus
Difficulty In Walking
Diplopia

Disease Recurrence
Dissociative Disorder
Disturbance In Attention
Dizziness
Drug Ineffective
Dysphemia
Dysstasia
Ear Discomfort
Electroencephalogram
Abnormal
Emotional Disorder
Enchondromatosis
Eosinophil Percentage
Increased

Freedom Of Information (FOI) Report

Ependymoma
Exostosis
Fall
Fatigue
Feeling Abnormal
Feeling Of Despair
Fibromyalgia
Folliculitis
Gait Disturbance
Gamma-Glutamyltransferase
Increased
Gastritis
Gastrooesophageal Reflux
Disease
Glioma
Gynaecomastia
Haematocrit Decreased
Hallucination
Headache
Hearing Impaired
Heart Rate Increased
Hepatic Enzyme Increased
Hepatic Steatosis
Hyperhidrosis
Hypersomnia
Impaired Driving Ability
Impaired Work Ability
Insomnia
Intervertebral Disc
Degeneration
Intervertebral Disc
Displacement
Intestinal Functional
Disorder
Irritability
Joint Injury
Joint Swelling
Learning Disorder
Lower Gastrointestinal
Haemorrhage
Lumbar Spinal Stenosis
Malaise
Marital Problem
Memory Impairment
Meniscus Lesion
Mobility Decreased
Monocyte Count Decreased

Monocyte Percentage
Increased
Muscle Rigidity
Musculoskeletal Stiffness
Nausea
Nervous System Disorder
Neurodegenerative
Disorder
Nuchal Rigidity
Obstructive Airways
Disorder
Osteoarthritis
Osteopenia
Pacemaker Complication

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration		Report Source	Product	Role	Manufacturer	Route
900 MG (1 D),		Pain					
		Paraesthesia					
ORAL		Paraneoplastic Syndrome	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Patellofemoral Pain Syndrome	Professional				
		Peptic Ulcer	Company Representative	Valproate Sodium (Valproate Sodium)	SS		
		Petit Mal Epilepsy		Buspirone Hydrochloride (Buspirone Hydrochloride)	C		
		Pharyngolaryngeal Pain		Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		
		Pigmented Naevus		Omeprazole (Omeprazole)	C		
		Plantar Fasciitis		Capsaicin (Capsaicin)	C		
		Post Procedural Complication		Celecoxib (Celecoxib)	C		
		Protein Total Decreased		Lansoprazole (Lansoprazole)	C		
		Radiculopathy					
		Reading Disorder					
		Red Blood Cell Sedimentation Rate Increased					
		Scar					
		Seborrhoeic Keratosis					
		Sensory Disturbance					
		Sick Sinus Syndrome					
		Soft Tissue Disorder					
		Soliloquy					
		Somnolence					
		Speech Disorder					
		Spinal Osteoarthritis					
		Staring					
		Stereotypy					
		Strabismus					
		Stress					
		Suicidal Ideation					
		Synovitis					
		Tension					
		Tinnitus					
		Toe Deformity					
		Tonsillar Disorder					
		Tremor					
		Urine Ketone Body Present					
		Vision Blurred					
		Visual Disturbance					
		Visual Tracking Test Abnormal					
		Weight Decreased					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 50 MG (50 MG, Initial or Prolonged 1 IN 1 D),		Activities Of Daily Living Impaired Blood Pressure Increased	Consumer	Zoloft (Sertraline)	PS		ORAL
ORAL		Dysphagia Dyspnoea Heart Rate Irregular		Neurontin (Gabapentin) (Gabapentin)	SS		ORAL
ORAL		Immobile Prescribed Overdose		Olanzapine (Olanzapine)	SS		ORAL
ORAL		Weight Decreased		Tolterodine L-Tartrate (Tolterodine			

Freedom Of Information (FOI) Report

ORAL	L-Tartrate)	SS	ORAL
	Rofecoxib (Rofecoxib)	SS	
	Atenolol (Atenolol)	C	
	Multivitamins (Ascorbic Acid/Ergocalciferol/ Folic Acid/Nicotinamide/Pa	C	
	Senna Fruit (Senna Fruit)	C	
	Donepezil Hydrochloride (Donepezil Hydrochloride)	C	
	Hemorid For Women Cream (Aloe Vera, Paraffin, Liquid Phenylephrine Hydrochloride, All Other Therapeutic Products	C	

Date:10/18/04ISR Number: 4480087-0Report Type:Expedited (15-DaCompany Report #2004056058
Age:84 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Adverse Drug Reaction Balance Disorder	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
600 MG (600 MG, 1 IN 1 D), ORAL		Disorientation Dizziness	Professional				
		Fall Gait Disturbance Hearing Impaired Vision Blurred		Vicodin (Hydrocodone Bitartrate, Paracetamol) Paracetamol (Paracetamol) Dyazide (Hydrochlorothiazide , Triamterene) Multivitamins (Ascorbic	C C C		

Date:10/18/04ISR Number: 4480088-2Report Type:Expedited (15-DaCompany Report #2004039112

Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain Upper
Initial or Prolonged	Activated Partial Thromboplastin Time Prolonged
	Anticoagulation Drug Level Above Therapeutic
	Anticoagulation Drug Level Below Therapeutic
	Bradyphrenia
	Cardiac Disorder
	Carpal Tunnel Syndrome

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness Fall Fatigue	Consumer	Neurontin (Gabapentin)	PS		
600 MG (200 MG, 3 IN 1 D)		Feeling Abnormal Gait Disturbance Gastric Haemorrhage					
400 MCG (200 MCG, 2 IN 1 D), ORAL		Head Injury Hypersomnia Rectal Haemorrhage Vaginal Haemorrhage		Celecoxib (Celecoxib)	SS		ORAL
				Warfarin Sodium (Warfarin Sodium)	C		
				Rabeprazole Sodium (Rabeprazole Sodium)	C		
				Ultracet (Paracetamol, Tramadol Hydrochloride)	C		
				Zolpidem Tartrate (Zolpidem Tartrate)	C		
				Estrogens Conjugated (Estrogens Conjugated)	C		
				Rabeprazole Sodium (Rabeprazole Sodium)	C		
				Montelukast Sodium (Montelukast Sodium)	C		

Date:10/18/04ISR Number: 4480091-2Report Type:Expedited (15-DaCompany Report #2004075222
Age:54 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG, ORAL		Amnesia Complex Regional Pain Syndrome	Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other 60 MG (15 MG,		Depression		Prednisone (Prednisone)	SS		ORAL

Disease Recurrence

Excitability

ORAL

Gastritis
 Headache
 Insomnia
 Mania
 Pain
 Pain In Extremity
 Sedation
 Suicidal Ideation
 Therapeutic Response
 Unexpected
 White Blood Cell Count
 Increased

Vicodin (Hydrocodon
 Bitartrate,
 Paracetamol) SS
 Escitalopram
 (Escitalopram) SS
 Provella-14
 (Estrogens
 Conjugated,
 Medroxyprogesterone
 Acetate) C
 Nifedipine
 (Nifedipine) C
 Propranolol
 Hydrochloride
 (Propranolol
 Hydrochloride) C
 Cetirizine
 Hydrochloride
 (Cetirizine
 Hydrochloride) C
 Lansoprazole
 (Lansoprazole) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/18/04ISR Number: 4480095-XReport Type:Expedited (15-DaCompany Report #2004048837

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anticonvulsant Drug Level Increased	Consumer Health	Dilantin Kapseals (Phenytoin Sodium)	PS		ORAL
SEE IMAGE							
		Asthenia	Professional	Neurontin (Gabapentin)	SS		ORAL
ORAL		Blood Pressure Diastolic					
		Increased		Cardura (Doxazosin Mesilate)	SS		ORAL
		Blood Pressure Increased					
		Carpal Tunnel Syndrome		(Doxazosin Mesilate)	SS		ORAL
6 MG (6 MG, 1							
IN 1 D), ORAL		Convulsion					
		Drug Level Decreased		Carbamazepine (Carbamazepine)	SS		
		Drug Toxicity		Oxcarbazepine (Oxcarbazepine)	SS		
		Fall		Estradiol (Estradiol)	C		
		Fatigue		Nabumetone (Nabumetone)	C		
		Feeling Abnormal		Fexofenadine Hydrochloride (Fexofenadine Hydrochloride)	C		
		Hip Arthroplasty		Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
		Hypersensitivity					
		Injury					
		Medical Device Implantation					
		Memory Impairment					
		Mental Impairment					
		Nausea					
		Nervous System Disorder					
		Neuropathy Peripheral					
		Spinal Osteoarthritis					
		Surgical Procedure Repeated					

Date:10/18/04ISR Number: 4480106-1Report Type:Expedited (15-DaCompany Report #K200401532

Age:44 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Levoxyl (Levothyroxine) Tablet, 25 Mcg	PS		ORAL
ORAL							

ORAL			Valproic Acid (Valproic Acid)	SS		ORAL
ORAL			Gabapentin (Gabapentin)	SS		ORAL

Date:10/18/04ISR Number: 4480118-8Report Type:Expedited (15-DaCompany Report #2004060020
Age:69 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Atrial Fibrillation	Foreign	Neurontin			
Hospitalization -		Blood Creatinine	Health	(Gabapentin)	PS		ORAL
300 MG (300		Increased	Professional				
Initial or Prolonged		Cardiac Failure	Company				
MG, 1 IN 1		Congestive	Representative	Antibiotics	C		
D), ORAL		Cerebral Atrophy		(Antibiotics)			
		Embolic Cerebral					
		Infarction					
		Liver Disorder					
		Renal Disorder					
		Somnolence					

Freedom Of Information (FOI) Report

Date:10/18/04ISR Number: 4480403-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200657
Age:44 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain Upper
Initial or Prolonged	Affect Lability
Disability	Agitation
Other	Anger
	Application Site
	Irritation
	Arthralgia
	Arthritis
	Arthropod Bite
	Asthenia
	Blood Glucose Increased
	Carpal Tunnel Syndrome
	Change Of Bowel Habit
	Chest Pain
	Chromaturia
	Constipation
	Contusion
	Convulsion
	Coordination Abnormal
	Deafness
	Decreased Appetite
	Depression
	Diarrhoea
	Disease Recurrence
	Dissociative Disorder
	Disturbance In Attention
	Dizziness
	Dry Mouth
	Dysarthria
	Dyspepsia
	Dysphagia
	Dyspnoea
	Dysuria
	Essential Tremor
	Eustachian Tube
	Dysfunction
	Fall
	Fibrocystic Breast
	Disease
	Flat Affect
	Gastrooesophageal Reflux
	Disease
	Goitre

Granuloma
Hallucination, Visual
Hand Fracture
Head Injury
Headache
Heart Rate Increased
Heart Rate Irregular
Hordeolum
Hot Flush
Hypoaesthesia
Hypothyroidism
Impaired Healing
Insomnia
Intervertebral Disc
Degeneration

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1800 MG	(DAILY), ORAL	Irritability				
		Joint Injury				
		Joint Sprain				
		Joint Swelling	Neurontin			
		Laryngitis	(Gabapentin)	PS		ORAL
		Legal Problem				
		Memory Impairment	Lamotrigine			
		Meniere'S Disease	(Lamotrigine)	C		
		Mental Impairment	Lithium Carbonate			
		Mouth Breathing	(Lithium Carbonate)	C		
		Nasal Ulcer	Clonazepam			
		Nausea	(Clonazepam)	C		
		Nervous System Disorder	Methylphenidate			
		Neuropathy	Hydrochloride			
		Neutrophil Count Abnormal	(Methylphenidate			
		Nystagmus	Hydrochloride)	C		
		Otitis Media	Levothyroxine Sodium			
		Pain In Extremity	(Levothyroxine			
		Paraesthesia	Sodium)	C		
		Pharyngitis	Liothyronine Sodium			
		Poor Peripheral	(Liothyronine			
		Circulation	Sodium)	C		
		Radiculopathy	Sertraline			
		Rash Erythematous	Hydrochloride			
		Red Blood Cell Count	(Sertraline			
		Decreased	Hydrochloride)	C		
		Red Cell Distribution	Pilocarpine			
		Width Increased	Hydrochloride			
		Rhinitis Allergic	(Pilocarpine			
		Road Traffic Accident	Hydrochloride)	C		
		Sedation	Metoprolol Succinate			
		Sinus Disorder	(Metoprolol			
		Skin Laceration	Succinate)	C		
		Sneezing	Lansoprazole			
		Somnolence	(Lansoprazole)	C		
		Stress	Hyoscyamine Sulfate			
		Suicidal Ideation	(Hyoscyamine			
		Swelling	Sulfate)	C		
		Syncope	Diltiazem			
		Tardive Dyskinesia	Hydrochloride			
		Tendonitis	(Diltiazem			
		Thirst	Hydrochloride)	C		
		Thyroid Disorder	Bupropion			
		Tinnitus	Hydrochloride			

Unresponsive To Verbal Stimuli	(Bupropion Hydrochloride	C
Urinary Tract Infection	Celebrex (Celecoxib)	C
Vaginal Candidiasis	Quetiapine Fumarate	
Vertigo Positional	(Quetiapine Fumarate)	C
Vestibular Neuronitis	Metoprolol Tartrate	
Vestibulitis	(Metoprolol Tartrate)	C
Viral Labyrinthitis	Citalopram	
Vision Blurred	Hydrobromide	
Weight Fluctuation	(Citalopram Hydrobromide)	C
White Blood Cell Count Decreased	Olanzapine	
Wound Infection	(Olanzapine)	C
	Omeprazole	
	(Omeprazole)	C

Freedom Of Information (FOI) Report

Nasal Preparations
 (Nasal Preparations) C
 Nefazodone
 Hydrochloride
 (Nefazodone
 Hydrochloride) C
 Totolin (Guiaenesin,
 Phenylpropanolamine
 Hydrochloride) C
 Meclozine
 (Meclozine) C
 Fluticasone
 Propionate
 (Fluticasone
 Propionate) C
 Trimethobenazamide
 Hydrochloride
 (Trimethobenzamide
 Hydrochloride) C
 Respaire-Sr-120
 (Guaifenesin,
 Pseudoephedrine
 Hydrochloride) C
 Risperidone
 (Risperidone) C

Date:10/18/04ISR Number: 4480444-2Report Type:Expedited (15-DaCompany Report #2004075431

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthritis	Consumer	Neurontin			
Other		Knee Arthroplasty		(Gabapentin)	PS		
600 MG		Osteoporosis		Fentanyl (Fentanyl)	SS		
75 MCG				Ibuprofen			
				(Ibuprofen)	SS		
				Rofecoxib			
				(Rofecoxib)	SS		
				Helianthus Tuberosus			
				(Helianthus			
				Tuberosus	SS		
				All Other			
				Therapeutic Products	SS		

Date:10/18/04ISR Number: 4480461-2Report Type:Expedited (15-DaCompany Report #2004076723

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Condition Aggravated	Consumer	Neurontin			
Initial or Prolonged		Depression		(Gabapentin)	PS		
		Suicidal Ideation					

Date:10/18/04ISR Number: 4480462-4Report Type:Expedited (15-DaCompany Report #2004066809

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Conversion Disorder	Consumer	Neurontin			
		Grand Mal Convulsion	Health	(Gabapentin)	PS		ORAL
		Petit Mal Epilepsy	Professional				

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Freedom Of Information (FOI) Report

D), ORAL

Valproate Semisodium
 (Valproate
 Semisodium) C
 Citalopram
 Hydrobromide
 (Citalopram
 Hydrobromide) C
 Estrogens Conjugated
 (Estrogens
 Conjugated) C

Date:10/19/04ISR Number: 4480533-2Report Type:Expedited (15-DaCompany Report #2004075203
 Age:38 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Pulmonary Embolism	Foreign	Neurontin			
Other		Thrombophlebitis Pelvic	Health	(Gabapentin)	PS		ORAL
600 MG (300							
MG, 2 IN 1		Vein	Professional				
D), ORAL							
5 MG (1 D),				Risperidone			
ORAL				(Risperidone)	SS		ORAL
40 DROP (20							
DROP 2 IN				Diazepam	SS		ORAL
1D)ORAL							
2 MG , 2 IN I				Hydromorphone			
DAY, ORAL				Hydrochloride	SS		ORAL

Date:10/19/04ISR Number: 4480537-XReport Type:Expedited (15-DaCompany Report #2004075544
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other 1800 MG (600 MG, 3 IN 1 D), ORAL		Cranial Nerve Disorder Diplopia Eye Pain Gaze Palsy Visual Disturbance	Foreign Health Professional	Gabapentin (Gabapentin) Lamotrigine (Lamotrigine) Acetylsalicylic Acid (Acetylsalicylic Acid) Atenolol (Atenolol) Ramipril (Ramipril) Simvastatin (Simvastatin)	PS C C C C C		ORAL

Date:10/19/04ISR Number: 4480617-9Report Type:Expedited (15-DaCompany Report #2004076131
Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 3000 MG, ORAL		Bone Marrow Depression Bone Marrow Toxicity	Foreign Health Professional	Neurontin (Tablets) (Gabapentin) Metamizole Sodium (Metamizole Sodium) Azathioprine (Azathioprine)	PS SS SS		ORAL ORAL ORAL

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Date:10/19/04
 ISR Number: 4480656-8
 Report Type:Expedited (15-DaCompany Report #2004067531
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anger	Consumer	Neurontin			
Hospitalization -		Anxiety	Health	(Gabapentin)	PS		ORAL
600 MG (300		Arthralgia	Professional				
Initial or Prolonged		Asthenia					
MG, 2 IN 1		Atherosclerosis		Benadryl			
Disability		Back Pain		(Diphenhydramine)	SS		ORAL
D), ORAL		Brain Oedema		Oxycodone			
Other		Burning Sensation		Hydrochloride			
ORAL		Cardio-Respiratory Arrest		(Oxycodone			
		Cardiomegaly		Hydrochloride)	SS		ORAL
1-3 TABLETS		Coma					
Q12H, ORAL		Coronary Artery		Hydrocodone			
		Atherosclerosis		(Hydrocodone)	SS		ORAL
1-2 EVERY 4-6		Discomfort					
HOURS (10		Drug Ineffective					
MG), ORAL		Drug Interaction		Zolpidem Tartrate			
		Erectile Dysfunction		(Zolpidem Tartrate)	SS		ORAL
10 MG (10 MG,		Excoriation					
1 IN 1 D),		Feeling Abnormal					
ORAL		Haematemesis		Citalopram			
		Hepatic Steatosis		Hydrobromide			
		Hypertensive Heart		(Citalopram			
80 MG (80 MG,		Disease		Hydrobromide)	SS		ORAL
1 IN 1 D),		Hypoaesthesia					
ORAL		Insomnia					
		Mood Swings		Amitriptyline			
		Multiple Drug Overdose		Hydrobromide			

Accidental	(Amitriptyline	
Muscle Spasms	Hydrobromide)	C
Muscle Spasticity	Fentanyl (Fentanyl)	C
Muscle Tightness	Cyclobenzaprine	
Nerve Root Lesion	Hydrochloride(Cyclob	
Pain	enzaprine	
Paraesthesia	Hydrochloride)	C
Pulmonary Oedema	Bupropion	
Radiculopathy	Hydrochloride(Buprop	
Renal Cyst	ion Hydrochloride)	C
Respiratory Depression	Quinine Sulfate	
Snoring	(Quinine Sulfate)	C
Spinal Myelogram Abnormal	Sertraline	
Ventricular Hypertrophy	Hydrochloride(Sertra	
	line Hydrochloride)	C
	Trazodone	
	(Trazodone)	C
	Hydroxyzine Embonate	
	(Hydroxyzine	
	Embonate)	C

Date:10/19/04ISR Number: 4480658-1Report Type:Expedited (15-DaCompany Report #2004075210

Age: Gender: I/FU:I

Outcome	PT
Hospitalization -	Alcoholism
Initial or Prolonged	Belligerence
Other	Cardiac Arrest
	Coma
	Drug Abuser

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Marital Problem Personality Change Road Traffic Accident					
600 MG (100 MG, 6 IN 1 D), ORAL		Suicidal Ideation Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Methamphetamine (Methamphetamine)	SS		
				Ethanol (Ethanol)	SS		
				Olanzapine (Olanzapine)	C		
				Lorazepam (Lorazepam)	C		
				Quetiapine Fumarate (Quetiapine Fumarate)	C		
				Clonazepam (Clonazepam)	C		
				Escitalopram (Escitalopram)	C		

Date:10/19/04ISR Number: 4480665-9Report Type:Expedited (15-DaCompany Report #2004075462

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Ineffective Grand Mal Convulsion Partial Seizures	Health Professional	Neurontin (Gabapentin)	PS		
				Dilantin Suspension (Phenytoin Sodium)	SS		
				Valproate Semisodium (Valproate Semisodium)	SS		
				Lamotrigine (Lamotrigine)	SS		
				Phenobarbital (Phenobarbital)	SS		
				Oxcarbazepine (Oxcarbazepine)	SS		
				Topiramate			

(Topiramate) C
Carbamazepine C
(Carbamazepine) C

Date:10/19/04ISR Number: 4480708-2Report Type:Expedited (15-DaCompany Report #2004010780
Age:69 YR Gender:Male I/FU:F

Outcome PT
Disability Actinic Keratosis
Other Blood Cholesterol
Increased
Blood Creatinine
Increased
Blood Glucose Increased
Blood Potassium Increased
Blood Sodium Increased
Bundle Branch Block Left
Difficulty In Walking
Diverticulum

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Emotional Disorder Fatigue Feeling Abnormal	Report Source				
10 MG (DAILY)		Feeling Cold Flushing	Consumer	Lipitor (Atorvastatin)	PS		
300 MG, ORAL		Hyperhidrosis Hypersensitivity		Neurontin (Gabapentin)	SS		ORAL
(DAILY), ORAL		Impaired Work Ability Injury Insomnia Joint Crepitation		Hyzaar (Hydrochlorothiazide , Losartan Potassium)	SS		ORAL
		Kyphosis Low Density Lipoprotein Increased Mental Disorder Neck Injury Nervousness Pain In Extremity Polymyositis Prostatic Specific Antigen Increased Rectal Polyp Sinus Headache Spinal Column Stenosis Visual Acuity Reduced		Influenza Vaccine (Influenza Vaccine) All Other Therapeutic Products (All Other Therapeutic Products) Vicodin (Hydrocodone Bitartrate, Paracetamol) Acetylsalicylic Acid (Acetylsalicylic Acid) Amlodipine Besilate (Amlodipine Besilate)	C C C C C		

Date:10/19/04ISR Number: 4480711-2Report Type:Expedited (15-DaCompany Report #DSA_70449_2004
Age:49 YR Gender:Not SpecifiI/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death DF PO		Multiple Drug Overdose	Literature	Methadone	PS		ORAL
DF PO			Health	Gabapentin	SS		ORAL
DF PO			Professional	Methocarbamol	SS		ORAL

Date:10/19/04ISR Number: 4480742-2Report Type:Expedited (15-DaCompany Report #DSA_70444-_2004
Age:38 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Arrest	Literature	Methadone	PS		ORAL
DF PO							
		Respiratory Arrest	Health	Fluoxetine	SS		ORAL
DF PO							
			Professional	Gabapentin	SS		ORAL
DF PO							

Date:10/19/04ISR Number: 4480896-8Report Type:Expedited (15-DaCompany Report #HQWYE542005OCT04
Age:49 YR Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Intentional Misuse	Literature	Methocarbamol (Methocarbamol, Tablet, 500 Mg)	PS		ORAL
ORAL							
				Gabapentin (Gabapentin,)	SS		ORAL
ORAL							
				Methadone (Methadone,)	SS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/19/04ISR Number: 4482224-0Report Type:Direct
 Age:73 YR Gender:Male I/FU:I

Company Report #CTU 229990

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	900 3XDAILY			Neurontin 300 Mg	PS		ORAL
		Pancreatic Carcinoma					
							ORAL

Date:10/19/04ISR Number: 4483520-3Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 229971

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300 MG HS			Gabapentin 300 Mg	PS		ORAL
		Cardiac Flutter					
		Dyspnoea					
				Oxycodone/Apap	C		
				Trazodone	C		
				Paroxetine	C		
				Propoxyphene/Apap	C		
				Phenytoin	C		
				Asa	C		

Date:10/20/04ISR Number: 4482089-7Report Type:Expedited (15-DaCompany Report #2004076849
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG (300 MG, 3 IN 1 D), UNKNOWN			Neurontin (Gabapentin)	PS		
		Fall	Foreign Health Professional Company Representative				
				Telmisartan (Telmisartan)	C		
				Loprazolam Mesilate (Loprazolam Mesilate)	C		

Calcium Carbonate
 (Calcium Carbonate) C
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C
 Fluoxetine
 Hydrochloride
 (Fluoxetine
 Hydrochloride) C
 Alendronate Sodium
 (Alendronate Sodium) C

Date:10/20/04ISR Number: 4483477-5Report Type:Expedited (15-DaCompany Report #2004055008
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia Pain In Extremity Surgery	Consumer Health Professional	Neurontin (Gabapentin) Hydrocortisone (Hydrocortisone)	PS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/20/04ISR Number: 4483486-6Report Type:Expedited (15-DaCompany Report #2004075594
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abasia	Consumer	Neurontin			
		Movement Disorder		(Gabapentin)	PS		ORAL
		Trance					
200 MG (100							
MG, 2 IN 1							
D), ORAL							

Date:10/21/04ISR Number: 4481522-4Report Type:Expedited (15-DaCompany Report #2004078685
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Foreign	Neurontin			
		Epilepsy	Health	(Gabapentin)	PS		
			Professional	Valproate Sodium			
				(Valproate Sodium)	SS		
				All Other			
				Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	C		

Date:10/21/04ISR Number: 4484177-8Report Type:Expedited (15-DaCompany Report #2004AL000810
 Age:50 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Literature	Chlorpromazine			
		Intentional Misuse		Hydrochloride Oral			
				Concentrate Usp,			
				100mg/Ml (Alpharma)	PS	Alpharma	ORAL
PO				Quetiapine	SS		ORAL
PO				Gabapentin	SS		ORAL
PO							

Date:10/21/04ISR Number: 4485529-2Report Type:Expedited (15-DaCompany Report #8007578
Age:99 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Asthenia	Health	Keppra	PS		
500 MG 2/D		Coordination Abnormal Dyspnoea Dysstasia Feeling Abnormal Medication Error	Professional	Neurontin	SS		

Date:10/21/04ISR Number: 4485934-4Report Type:Expedited (15-DaCompany Report #HQWYE542005OCT04
Age:49 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Intentional Misuse	Literature	Methocarbamol (Methocarbamol, Tablet, 500 Mg)	PS		ORAL
ORAL				Gabapentin (Gabapentin,)	SS		ORAL
ORAL				Methadone (Methadone,)	SS		ORAL
ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/21/04ISR Number: 4486066-1Report Type:Expedited (15-DaCompany Report #2004060002

Age:90 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour Circulatory Collapse	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
400 MG (200 Disability MG, 2 IN 1 Other D),ORAL		Creatinine Renal Clearance Decreased	Professional				
		Fall		Tolterodine (Tolterodine)	C		
		Flatulence		Carbamazepine (Carbamazepine)	C		
		Loss Of Consciousness		Paracetamol (Paracetamol)	C		
				Frumil (Amiloride Hydrochloride, Furosemide)	C		
				Nulytely (Macrogol, Potassium Chloride, Sodium Bicarbonate, Sodium Chloride)	C		
				Lekovit Ca (Calcium Carbonate, Colecalciferol)	C		
				Amoxicillin (Amoxicilline)	C		
				Ciprofloxacin (Ciprofloxacin)	C		
				Tramadol (Tramadol)	C		

Date:10/21/04ISR Number: 4486665-7Report Type:Expedited (15-DaCompany Report #2004066878

Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 900 MG (300 MG, 3 IN 1		Apathy Heart Valve Insufficiency	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Major Depression	Professional				

Myocardial Infarction

D), ORAL

Post Procedural
Complication

Atenolol (Atenolol)	C
Simvastatin (Simvastatin)	C
Fluoxetine (Fluoxetine)	C
Clopidogrel Sulfate (Clopidogrel Sulfate)	C
Mirtazapine (Mirtazapine)	C
Alendronate Sodium (Alendronate Sodium)	C
Tolterodine L-Tartrate (Tolterodine L-Tartrate)	C
Ramipril (Ramipril)	C

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Freedom Of Information (FOI) Report

Date:10/21/04ISR Number: 4486667-0Report Type:Expedited (15-DaCompany Report #2004010780
Age:69 YR Gender:Male I/FU:F

Outcome	PT
Disability	Actinic Keratosis
Other	Aldolase Increased
	Anhedonia
	Arthralgia
	Asthenia
	Blood Cholesterol Increased
	Blood Glucose Increased
	Blood Potassium Increased
	Blood Sodium Increased
	Blood Urine Present
	Bundle Branch Block Left
	C-Reactive Protein Increased
	Crepitations
	Difficulty In Walking
	Diverticulum
	Emotional Distress
	Fatigue
	Feeling Cold
	Flushing
	Gait Disturbance
	Head Injury
	Heart Rate Increased
	Hyperhidrosis
	Impaired Work Ability
	Injury
	Insomnia
	Joint Range Of Motion Decreased
	Kyphosis
	Limb Injury
	Low Density Lipoprotein Increased
	Mental Disorder
	Muscle Atrophy
	Muscle Spasms
	Muscular Weakness
	Myalgia
	Myopathy
	Myositis
	Neck Injury
	Nervousness

Neuralgia
Neuropathy
Pain
Pain In Extremity
Ph Urine Decreased
Polymyalgia
Polymyositis
Prostatic Specific
Antigen Increased
Protein Urine
Rectal Polyp
Red Blood Cell
Sedimentation Rate
Increased
Sinus Headache

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Spinal Column Stenosis Tenderness Visual Acuity Reduced	Report Source	Product	Role	Manufacturer	Route
10 MG (DAILY)		Weight Decreased	Consumer	Lipitor (Atorvastatin)	PS		
300 MG, ORAL				Neurontin (Gabapentin)	SS		ORAL
(DAILY), ORAL				Hyzaar (Hydrochlorothiazide , Losartan Potassium)	SS		ORAL
				Influenza Vaccine (Influenza Vaccine)	C		
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Amlodipine Besilate (Amlodipine Besilate)	C		

Date:10/21/04ISR Number: 4486746-8Report Type:Expedited (15-DaCompany Report #2004076852

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hyperpyrexia	Foreign	Neurontin			
Other		Treatment Noncompliance	Health	(Gabapentin)	PS		ORAL
ORAL			Professional				

Date:10/22/04ISR Number: 4484804-5Report Type:Expedited (15-DaCompany Report #2004AL000916

Age:35 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Tramadol			

	Completed Suicide	Hydrochloride		
	Multiple Drug Overdose	Tablets, 50 Mg		
PO		(Purepac)	PS	ORAL
PO		Citalopram	SS	ORAL
PO		Gabapentin	SS	ORAL

Date:10/25/04ISR Number: 4484286-3Report Type:Expedited (15-DaCompany Report #SE-JNJFOC-20041005449
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Drug Exposure During		Topamax	PS		
INTRA-UTERINE						
Initial or Prolonged	Pregnancy		Neurontin	SS		
INTRA-UTERINE						
Congenital Anomaly	Ventricular Septal Defect		Temesta	SS		
INTRA-UTERINE						

Date:10/25/04ISR Number: 4484344-3Report Type:Expedited (15-DaCompany Report #GB-BRISTOL-MYERS SQUIBB COMPANY-12732475
Age:80 YR Gender:Male I/FU:I

Outcome	PT
Death	Atrial Fibrillation
Other	Blood Creatinine
	Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Blood Ph Increased				
		Blood Potassium Decreased				
		Blood Sodium Decreased				
		Cardiomegaly	Irbesartan	PS	Bristol-Myers Squibb	
		Diarrhoea			Company	
		Myocardial Infarction	Morphine	SS		
		Neutrophil Count Increased	Feldene	SS		
		Oliguria	Celebrex	SS		
		Pulmonary Oedema	Neurontin	SS		
		Respiratory Alkalosis	Prozac	SS		
		Troponin Increased				
		Ventricular Fibrillation				
		Vomiting				
		White Blood Cell Count Increased				

Date:10/25/04ISR Number: 4485315-3Report Type:Direct Company Report #CTU 230358
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Other		Dysarthria	Neurontin	PS		
		Dyskinesia	Famvir	SS		
		Nausea				

Date:10/25/04ISR Number: 4485420-1Report Type:Direct Company Report #CTU 230410
 Age: Gender:Male I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				
Other		Condition Aggravated	Neurontin 300mg			
		Pain	4x1day	PS		ORAL
300MG PO QD						

Date:10/25/04ISR Number: 4487614-8Report Type:Expedited (15-DaCompany Report #2004077254
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose		PT				

Other	Abnormal Dreams	Consumer	Neurontin (Gabapentin)	PS	ORAL
ORAL	Crying				
	Hypoaesthesia		Fluoxetine		
	Lyme Disease		Hydrochloride	C	
	Pain		Montelukast Sodium	C	
			Beclometasone		
			Dipropionate	C	
			Fluticasone		
			Propionate	C	

Date:10/25/04ISR Number: 4487629-XReport Type:Expedited (15-DaCompany Report #2004077855

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Deformity	Consumer	Neurontin			
ORAL		Pain		(Gabapentin)	PS		ORAL
		Polytraumatism					
		Suicide Attempt					
		Unevaluable Event					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/25/04ISR Number: 4487632-XReport Type:Expedited (15-DaCompany Report #2004079228

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Ineffective Knee Operation	Consumer	Neurontin (Gabapentin)	PS		
900 MG (300 MG , 3 IN 1 D)		Pain Pain In Extremity					
		Shoulder Operation Spinal Fusion Surgery		Valdecoxib (Valdecoxib) Rofecoxib (Rofecoxib)	SS SS		

Date:10/25/04ISR Number: 4487804-4Report Type:Expedited (15-DaCompany Report #2004077261

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Other		Drug Toxicity Sudden Cardiac Death	Consumer	Neurontin (Gabepentin)	PS		
900 MG (300 MG,3 IN 1 D)				All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:10/25/04ISR Number: 4488197-9Report Type:Expedited (15-DaCompany Report #2004076841

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Condition Aggravated Mental Disorder	Consumer	Neurontin (Gapabentin)	PS		
1800 MG (600 MG, 3 IN 1 D)		Suicidal Ideation					

Date:10/25/04ISR Number: 4488231-6Report Type:Expedited (15-DaCompany Report #2004067531
Age:42 YR Gender:Male I/FU:F

Outcome	PT
Death	Accidental Death
Hospitalization -	Anger
Initial or Prolonged	Anxiety
Disability	Arthralgia
Other	Atherosclerosis
	Brain Oedema
	Burning Sensation
	Cardio-Respiratory Arrest
	Cardiomegaly
	Coma
	Coronary Artery
	Atherosclerosis
	Decreased Activity
	Drug Abuser
	Drug Ineffective
	Drug Interaction
	Drug Level Above
	Therapeutic
	Drug Screen Positive
	Erectile Dysfunction

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, 2 IN 1 D), ORAL		Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Professional				
			Benadryl (Diphenhydramine)	SS		ORAL
			Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL
1-3 TABLETS Q12H, ORAL						
1-2 EVERY 4-6 HOURS (10 MG), ORAL			Hydrocodone (Hydrocodone)	SS		ORAL
10 MG (10 MG, 1 IN 1 D), ORAL			Zolpidem Tartrate (Zolpidem Tartrate)	SS		ORAL
80 MG (80 MG, 1 IN 1 D), ORAL			Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		ORAL
			Amitriptyline Hydrochloride (Amitriptyline			

Hydrochloride)	C
Fentanyl (Fentanyl)	C
Cyclobenzaprine	
Hydrochloride	
(Cyclobenzaprine	
Hydrochloride)	C
Bupropion	
Hydrochloride	
(Bupropion	
Hydrochloride)	C
Quinine Sulfate	
(Quinine Sulfate)	C
Sertraline	
Hydrochloride	
(Sertraline	
Hydrochloride)	C
Trazodone	
(Trazodone)	C
Hydroxyzine Embonate	
(Hydroxyzine	
Embonate)	C

Date:10/25/04ISR Number: 4488733-2Report Type:Expedited (15-DaCompany Report #2004077798
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Crying	Foreign	Gabapentin (Tablets)			
		Fear	Consumer	(Gabapentin)	PS		ORAL
ORAL		Hallucination					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/25/04ISR Number: 4488747-2Report Type:Expedited (15-DaCompany Report #2004076847
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Zoloft (Sertraline) Dantrolene Sodium (Dantrolene Sodium)	PS SS C		

Date:10/25/04ISR Number: 4489384-6Report Type:Expedited (15-DaCompany Report #KII-2004-0013636
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Acidosis	Study Health Professional	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		ORAL
Other		Alanine Aminotransferase Increased	Other				
ORAL		Aspartate Aminotransferase Increased		Acetaminophen W/Hydrocodone Bitartrate			
ORAL		Aspiration		(Paracetamol, Hydrocodone	SS		ORAL
ORAL		Blood Bicarbonate Decreased		Neurontin (Gabapentin)	SS		ORAL
ORAL		Blood Calcium Decreased					
ORAL		Blood Creatine Phosphokinase Increased		Tetracycline (Tetracycline)	SS		ORAL
ORAL		Blood Creatine Phosphokinase Mb Increased		Benzodiazepine (Derivatives)	SS		
ORAL		Blood Creatinine Increased					
ORAL		Blood Glucose Increased					
ORAL		Blood Ph Decreased					
ORAL		Blood Potassium Decreased					
ORAL		Blood Potassium Increased					
ORAL		Body Temperature Increased					
ORAL		Depressed Level Of					

Consciousness
Drug Screen Positive
Haematocrit Abnormal
Inappropriate Affect
Intentional Misuse
Lipase Increased
Multiple Drug Overdose
Pco2 Increased
Pneumonia
Rhonchi
Sinus Tachycardia
Snoring
Suicide Attempt
Troponin Increased
Vomiting
Wheezing
White Blood Cell Count
Increased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/25/04ISR Number: 4489958-2Report Type:Expedited (15-DaCompany Report #2004075607

Age:94 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 50 MG (50 MG Initial or Prolonged , 1 IN 1 D), Other ORAL	Activities Of Daily Living Impaired Aphagia	Consumer	Zoloft (Sertraline)	PS		ORAL
ORAL	Blood Pressure Increased Dysphagia		Neurontin (Gabapentin)	SS		ORAL
ORAL	Dyspnoea Mobility Decreased		Olanzapine (Olanzapine)	SS		ORAL
ORAL	Pulse Abnormal Unevaluable Event Weight Decreased		Tolterodine L-Tartrate (Tolterodine L-Tartrate)	SS		ORAL
			Rofecoxib (Rofecoxib)	SS		
			Atenolol (Atenolol)	C		
			Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotinamide, Senna Fruit (Senna Fruit)	C C		
			Donepezil Hydrochloride (Donepezil Hydrochloride)	C		
			All Other Therapeutic Products (All Other Therapeutic Products)	C		
			Hemorid For Women Cream (Aloe Vera, Paraffin, Liquid, Phenylephrine Hydrochloride,	C		

Date:10/26/04ISR Number: 4488301-2Report Type:Expedited (15-DaCompany Report #2004065315
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2400	MG (800)	Health Professional	Neurontin (Gabapentin)	PS		
Other MG, 3 IN 1		Drug Withdrawal Syndrome	Company				
D), PLACENTAL		Neonatal	Representative				
TRANSPLACENTAL	PLACENTAL	Maternal Use Of Illicit Drugs		Cannabis (Cannabis)	SS		
TRANSPLACENTAL	PLACENTAL	Premature Baby		Hydrocodone (Hydrocodone)	SS		

Date:10/26/04ISR Number: 4488302-4Report Type:Expedited (15-DaCompany Report #2004077846
 Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1200 MG (600)		Blood Pressure Increased Blood Sodium Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
MG, 2 IN 1		Convulsion					
D), ORAL		Drug Ineffective					

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Freedom Of Information (FOI) Report

Valdecoxib
 (Valdecoxib) SS
 Cyclobenzaprine
 (Cyclobenzaprine) SS
 Levetiracetam
 (Levetiracetam) C
 Alprazolam
 (Alprazolam) C
 Dextropropoxyphene
 (Dextropropoxyphene) C

Date:10/26/04ISR Number: 4488339-5Report Type:Expedited (15-DaCompany Report #2004057241
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin			
Other		Constipation		(Gabapentin)	PS		
		Dizziness		Ibuprofen			
		Drug Ineffective		(Ibuprofen)	SS		
		Epistaxis		Acetylsalicylic Acid			
		Fatigue		(Acetylsalicylic Acid)	SS		
		Hypoaesthesia		Naproxen Sodium			
		Insomnia		(Naproxen Sodium)	SS		
		Intervertebral Disc Compression		Vicodin (Hydrocodone Bitartrate,			
		Localised Infection		Paracetamol)	SS		
		Pollakiuria		Codeine (Codeine)	SS		
		Renal Pain		Rosiglitazone Maleate			
		Sinus Disorder		(Rosiglitazone Maleate)	C		
		Treatment Noncompliance		Furosemide			
				(Furosemide)	C		
				Metolazone			
				(Metolazone)	C		
				Esomeprazole			
				(Esomeprazole)	C		
				Potassium Chloride			
				(Potassium Chloride)	C		
				Rofecoxib			
				(Rofecoxib)	C		
				Metolazone			
				(Metolazone)	C		

Date:10/26/04ISR Number: 4488417-0Report Type:Expedited (15-DaCompany Report #2004079234
Age:83 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Abnormal Behaviour	Consumer	Neurontin			
Other		Delirium		(Gabapentin)	PS		
		Dizziness					
		Fall					
		Somnolence					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/26/04ISR Number: 4489407-4Report Type:Expedited (15-DaCompany Report #2004081132

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bone Marrow Depression	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:10/26/04ISR Number: 4489875-8Report Type:Expedited (15-DaCompany Report #2004077781

Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Agranulocytosis Asthenia	Health Professional Company Representative	Gabapentin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL		Leukopenia Pyrexia		Omeprazole (Omeprazole) Nifedipine (Nifedipine) Paracetamol (Paracetamol) Fentanyl (Fentanyl)	C C C C		

Date:10/26/04ISR Number: 4489878-3Report Type:Expedited (15-DaCompany Report #2004077869

Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Eye Oedema Generalised Oedema	Foreign Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
900 MG (900 MG, 1 IN 1 D), ORAL		Headache Insomnia					

Myelodysplastic Syndrome
Pain In Extremity
Refractory Anaemia

Tramadol
Hydrochloride
(Tramadol
Hydrochloride)

C

Date:10/27/04ISR Number: 4489643-7Report Type:Expedited (15-DaCompany Report #2004-UK-01033UK
Age:80 YR Gender:Male I/FU:I

Outcome PT
Death Atrial Fibrillation
Blood Creatinine
Increased
Blood Potassium Decreased
Blood Sodium Decreased
Body Temperature
Increased
Cardiomegaly
Diarrhoea
Drug Ineffective
Electrolyte Imbalance
Myocardial Infarction
Myocardial Ischaemia
Neutrophil Count
Increased
Oliguria

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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 Oedema Respiratory Alkalosis Vasodilatation					
		Ventricular Fibrillation Vomiting White Blood Cell Count Increased	Foreign	Morphine (0015/0122/0208/0157)) (Morphine)	PS		
INTRAVENOUS	IV			Feldene (Piroxicam) Celebrex (Celecoxib) Neurontin (Gabapentin) Irbesartan (Irbesartan) Prozac (Fluoxetine Hydrochloride)	SS SS SS SS SS		

Date:10/27/04ISR Number: 4490002-1Report Type:Expedited (15-DaCompany Report #2004079364
Age:80 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other			Myocardial Ischaemia	Foreign Health Professional	Gabapentin (Gabapentin) Tamsulosin (Tamsulosn)	PS SS		ORAL
	400 MCG (400 MCG, 1 IN 1 D), ORAL							

Date:10/27/04ISR Number: 4490175-0Report Type:Expedited (15-DaCompany Report #2004076197
Age:60 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Abdominal Pain Abnormal Dreams Alcoholism Amnesia	Consumer	Lipitor (Atorvastatin) Neurontin (Gabapentin)	PS SS		
	300 MG (100							

Anxiety
Arthritis
Back Pain
Chest Discomfort
Depression
Dizziness
Drug Ineffective
Electromyogram Abnormal
Fatigue
Headache
Impaired Driving Ability
Impaired Work Ability
Migraine
Muscle Disorder
Muscular Weakness
Musculoskeletal Stiffness
Neck Pain
Nerve Conduction Studies
Abnormal
Photosensitivity Reaction
Sleep Disorder
Spinal Osteoarthritis
Tinnitus
Transient Ischaemic
Attack

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/27/04ISR Number: 4490186-5Report Type:Expedited (15-DaCompany Report #2004075222

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG (15 Other MG, 4 IN 1 D), ORAL	Amnesia Complex Regional Pain Syndrome Depression Disease Recurrence Drug Effect Decreased Excitability Gastritis	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
60 MG (15 MG , 4 IN 1 D), ORAL	Headache Insomnia Mania Pain In Extremity Rheumatoid Arthritis Sedation Suicidal Ideation White Blood Cell Count Increased		Prednisone (Prednisone)	SS		ORAL
			Vicodin (Hydrocodone Bitartrate, Paracetamol) Escitalopram (Escitalopram) Provella-14 (Estrogens Conjugated, Mdedoxyprogesterone Acetate) Nifedipine (Nfiedipine) Propranolol Hydrochloride (Propranolol Hydrochloride) Cetirizine Hydrochlrodie (Cetirizine Hydrochloride) Lansoprazole (Lansoprazole)	SS SS C C C C		

Date:10/27/04ISR Number: 4490189-0Report Type:Expedited (15-DaCompany Report #2004079252

Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Alopecia Aphonia	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other 400 MG (100 MG, 4 IN 1 D), ORAL		Drug Effect Increased Drug Hypersensitivity					
ORAL		Drug Ineffective Dysgeusia		Gabapentin (Gabapentin)	SS		ORAL
		Dysphagia Feeling Abnormal Multiple Allergies Oral Intake Reduced Pain Of Skin Pharmaceutical Product Complaint Pharyngolaryngeal Pain Transient Ischaemic Attack Viral Load Increased		Tetracycline (Tetracycline) Ciprofloxacin (Ciprofloxacin) Diazepam (Diazepam) Estrogens Conjugated (Estrogens Conjugated) Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride) All Other Therapeutic Products (All Other Therapeutic	SS SS C C C		

Freedom Of Information (FOI) Report

Products) C
 Hydrochlorothiazide
 (Hydrochlorothiazide
) C

Date:10/27/04ISR Number: 4490212-3Report Type:Expedited (15-DaCompany Report #2004079287

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (200 Other MG, 1 IN 1 D)		Drug Ineffective Liver Function Test Abnormal	Foreign Consumer	Dilantin Suspension (Phenytoin Sodium)	PS		
400 MG		Panic Attack Renal Disorder		Neurontin (Gabapentin)	SS		
		Surgery White Blood Cell Count Abnormal					

Date:10/27/04ISR Number: 4490877-6Report Type:Expedited (15-DaCompany Report #2004068343

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 800 MG (400 MG,2 IN 1 D), ORAL		Confusional State Movement Disorder Neuropathy Pain	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
		Pyrexia Somnolence		Atorvastatin Calcium (Atorvastatin Calcium) Bromazepam (Bromazepam) Amitriptyline Hydrochloride (Amitriptyline Hydrochloride) Carbamazepine	C C C		

(Carbamazepine) C
Diclofenac Sodium C
(Diclofenac Sodium) C

Date:10/28/04ISR Number: 4488528-XReport Type:Direct
Age:49 YR Gender:Male I/FU:I

Company Report #CTU 230669

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased Hypoaesthesia		Gabapentin 400 Mg -Ivax	PS	Ivax	
400 MG QID		Pain In Extremity Paraesthesia Pharmaceutical Product Complaint		Eskalith Paxil Topamax Premarin Cydominzapine	C C C C C		

Date:10/28/04ISR Number: 4490060-4Report Type:Expedited (15-DaCompany Report #2004082345
Age: Gender:Female I/FU:I

Outcome PT
Other Cutaneous Lupus
Erythematosis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Fibromyalgia Rheumatoid Arthritis Sjogren'S Syndrome	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin)	PS		
				Zoloft (Sertraline)	SS		
				Methotrexate (Methotrexate)	SS		
				Infliximab (Infliximab)	SS		
				Hydroxychloroquine Phosphate (Hydroxychloroquine Phosphate)	SS		
				Tizanidine Hydrochloride (Tizanidine Hydrochloride)	SS		
				Morphine Sulfate (Morphine Sulfate)	C		

Date:10/28/04ISR Number: 4490252-4Report Type:Expedited (15-DaCompany Report #2004081489
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin (Gabapentin)	PS		ORAL
Life-Threatening ORAL		Blood Pressure					
Disability ORAL		Fluctuation Cardiovascular Disorder		Rofecoxib (Rofecoxib)	SS		ORAL
		Cerebrovascular Disorder Injury Oedema Overdose Pain					

Date:10/28/04ISR Number: 4490624-8Report Type:Expedited (15-DaCompany Report #2004079435
Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Delirium	Foreign	Neurontin			
Initial or Prolonged		Depressed Level Of	Health	(Gabapentin)	PS		
300 MG							
Other		Consciousness	Professional	Tramadol			
		Drug Interaction	Company	Hydrochloride			
			Representative	(Tramadol			
				Hydrochloride)	SS		
150 MG,							

Date:10/29/04ISR Number: 4491929-7Report Type:Direct Company Report #CTU 230793
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disorientation		Gabapentin	PS		ORAL
300MG	TID			300mg			
		Dizziness					
ORAL		Somnolence					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:10/29/04ISR Number: 4492036-XReport Type:Expedited (15-DaCompany Report #2004-BP-10224YA

Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Myocardial Ischaemia	Health	Flomax Mr	PS		ORAL
0.4 MG			Professional Other	Gabapentin (Gabapentin)	SS		

Date:10/29/04ISR Number: 4492496-4Report Type:Expedited (15-DaCompany Report #2004081303

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Knee Arthroplasty	Consumer	Neurontin (Gabapentin)	PS		
				Zoloft (Sertraline)	SS		
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		
				Rofecoxib (Rofecoxib)	SS		
				All Other Therapeutic Products	SS		

Date:10/29/04ISR Number: 4492680-XReport Type:Expedited (15-DaCompany Report #2004062853

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Blood Alkaline Phosphatase Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		Blood Calcium Increased	Professional				
200 MG (100 MG, 2 IN 1 D), ORAL		C-Reactive Protein Increased Gait Disturbance Mammary Duct Ectasia Pancreatic Atrophy Renal Cyst		Clomipramine (Clomipramine)	C		

Date:10/29/04ISR Number: 4492688-4Report Type:Expedited (15-DaCompany Report #2004080565

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pancreatitis Haemorrhagic	Foreign Health	Neurontin(Gabapentin)	PS		ORAL
ORAL			Professional Company Representative				

Date:10/29/04ISR Number: 4492691-4Report Type:Expedited (15-DaCompany Report #2004081142

Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Drug Ineffective Epilepsy	Foreign Health	Neurontin(Gabapentin)	PS		
1800 MG(600 MG, 3 IN 1 D)		Hyponatraemia	Professional Company Representative	Pravastatin Sodium(Pravastatin Sodium)	C		
		Leukopenia Neutropenia		Phenobarbital Sodium			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Phenobarbital Sodium) C
 Risperidone (Risperidone) C
 Noctran 10 (Acepromazine, Acepromatazine, Clorazepate Dipotassium) C

Date:10/29/04ISR Number: 4493041-XReport Type:Expedited (15-DaCompany Report #2004080275
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Renal Cyst	Consumer	Neurontin (Gabapentin)	PS		
200 MG (100 MG, 2 IN 1 D)							
				Donepezil Hydrochloride (Donepezil Hydrochloride)	SS		

Date:10/29/04ISR Number: 4493047-0Report Type:Expedited (15-DaCompany Report #2004062834
 Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthropathy Back Disorder	Consumer Health Professional	Neurontin (Gabapentin)	PS		
				Ibuprofen (Ibuprofen)	SS		
				Paracetamol (Paracetamol)	C		

Date:10/29/04ISR Number: 4534118-XReport Type:Direct Company Report #USP 56912
 Age:10 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

SOLUTION Medication Error Zarontin PS Pfizer Us Pharm
SOLUTION Neurontin 250 Mg/5ml SS Parke-Davis

Date:11/01/04ISR Number: 4491237-4Report Type:Direct Company Report #CTU 230849
Age:74 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 900 MG /DAY Initial or Prolonged	Generalised Oedema		Gabapentin	PS		

Date:11/01/04ISR Number: 4491388-4Report Type:Direct Company Report #CTU 230914
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration 600 MG QID PO	Convulsion Pharmaceutical Product Complaint		Neurontin	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/01/04ISR Number: 4491532-9Report Type:Expedited (15-DaCompany Report #2004080583
Age:88 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - ORAL		Fall Hypotension	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
Initial or Prolonged		Renal Failure Acute	Professional	Amiodarone (Amiodarone)	C		
				Carbocisteine (Carbocisteine)	C		
				Tiotropium Bromide (Tiotropium Bromide)	C		
				Candesartan (Candesartan)	C		
				Salbutamol (Salbutamol)	C		
				Beclometasone (Beclometasone)	C		
				Furosemide (Furosemide)	C		

Date:11/01/04ISR Number: 4491570-6Report Type:Expedited (15-DaCompany Report #2004080543
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cerebellar Ataxia Dizziness	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL			Professional	Calcium Carbonate	C		

Date:11/01/04ISR Number: 4491581-0Report Type:Expedited (15-DaCompany Report #2004081134
Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Alkalosis	Foreign	Neurontin			

Other ORAL	Blood Creatinine Increased	Health Professional	(Gabapentin)	PS	ORAL
	Blood Potassium Decreased		Feldene (Piroxicam)	SS	
	Blood Sodium Decreased		Celecoxib (Celecoxib)	SS	ORAL
ORAL	Body Temperature Increased		Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	SS	
	Cardiomegaly		Irbesartan (Irbesartan)	SS	
	Electrolyte Imbalance		Morphine (Morphine)	SS	
	Myocardial Infarction		Prochlorperazine	C	
	Myocardial Ischaemia				
	Neutrophil Count Increased				
	Pulmonary Oedema				
	Ventricular Fibrillation				
	Vomiting				
	White Blood Cell Count Increased				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/01/04ISR Number: 4491764-XReport Type:Expedited (15-DaCompany Report #2004084209
 Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diaphragmatic Paralysis	Health Professional	Neurontin (Gabapentin)	PS		
UNKNOWN	600 MG (300		Company				
MG, 2 IN 1 D)			Representative				
UNKNOWN							

Date:11/01/04ISR Number: 4491766-3Report Type:Expedited (15-DaCompany Report #2004080553
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Accidental Overdose	Health Professional	Neurontin (Gabapentin)	PS		
Initial or Prolonged		Dialysis					
UNKNOWN	900 MG (300						
Other		Mental Status Changes					
MG,3 IN 1 D),							
UNKNOWN							
UNKNOWN	UNKNOWN			Fentanyl (Fentanyl) (Fentanyl)	SS		
				Mycophenolate Mofetil (Mycophenolate Mofetil)	C		
				Tacrolimus (Tacrolimus)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Prednisone (Prednisone)	C		
				Calcium Acetate (Calcium Acetate)	C		
				Labetalol (Labetalol)	C		
				Isosorbide Dinitrate (Isosorbide			

Dinitrate) C
 Enalapril Maleate C
 (Enalapril Maleate)
 Paroxetine
 Hydrochloride
 (Paroxetine)
 Hydrochloride) C
 Clonazepam C
 (Clonazepam)
 Valproate Semisodium C
 (Valproate
 Semisodium) C

Date:11/01/04ISR Number: 4492086-3Report Type:Expedited (15-DaCompany Report #2004081136
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetes Mellitus	Consumer	Neurontin			
		Fibromyalgia		(Gabapentin)	PS		
		Multiple Sclerosis		Fentanyl (Fentanyl)	SS		
				Oxycocet (Oxycodone Hydrochloride, Paracetamol)	SS		
				Lidocaine			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Lidocaine) SS
 Baclofen (Baclofen) SS
 Amantadine
 Hydrochloride
 (Amantadine
 Hydrochloride) SS
 All Other
 Therapeutic Prouducts
 (All Other
 Therapeutic
 Products) SS
 Insulin (Insulin) SS

Date:11/01/04ISR Number: 4492153-4Report Type:Expedited (15-DaCompany Report #2004080073
 Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Nausea Therapy Non-Responder	Consumer	Neurontin (Gabapentin) All Other Therapeutic Products (All Other Therapeutic Products)	PS C		

Date:11/01/04ISR Number: 4492162-5Report Type:Expedited (15-DaCompany Report #2004077846
 Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1200 MG (600 MG, 2 IN 1 D), ORAL		Blood Pressure Increased Blood Sodium Decreased Condition Aggravated Convulsion Drug Ineffective Headache Neck Pain	Consumer	Neurontin (Gabapentin) Valdecoxib (Valdecoxib) Cyclobenzaprine (Cyclobenzaprine) Levetiracetam (Levetiracetam)	PS SS SS C		ORAL

Alprazolam
(Alprazolam) C
Dextropropoxyphene
(Dextropropoxyphene) C

Date:11/01/04ISR Number: 4492176-5Report Type:Expedited (15-DaCompany Report #2004080083

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Nausea	Consumer	Neurontin (Gabapentin)	PS		

Date:11/02/04ISR Number: 4492857-3Report Type:Direct Company Report #CTU 231044

Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - 600 MG/900 MG Initial or Prolonged QAM/QPM ORAL		Syncope		Gabapentin 300 Mg	PS		ORAL

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Freedom Of Information (FOI) Report

Date:11/02/04ISR Number: 4492926-8Report Type:Expedited (15-DaCompany Report #9042

Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Atrial Fibrillation		Morphine	PS		
		Blood Ph Increased		Piroxicam	SS		
		Blood Potassium Decreased		Celecoxib	SS		
		Blood Sodium Decreased		Gabapentin	SS		
		Body Temperature Increased		Irbesartan	SS		
		Cardiomegaly		Fluoxetine	SS		
		Diarrhoea					
		Electrolyte Imbalance					
		Heart Rate Increased					
		Myocardial Infarction					
		Myocardial Ischaemia					
		Neutrophil Count Increased					
		Oliguria					
		Pulmonary Oedema					
		Respiratory Alkalosis					
		Respiratory Rate Increased					
		Resuscitation					
		Ventricular Fibrillation					
		Vomiting					
		White Blood Cell Count Increased					

Date:11/02/04ISR Number: 4493715-0Report Type:Expedited (15-DaCompany Report #2004081169

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Tolerance Decreased	Health	Dilantin Suspension			
Other		Grand Mal Convulsion	Professional	(Phenytoin Sodium)	PS		
				Neurontin			
				(Gabapentin)	SS		
				Lamotrigine			
				(Lamotrigine)	SS		
				Phenobarbital			
				(Phenobarbital)	SS		
				Carbamazepine			
				(Carbamazepine)	SS		

Date:11/02/04ISR Number: 4494218-XReport Type:Expedited (15-DaCompany Report #A02200403140
Age:63 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Alanine Aminotransferase
Initial or Prolonged	Increased
	Aspartate
	Aminotransferase
	Increased
	C-Reactive Protein
	Increased
	Epstein-Barr Virus
	Infection
	Haematoma

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Freedom Of Information (FOI) Report

Dose	Duration	Neutropenia Normochromic Normocytic Anaemia	Report Source	Product	Role	Manufacturer	Route
10 MG OD ORAL		Thrombocytopenic Purpura	Foreign	Stilnox - (Zolpidem)	PS		ORAL
ORAL		White Blood Cell Count Decreased	Health Professional Other	Aprovel - (Irbesartan) - Tablet	SS		ORAL
ORAL	14 WK			Veinamitol - Troloxerutin) -	SS		ORAL
150 MG BID, ORAL	7 WK			Topalgic - (Tramadol Hydrochloride)	SS		ORAL
ORAL	14 WK			Lamaline - (Paracetamol/Opium/C afeine)	SS		ORAL
300 MG TID, ORAL	2 YR			Neurontin - (Gabapentin) - Capsule - 300 Mg	SS		ORAL
TOPICAL	TOPICAL			Daktarin - (Miconazole Nitrate) - Ointment	SS		
1 UNIT OD ORAL				Cokenzen - (Candesartan+Hydroch lorothiazide)	SS		ORAL
0.5 MG BID ORAL				Rivotril - (Clonazepam) - Drops - 0.1 Mg	SS		ORAL

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (300 Other MG, 3 IN 1 D)	Delirium Depressed Level Of Consciousness	Foreign Health Professional	Neurontin (Gabapentin)	PS		
150 MG	Drug Interaction Hallucination, Visual	Company Representative	Tramadol Hydrochloride (Tramadol Hydrochloride)	SS		
			Omeprazole(Omeprazol e)	SS		
			Digoxin (Digoxin)	C		
			Spasfon (Phloroglucinol, Trimethylphlorogluci nol)	C		
			Carbosylane (Dimeticone)	C		
			Macrogol (Macrogol)	C		

Outcome	PT
Hospitalization - Initial or Prolonged	Haematoma Thrombocytopenia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Thrombocytopenic Purpura

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Health Professional	Aprovel Tabs	PS	Bristol-Myers Squibb Company	ORAL
102 DAY			Veinamitol	SS		ORAL
			Stilnox	SS		ORAL
55 DAY			Topalgic	SS		ORAL
102 DAY			Lamaline	SS		ORAL
30 MON			Neurontin	SS		ORAL

Date:11/03/04ISR Number: 4494619-XXReport Type:Expedited (15-DaCompany Report #2004082912

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hyperchlorhydria	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
(100 MG),		Mood Altered	Professional				
ORAL		Neutropenia					
				Ranitide Hydrochloride (Ranitidine Hydrochloride)	SS		
150 MG (150							
MG, 1 IN 1 D)							
				Clozapine (Clozapine)	SS		ORAL
(162.5 MG),							
ORAL							
				Lamotrigine (Lamotrigine)	SS		ORAL
ORAL							
				Risperidone (Risperidone)	SS		ORAL
(100 MG, 1 IN							

1 D), ORAL

Date:11/03/04ISR Number: 4494622-XReport Type:Expedited (15-DaCompany Report #2004081134
Age:80 YR Gender:Male I/FU:F

Outcome	PT
Death	Alkalosis
Other	Atrial Fibrillation
	Blood Creatinine
	Increased
	Blood Potassium Decreased
	Blood Sodium Decreased
	Body Temperature
	Increased
	Cardiomegaly
	Diarrhoea
	Drug Ineffective
	Heart Rate Increased
	Myocardial Infarction
	Myocardial Ischaemia
	Neutrophil Count
	Increased
	Oliguria
	Pulmonary Oedema
	Respiratory Rate
	Increased
	Ventricular Fibrillation
	Vomiting
	White Blood Cell Count

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Increased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
ORAL		Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Professional	Feldene (Piroxicam) Celecoxib (Celecoxib)	SS SS		ORAL
			Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	SS		
			Irbesartan (Irbesartan)	SS		
			Morphine (Morphine)	SS		
			Prochlorperazine	C		

Date:11/03/04ISR Number: 4494687-5Report Type:Expedited (15-DaCompany Report #2004-BP-10189RO
Age:20 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Abuser Multiple Drug Overdose	Literature	Methadone Hcl Tablets Usp, 10 Mg (Methadone Hydrochloride)	PS		
				Sertraline (Sertraline)	SS		
				Gabapentin	SS		

Date:11/03/04ISR Number: 4494732-7Report Type:Expedited (15-DaCompany Report #2004-BP-10183RO
Age:38 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Multiple Drug Overdose	Literature	Methadone Hcl Tablets Usp, 10 Mg (Methadone Hydrochloride)	PS		
				Fluoxetine			

(Fluoxetine
Hydrochloride) SS
Gabapentin
(Gabapentin) SS

Date:11/03/04ISR Number: 4494734-0Report Type:Expedited (15-DaCompany Report #2004-BP-10185RO

Age:49 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Multiple Drug Overdose	Literature	Methadone Hcl Tablets Usp, 10 Mg (Methadone Hydrochloride) Gabapentin (Gabapentin) Methacarbamol (Methocarbamol)	PS SS SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/03/04ISR Number: 4495004-7Report Type:Expedited (15-DaCompany Report #8007778
Age:45 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Acetaminophen W/Hydrocodone Bitartrate	PS		ORAL
PO				Gabapentin	SS		ORAL

Date:11/03/04ISR Number: 4495006-0Report Type:Expedited (15-DaCompany Report #8007777
Age:23 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest Multiple Drug Overdose Respiratory Arrest	Literature Health Professional	Acetaminophen W/Hydrocodone Bitartrate	PS		ORAL
PO				Gabapentin	SS		ORAL

Date:11/04/04ISR Number: 4495510-5Report Type:Expedited (15-DaCompany Report #2004-BP-10504RO
Age:31 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Drug Abuser Intentional Misuse	Literature	Cocaine Hcl Topical Solution, 10% (Cocaine) Gabapentin (Gabapentin) Ethanol (Ethanol)	PS SS SS		

Date:11/04/04ISR Number: 4497408-5Report Type:Expedited (15-DaCompany Report #2004082915
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged 900 MG (300 Other MG, 3 IN 1 D), ORAL ORAL	Back Disorder Body Height Decreased Bursitis Cyst Road Traffic Accident Scoliosis Wheelchair User	Health Professional	Neurontin (Gabapentin) Lipitor (Atorvastatin) Calcium Carbonate 9calcium Carbonate) Centrum Silver (Ascorbic Acid, Calcium, Minerals Nos, Retinol, Tocopheryl Acetate,	PS SS C C	ORAL ORAL
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Date:11/04/04ISR Number: 4497409-7Report Type:Expedited (15-DaCompany Report #2004062831
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		No Adverse Drug Effect	Consumer	Neurontin (Gabapentin) Oxycocet (Oxycodone Hydrochloride, Paracetamol) Paroxetine Hydrochloride	PS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Paroxetine
Hydrochloride) SS
Rofecoxib
(Rofecoxib) SS
Methylphenidate
Hydrochloride
(Methylphenidate
Hydrochloride) SS
Fentanyl (Fentanyl) SS

Date:11/04/04ISR Number: 4497410-3Report Type:Expedited (15-DaCompany Report #2004083706
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (300 MG, 2 IN 1 D), ORAL		Apathy Erythema Fatigue Feeling Abnormal Loss Of Employment Nervousness	Consumer	Neurontin (Gabapentin) Lithium (Lithium) Esomeprazole (Esomeprazole)	PS SS C		ORAL

Date:11/04/04ISR Number: 4497411-5Report Type:Expedited (15-DaCompany Report #2004047157
Age:14 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (200 MG, 3 IN 1 D), ORAL		Drug Effect Decreased Emotional Disorder Growth Accelerated Medication Error Pharmaceutical Product Complaint	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin			
ORAL		Completed Suicide		(Gabapentin)	PS		ORAL
		Fear					
		Pain					
		Suicidal Ideation					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abdominal Pain Upper	Consumer	Neurontin			
		Chemical Abuser		(Gabapentin)	PS		
		Suicide Attempt		All Other			
		Weight Decreased		Non-Therapeutic			
				Products (All Other			
				Therapeutic			
				Products)	SS		
RESPIRATORY							
(INHALATION)	INHALATION			Diazepam (Diazepam)	C		
				All Other			
				Therapeutic Products			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(All Other
Therapeutic
Products) C

Date:11/04/04ISR Number: 4497415-2Report Type:Expedited (15-DaCompany Report #2004063259

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oesophageal Ulcer	Consumer	Neurontin (Gabapentin)	PS		
				Motrin (Ibuprofen)	SS		
				Rofecoxib (Rofecoxib)	SS		

Date:11/04/04ISR Number: 4497551-0Report Type:Expedited (15-DaCompany Report #2004060020

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Atrial Fibrillation	Foreign	Neurontin			
Hospitalization -		Blood Creatinine	Health	(Gabapentin)	PS		ORAL
300 MG (300							
Initial or Prolonged		Increased	Professional				
MG, 1 IN 1							
D), ORAL		Cardiac Failure	Company				
		Congestive	Representative	Antibiotics			
		Cerebral Atrophy		(Antibiotics)	C		
		Embolic Cerebral		Acetylsalicylic Acid			
		Infarction		(Acetylsalicylic			
		Somnolence		Acid)	C		
				Furosemide			
				(Furosemide)	C		
				Ace Inhibitor Nos			
				(Ace Inhibitor Nos)	C		

Date:11/04/04ISR Number: 4497556-XReport Type:Expedited (15-DaCompany Report #2004082916

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other	Cardiac Pacemaker	Foreign	Neurontin (Tablets)		
2400 MG (2400	Insertion	Consumer	(Gabapentin)	PS	ORAL
MG, 1 IN 1		Health			
D), ORAL		Professional			
		Distributor			
		Other			

Date:11/04/04ISR Number: 4498329-4Report Type:Expedited (15-DaCompany Report #2004082914
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Foreign	Gabapentin			
Other		Cholestasis	Literature	(Gabapentin)	PS		
		Hepatic Steatosis	Health	Initard (Insulin,			
		Hepatitis	Professional	Insulin Injection,			
				Isophane)	C		
				Metformin			
				(Metformin)	C		
				Amitriptyline			
				(Amitriptyline)	C		
				Dihydrocodeine			
				(Dihydrocodeine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ramipril (Ramipril) C

Date:11/05/04ISR Number: 4493784-8Report Type:Expedited (15-DaCompany Report #US-ABBOTT-04P-163-0279130-00
Age:23 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Cardio-Respiratory Arrest	Hydrocodone/Acetamin ophen	PS		ORAL
				Gabapentin	SS		ORAL

Date:11/05/04ISR Number: 4493785-XReport Type:Expedited (15-DaCompany Report #US-ABBOTT-04P-163-0279137-00
Age:45 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide	Hydrocodone/Acetamin ophen	PS		ORAL
				Gabapentin	SS		ORAL

Date:11/05/04ISR Number: 4493799-XReport Type:Expedited (15-DaCompany Report #US-ABBOTT-04P-163-0279300-00
Age:44 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide	Valproic Acid	PS		ORAL
				Levothyroxine	SS		ORAL
				Gabapentin	SS		ORAL

Date:11/05/04ISR Number: 4497891-5Report Type:Expedited (15-DaCompany Report #2004083497
Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG (100 Other MG, 2 IN 1			Dizziness				

D), ORAL

Drug Ineffective

Dyspnoea

Hemiparesis

Insomnia

Nocturnal Dyspnoea

Transient Ischaemic

Attack

Omeprazole

(Omeprazole)

C

Furosemide

(Furosemide)

C

Date:11/05/04ISR Number: 4497896-4Report Type:Expedited (15-DaCompany Report #2004084794

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Diabetes Mellitus	Consumer	Neurontin			
Other		Double Vessel Bypass Graft		(Gabapentin)	PS		
		Limb Operation		Zoloft (Sertraline)	SS		
		Neuropathy Peripheral		Procardia			
				(Nifedipine)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/05/04ISR Number: 4497899-XReport Type:Expedited (15-DaCompany Report #2004085957

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
		General Physical Health		(Gabapentin)	PS		
		Deterioration		Dilantin Suspension			
		Hepatitis C		(Phenytoin Sodium)	SS		
		Ill-Defined Disorder					

Date:11/05/04ISR Number: 4498418-4Report Type:Expedited (15-DaCompany Report #2004080565

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pancreatitis Haemorrhagic	Foreign Health	Neurontin			
			Professional Company Representative	(Gabapentin)	PS		ORAL
ORAL							

Date:11/05/04ISR Number: 4498419-6Report Type:Expedited (15-DaCompany Report #2004083821

Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Blood Glucose Fluctuation	Foreign Health	Neurontin			
1200 MG (1		Diabetes Mellitus		(Gabapentin)	PS		ORAL
D), ORAL		Dizziness	Professional				
		Drug Ineffective		Lisinopril			
20 MG (1 D)		Drug Interaction		(Lisinopril)	SS		
		Fall		Hydrochlorothiazide			
12.5 MG (1 D)		Grand Mal Convulsion		(Hydrochlorothiazide			
		Hypoglycaemia)	SS		
5 MG (1 D)		Status Epilepticus		Baclofen (Baclofen)	SS		
				Pantoprazole Sodium			

20 MG (1 D)

(Pantoprazole Sodium) SS

SUBCUTANEOUS 1 AMPULE (2

Interferon Beta (Interferon Beta) SS

D),

SUBCUTANEOUS

Amitriptyline Hydrochloride (Amitriptyline Hydrochloride) C
Carbamazepine (Carbamazepine) C
Zopiclone (Zopiclone) C

Date:11/05/04ISR Number: 4498556-6Report Type:Expedited (15-DaCompany Report #2004051151

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crohn'S Disease	Consumer	Neurontin (Tablets)			
1800 MG (600		Drug Effect Decreased	Health	(Gabapentin)	PS		ORAL
MG, 3 IN 1		Drug Ineffective	Professional				
D), ORAL		Osteosclerosis					
				Celecoxib (Celecoxib)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Rofecoxib
(Rofecoxib) C

Date:11/08/04ISR Number: 4495946-2Report Type:Expedited (15-DaCompany Report #FR-JNJFOC-20041008398
Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Drug Interaction	Topalgic	PS		
OROPHARINGEAL							
Initial or Prolonged			Epstein-Barr Virus	Veinamitol	SS		
OROPHARINGEAL							
			Antibody Positive	Stilnox	SS		
OROPHARINGEAL							
			Laboratory Test Abnormal	Aprovel	SS		
OROPHARINGEAL							
			Thrombocytopenia	Lamaline	SS		
OROPHARINGEAL							
			Thrombocytopenic Purpura	Lamaline	SS		
OROPHARINGEAL							
				Lamaline	SS		
OROPHARINGEAL							
				Lamaline	SS		
OROPHARINGEAL							
				Lamaline	SS		
OROPHARINGEAL							
				Neurontin	SS		
OROPHARINGEAL							

Date:11/08/04ISR Number: 4496613-1Report Type:Direct Company Report #CTU 231427
Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Nausea Pruritus Vomiting	Gabapentin	PS		

Date:11/08/04ISR Number: 4497585-6Report Type:Expedited (15-DaCompany Report #GB-JNJFOC-20041007637
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other
 OROPHARINGEAL 1.5 MG bd - Haemoglobin Decreased Risperdal PS
 Neutropenia
 100 MG daily, White Blood Cell Count
 per source Decreased
 documents 21 DAY
 OROPHARINGEAL Clozapine SS
 OROPHARINGEAL Ranitidine SS
 Gabapentin SS
 OROPHARINGEAL Lamotrigine SS
 OROPHARINGEAL

Date:11/08/04ISR Number: 4498622-5Report Type:Expedited (15-DaCompany Report #2004076723
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Condition Aggravated Depression Emotional Distress Mental Disorder Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		

Date:11/08/04ISR Number: 4498753-XReport Type:Expedited (15-DaCompany Report #2004082946
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Hallucination, Auditory Self Mutilation	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/08/04ISR Number: 4498828-5Report Type:Expedited (15-DaCompany Report #2004084796
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (200 Other MG, 1 IN 1 D)		Drug Ineffective Grand Mal Convulsion	Consumer	Neurontin (Gabapentin)	PS		
				Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		

Date:11/08/04ISR Number: 4498865-0Report Type:Expedited (15-DaCompany Report #2004083712
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 800 MG (400 MG, 2 IN 1 D), ORAL		Cognitive Disorder Delusion Transient Ischaemic Attack	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:11/08/04ISR Number: 4498941-2Report Type:Expedited (15-DaCompany Report #2004086532
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Disability	Consumer	Neurontin (Gabapentin)	PS		
				Alprazolam (Alprazolam)	SS		
				Clonazepam (Clonazepam)	SS		
				Methadone (Methadone)	SS		
				Ibuprofen (Ibuprofen)	SS		

Date:11/08/04ISR Number: 4498943-6Report Type:Expedited (15-DaCompany Report #2004085913

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Jittery Irritability	Health Professional	Neurontin (Gabapentin)	PS		
1600 MG (800 MG, 2 IN 1 D)		Treatment Noncompliance	Company Representative	Antihypertensives (Antihypertensives) Celecoxib (Celecoxib)	C C		

Date:11/08/04ISR Number: 4498968-0Report Type:Expedited (15-DaCompany Report #2004085912

Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Partial Seizures Pharmaceutical Product	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
2400 MG (600 MG, 4 IN 1 D)		Complaint Vomiting		Metoclopramide (Metoclopramide)	C		
ORAL							

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Freedom Of Information (FOI) Report

Budesonide
 (Budesonide) C
 Fexofenadine
 Hydrochloride
 (Fexofenadine
 Hydrochloride) C

Date:11/08/04ISR Number: 4498975-8Report Type:Expedited (15-DaCompany Report #2004075239
 Age:71 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Chest Pain Coma	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
ORAL Other		Nausea Renal Failure	Professional	Pantoprazole (Pantoprazole)	C		
				Magnesium (Magnesium)	C		
				Oxycocet (Oxycodone Hydrochloride, Paracetamol)	C		
				Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
				Losartan Potassium (Losartan Potassium)	C		
				Furosemide (Furosemide)	C		
				Isosorbide (Isosorbide)	C		
				Potassium (Potassium)	C		
				Simvastatin (Simvastatin)	C		
				Escitalopram (Escitalopram)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride)	C		
				All Other			

Therapeutic Products	
(All Other	
Therapeutic	
Products)	C
Methylphenidate	
Hydrochloride	
(Methylphenidate	
Hydrochloride)	C
Seretide Mite	
(Fluticasone	
Propionate,	
Salmeterol	
Xinafoate)	C
Oxygen (Oxygen)	C
Fentanyl (Fentanyl)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/09/04ISR Number: 4498030-7Report Type:Direct
Age:69 YR Gender:Female I/FU:I

Company Report #CTU 231603

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
	100 MCG QD	2	MON		Levothyroxine 100 Mcg	PS		
	100 MG PO BID	2	MON	Pharmaceutical Product	Gabapentin	SS		ORAL
				Complaint				

Date:11/09/04ISR Number: 4500889-1Report Type:Expedited (15-DaCompany Report #2004042614
Age: Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged				Consumer	Nardil (Phenelzine Sulfate)	PS		
					Neurontin (Gabapentin)	SS		
					Lorazepam (Lorazepam)	SS		
					Clozapine (Clozapine)	C		
					Lamotrigine (Lamotrigine)	C		
				Pharmaceutical Product				
				Complaint				

Date:11/09/04ISR Number: 4502706-2Report Type:Expedited (15-DaCompany Report #9042
Age:80 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death					Morphine	PS		
					Piroxicam	SS		
					Celecoxib	SS		
					Gabapentin	SS		
					Irbesartan	SS		
					Fluoxetine	SS		
				Blood Creatinine Increased				
				Blood Ph Increased				
				Blood Potassium Decreased				
				Blood Sodium Decreased				
				Cardiomegaly				
				Diarrhoea				
				Electrolyte Imbalance				
				Myocardial Infarction				

Neutrophil Count
 Increased
 Oliguria
 Pulmonary Oedema
 Respiratory Alkalosis
 Ventricular Fibrillation
 Vomiting
 White Blood Cell Count
 Increased

Date:11/10/04ISR Number: 4497545-5Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040908820
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest		Ultram	PS		
OROPHARINGEAL		Completed Suicide		Citalopram	SS		
OROPHARINGEAL		Convulsion		Metoclopramide	SS		
		Hypotension		Omeprazole	SS		
		Hypoxic Encephalopathy		Gabapentin	SS		
OROPHARINGEAL		Intentional Misuse					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/10/04ISR Number: 4498777-2Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 231708

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypoaesthesia		Neurontin 100 Mg Pfizer	PS	Pfizer	
2 X DAILY							
				Neurontin 300 Mg Pfizer	SS	Pfizer	
2 X DAILY							

Date:11/10/04ISR Number: 4502351-9Report Type:Expedited (15-DaCompany Report #2004085994
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Blood Alkaline Phosphatase Increased	Foreign Health	Gabapentin (Tablets) (Gabapentin)	PS		ORAL
2000 MG, ORAL							
Other		Bronchiectasis Gamma-Glutamyltransferase Increased	Professional Company Representative	Lidocaine (Lidocaine) (Lidocaine)	SS		
INTRAVENOUS	INTRAVENOUS	Pyrexia Respiratory Tract		Carbamazepine (Carbamazepine)	SS		
UNKNOWN	UNKNOWN	Infection		Clonazepam (Clonazepam)	SS		
UNKNOWN	UNKNOWN			Tramadol Hydrochloride (Tramadol Hydrochloride)	SS		
UNKNOWN	UNKNOWN						

Date:11/10/04ISR Number: 4502376-3Report Type:Expedited (15-DaCompany Report #2004086032
 Age:8 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Oedema Peripheral	Foreign	Gabapentin			

750 MG (250
MG, 3 IN 1
D), ORAL

Pain	Health	(Gabapentin)	PS	ORAL
Radial Nerve Palsy	Professional			

Chlorhexidine (Chlorhexidine)	C
Codeine Phosphate (Codeine Phosphate)	C
Lactulose (Lactulose)	C
Senna (Senna)	C
Diclofenac (Diclofenac)	C
Granisetron (Granisetron)	C
Paracetamol (Paracetamol)	C
Morphine Sulfate (Morphine Sulfate)	C
Lorazepam (Lorazepam)	C
Bethanechol Chloride (Bethanechol Chloride)	C
Fentanyl (Fentanyl)	C
Cefuroxime (Cefuroxime)	C
Propofol (Propofol)	C
Cyclizine	

Freedom Of Information (FOI) Report

(Cyclizine) C
 Other Therapeutic
 Products (All Other
 Therapeutic
 Products) C
 Antineoplastic
 Agents
 (Antineoplastic
 Agents) C
 Vincristine
 (Vincristine) C
 Cyclophosphamide
 (Cyclophosphamide) C
 Carboplatin
 (Carboplatin) C
 Dactinomycin
 (Dactinomycin) C
 Etoposide
 (Etoposide) C

Date:11/10/04ISR Number: 4502668-8Report Type:Expedited (15-DaCompany Report #A02200403207
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Aggression Agitation Anxiety	Health Professional	Stilnox (Zolpidem)-Tablet-10 Mg	PS		ORAL
10 MG OD, ORAL		Cerebral Disorder					
		Cerebral Haemangioma Delirium Lacrimation Increased Nightmare		Laroxyl- (Amitripyline Hydrochloride)- Tablet - 25 Mg	SS		ORAL
25MG OD, ORAL	9 DAY	Suicidal Ideation Trismus Weight Decreased		Athymil - (Mianserin Hydrochloride) - Tablet - 30 Mg	SS		ORAL
30 MG OD, ORAL				Neurontin - (Gabapentin) - Tablet - 600 Mg	SS		ORAL
600 MG TID,							

ORAL 53 DAY

Duragesic -
(Fentanyl)- Patch-
25 Mcg SS

TRANSDERMAL 25 MCG OD,

TRANSDERMAL 2 DAY

Actiskenan-
(Morphine Sulfate)-
Capsule- 10 Mg SS

10 MG >QID;

ORAL

ORAL 9 DAY

Forlax (Macrogol) C
Minidril
(Levonorgestrel/Ethi
nylestradiol) C

Date:11/12/04ISR Number: 4499743-3Report Type:Expedited (15-DaCompany Report #200418611US

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Chest Pain	Health	Lantus	PS	Aventis	
Initial or Prolonged		Dizziness	Professional			Pharmaceuticals Inc.	
SUBCUTANEOUS	Dose unit:						

units

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dose: 20/125

Glucotrol Xl	SS	ORAL
Metformin	SS	ORAL
Lotensin Hct	SS	ORAL
Plavix	SS	ORAL
Crestor	SS	ORAL
Celebrex	SS	ORAL
Ecotrin	SS	ORAL
Isosorbide	SS	ORAL
Neurontin	SS	ORAL
Ssri	SS	ORAL
Protonix	C	ORAL

Date:11/12/04ISR Number: 4499793-7Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040908887
 Age:70 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Alanine Aminotransferase	Health	Tylox	PS		
OROPHARINGEAL	Increased	Professional	Valproic Acid	SS		
OROPHARINGEAL	Aspartate		Simvastatin	SS		
OROPHARINGEAL	Aminotransferase		Lorazepam	SS		
	Increased		Gabapentin	SS		
	Blood Creatine		Diclofenac/Misoprost			
	Phosphokinase Increased		ol	SS		
	Blood Creatine		Topiramate	SS		
	Phosphokinase Mb					
	Increased					
	Blood Creatinine					
	Increased					
	Blood Glucose Decreased					
	Blood Pressure Systolic					
	Increased					
	Brain Oedema					
	Coma					
	Completed Suicide					
	Coronary Artery Disease					
	Haematemesis					
	Hyperkalaemia					
	Hypertensive					
	Cardiomyopathy					
	Hypertensive Heart					

Disease
Intentional Misuse
Nephrosclerosis
Pulmonary Oedema

Date:11/12/04ISR Number: 4500774-5Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 231809

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated		Gabapentin 300mg			
Other		Feeding Disorder		Ivax	PS	Ivax	ORAL
TWO FOUR		Pharmaceutical Product					
TIMES/DAY		Complaint					
ORAL		Speech Disorder		Tegrotol	C		
		Trigeminal Neuralgia		Prednisone	C		
				Travatan	C		
				Voltaren	C		
				Avonex	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/12/04ISR Number: 4502861-4Report Type:Expedited (15-DaCompany Report #2004085157
Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Quadriplegia	Health Professional	Dilantin Suspension (Phenytoin Sodium) Neurontin (Gabapentin)	PS SS		

Date:11/12/04ISR Number: 4502968-1Report Type:Expedited (15-DaCompany Report #2004083706
Age:62 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Apathy Erythema	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 2 IN 1 D), ORAL		Fatigue Feeling Abnormal Impaired Work Ability Loss Of Employment Nervousness Personality Change Self Esteem Decreased Somnolence		Lithium (Lithium) Esomeprazole (Esomeprazole)	C C		

Date:11/15/04ISR Number: 4500607-7Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20040502832
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Apnoea		Duragesic	PS		
TRANSDERMAL		Blood Urine Present		Duragesic	SS		
TRANSDERMAL		Cardio-Respiratory Arrest		Duragesic	SS		
TRANSDERMAL		Coma Completed Suicide		Effexor Neurontin	SS SS		

TRANSDERMAL

Dizziness	Duragesic	SS
Fall	Atenolol	C
Foreign Body Aspiration	Lorazepam	C
Gastric Perforation	Flexeril	C
Haematemesis	Bextra	C
Headache	Temazepam	C
Hepatic Congestion	Topamax	C
Injury	Ambien	C
Insomnia	Zyban	C
Medication Error		
Muscle Spasms		
Narcotic Intoxication		
Overdose		
Pain		
Pain In Extremity		
Petechiae		
Pharmaceutical Product Complaint		
Precancerous Cells		
Present		
Pulmonary Congestion		
Pulse Absent		
Rash		
Ulnar Nerve Injury		
Ulnar Nerve Palsy		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/15/04ISR Number: 4503882-8Report Type:Expedited (15-DaCompany Report #2004090318
Age:59 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2400 MG (800 MG, 3 IN 1 D)	Blood Bilirubin Abnormal Gamma-Glutamyltransferase Increased Multiple Sclerosis Oedema Peripheral	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Paracetamol (Paracetamol) Atorvastatin Calcium (Atorvastatin Calcium) Propofan (Caffeine, Carbasalate Calcium, Chlorphenamine Maleate, Dextropropoxyphene,	PS C C C		

Date:11/15/04ISR Number: 4504354-7Report Type:Expedited (15-DaCompany Report #2004086847
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 100 MG (100 MG, 1 IN 1 D), ORAL	Abasia Anuria Blood Pressure Increased Confusional State Depression Diverticulitis Dysarthria Feeling Abnormal Metrorrhagia Nervous System Disorder Oedema Peripheral Overdose Pain In Extremity Suicidal Ideation Uterine Leiomyoma Vaginal Disorder	Consumer	Neurontin (Gabapentin) Zolmitriptan (Zolmitriptan) Sumatriptan Succinate (Sumatriptan Succinate) Axotal (Old Form) (Butalbital, Caffeine, Paracetamol) Vicodin (Hydrocodone Bitartrate,	PS C C C		ORAL

Vision Blurred
White Blood Cell Count
Increased

Paracetamol) C
Alprazolam
(Alprazolam) C
Paroxetine
Hydrochloride
(Paroxetine
Hydrochloride) C
Promethazine
(Promethazine) C

Date:11/16/04ISR Number: 4502235-6Report Type:Expedited (15-DaCompany Report #AU-JNJFOC-20041101687

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged	Confusional State Drug Interaction		Tramadol Hydrochloride	PS		
OROPHARINGEAL						
	Hyponatraemia		Gabapentin	SS		
OROPHARINGEAL						
			Trimethoprim	SS		
OROPHARINGEAL						
			Bekunis	SS		
OROPHARINGEAL						
			Bekunis	SS		
OROPHARINGEAL						
			Bekunis	SS		
OROPHARINGEAL						
			Bekunis	SS		

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FDA - Adverse Event Reporting System (AERS)

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Diltiazem
 Hydrochloride C
 Hydrochlorothiazide
 Irbesartan C
 Hydrochlorothiazide
 Irbesartan C
 Sofolol C
 Aspirin Compound C
 Aspirin Compound C
 Aspirin Compound C
 Coloxyl With Senna C
 Coloxyl With Senna C

Date:11/16/04ISR Number: 4503100-0Report Type:Direct Company Report #CTU 232099
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea Gastrointestinal Disorder Nausea Vomiting		Gabapentin	PS		

Date:11/16/04ISR Number: 4503116-4Report Type:Direct Company Report #CTU 232103
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pain In Extremity		Gabapentin 800 Mg Tab	PS		

Date:11/16/04ISR Number: 4504547-9Report Type:Expedited (15-DaCompany Report #2004084796
 Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 200 MG (200 Other MG, 1 IN 1 D)		Drug Ineffective Feeling Abnormal Grand Mal Convulsion	Consumer	Neurontin (Gabapentin)	PS		

Fluoxetine
Hydrochloride
(Fluoxetine
Hydrochloride) C

Date:11/16/04ISR Number: 4504642-4Report Type:Expedited (15-DaCompany Report #044-0945-M0100204
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG (300	Hepatic Steatosis Hepatitis	Foreign Literature	Neurontin (Gabapentin)	PS		ORAL
Other MG, 3 IN 1		Jaundice Cholestatic	Health Professional				

Human Mixtard
(Insulin Human,
Insulin Human
Injection, Isophane) C
Metformin
(Metformin) C
Amitriptyline
(Amitriptyline) C

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Freedom Of Information (FOI) Report

Dihydrocodeine
 (Didhydrocodeine) C
 Ramipril (Ramipril) C

Date:11/16/04ISR Number: 4505279-3Report Type:Expedited (15-DaCompany Report #2004079435
 Age:82 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG, 3 IN 1 D), ORAL		Bedridden Confusional State Delirium Depressed Level Of Consciousness Drug Interaction Hallucination, Visual	Foreign Health Professional Company Representative	Neurontin (Tablets) (Gabapentin) Tramadol Hydrochloride (Tramadol Hydrochloride)	PS SS		ORAL ORAL
300 MG (150 MG, 2 IN 1 D), ORAL				Digoxin (Digoxin) Omeprazole (Omeprazole) Spasfon (Phloroglucinol, Trimethylphlorglucin ol) Carbosylane (Dimeticone) Macrogol (Macrogol) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Lansoprazole (Lansoprazole) Oxazepam (Oxazepam) Calcium Carbonate (Calcium Carbonate)	C C C C C C C C C C C C		

Date:11/16/04ISR Number: 4505340-3Report Type:Expedited (15-DaCompany Report #2004089117
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Condition Aggravated Dyspnoea Guillain-Barre Syndrome Respiratory Disorder	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:11/16/04ISR Number: 4505345-2Report Type:Expedited (15-DaCompany Report #2004090318
Age:59 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2400 MG (800 Other MG, 3 IN 1 D)	Gamma-Glutamyltransferase Increased Multiple Sclerosis Oedema Peripheral	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Propofan (Caffeine, Carbasalate Calcium, Chlorphenamine	PS		

Hospitalization - Hallucination
 900 MG QD
 Initial or Prolonged Muscle Twitching
 (300 MG TID)
 Somnolence

Gabapentin PS
 Hyoscyamine C
 Sertraline C
 Hctz C
 Terazosin C
 Loratadine C
 Zocor C
 Lortab C
 Insulin Nph C
 Metoprolol C
 Methocarbamol C
 Fosinopril C
 Zantac C

Date:11/17/04ISR Number: 4505086-1Report Type:Direct Company Report #CTU 232248
 Age:66 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 300 MG TID Initial or Prolonged	Muscle Twitching		Gabapentin	PS		
			Fosinopril	C		
			Metformin	C		
			Loratadine	C		
			Psyllium	C		
			Sennakot	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Insulin Nph C

Date:11/17/04ISR Number: 4506444-1Report Type:Expedited (15-DaCompany Report #2004034121
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bedridden	Consumer	Neurontin			
300 MG		Cellulitis		(Gabapentin)	PS		
		Cystitis					
		Depression					
		Diabetes Mellitus					
		Dizziness					
		Lung Neoplasm Malignant					
		Lymphoedema					
		Muscle Spasms					
		Oedema Peripheral					
		Pain					
		Unevaluable Event					
		Vaginal Pain					
		Vomiting					

Date:11/17/04ISR Number: 4506459-3Report Type:Expedited (15-DaCompany Report #2004068966
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Consumer	Neurontin			
		Pain In Extremity	Health	(Gabapentin)	PS		
			Professional	Hydrocortisone			
				(Hydrocortisone)	SS		

Date:11/17/04ISR Number: 4506613-0Report Type:Expedited (15-DaCompany Report #2004085912
Age:37 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Partial Seizures	Consumer	Neurontin (Tablets)			
2400 MG (600		Vomiting		(Gabapentin)	PS		ORAL

D), ORAL

Metoclopramide
(Metoclopramide) C
Budesonide (Budesonide)
e) C
Fexofenadine
Hydrochloride
(Fexofenadine
Hydrochloride) C

Date: 11/17/04 ISR Number: 4506641-5 Report Type: Expedited (15-DaCompany Report #2004014973

Age: Gender: Female I/FU: F

Outcome PT
Disability Abdominal Hernia
Other Brain Damage
Circulatory Collapse
Coma
Diabetes Mellitus
Dysgraphia

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FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Economic Problem					
		Gangrene					
		Medication Error					
		Mental Impairment	Consumer	Zoloft (Sertraline)	PS		
		Nervous System Disorder		Celebrex (Celecoxib)			
		Pain		(Celecoxib)	SS		
		Parosmia		Neurontin			
		Post Procedural		(Gabapentin)	SS		
		Complication		Lipitor			
		Post Procedural Discharge		(Atorvastatin)	SS		
		Postoperative Infection		Zyrtec (Tablets)			
		Reading Disorder		(Cetirizine)	SS		
		Staphylococcal Infection		Glucotrol			
		Umbilical Hernia		(Glipizide)	SS		
		Weight Increased		Valdecoxib			
		Wheelchair User		(Valdecoxib)	C		

Date:11/17/04ISR Number: 4506857-8Report Type:Expedited (15-DaCompany Report #RS004775-USA

Age:52 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Balance Disorder	Consumer	Aciphex			
		Chills		(Rabeprazole)			
		Disorientation		(Rabeprazole Sodium)	PS		ORAL
60 MG, 1 IN 1		Influenza Like Illness					
D, ORAL		Nausea		Provigil (Modafinil)	SS		ORAL
SEE IMAGE		Vomiting		Klonopin			
2.25 MG, 1 IN				(Clonazepam)	SS		ORAL
1 D, ORAL				Neurontin			
1200 MG, 1 IN				(Gabapentin)	SS		ORAL
1 D, ORAL				Hydrochlorothiazide			
100 MG, 1 IN				(Hydrochlorothiazide)	SS		ORAL

1 D, ORAL					
100 MG, 1 IN			Doryx (Doxycycline)	SS	ORAL
1 D, ORAL					
10 MG, 1 IN 1			Lexapro	SS	ORAL
D, ORAL					

Date:11/17/04ISR Number: 4506889-XReport Type:Expedited (15-DaCompany Report #2004089442
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Arthritis	Consumer	Neurontin			
Initial or Prolonged	Breast Cancer		(Gabapentin)	PS		ORAL
ORAL						
Other	Cardiac Arrest		Norvasc (Amlodipine)	SS		
	Cataract		Celebrex (Celecoxib)	SS		
	Chest Pain		Donepezil			
	Dementia Alzheimer'S Type		Hydrochloride			
	Dyspepsia		(Donepezil			
	Feeling Abnormal		Hydrochloride)	SS		
	Kidney Infection					
	Malaise					
	Rotator Cuff Syndrome					
	Urinary Tract Infection					
	Weight Decreased					

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Freedom Of Information (FOI) Report

Date:11/17/04ISR Number: 4506894-3Report Type:Expedited (15-DaCompany Report #2004088763

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis	Consumer	Neurontin			
		Coronary Artery Disease		(Gabapentin)	PS		
		Feeling Abnormal		All Other			
		Hypertension		Therapeutic Products			
		Myocardial Infarction		(All Other			
		Neuropathy Peripheral		Therapeutic			
				Products)	SS		

Date:11/17/04ISR Number: 4506898-0Report Type:Expedited (15-DaCompany Report #2004082358

Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Asthenia	Consumer	Neurontin			
Initial or Prolonged		Balance Disorder		(Gabapentin)	PS		ORAL
100 MG (100							
Other		Carotid Artery Stenosis					
MG, 1 IN 1							
D), ORAL		Cerebellar Atrophy					
		Cerebral Atrophy		Valproate Semisodium			
		Cerebrovascular Accident		(Valproate			
		Clumsiness		Semisodium)	SS		ORAL
1000 MG (500							
MG, 2 IN 1		Condition Aggravated					
D), ORAL		Cystitis Interstitial					
		Demyelination		Topiramate			
		Difficulty In Walking		(Topiramate)	SS		ORAL
30 MG (30							
MG), ORAL		Disturbance In Attention					
		Dizziness		Marvelon			
		Drooling		(Desogestrel,			
		Drug Ineffective		Ethinylestradiol)	C		
		Dysarthria		Amlodipine Besilate			
		Dysgeusia		(Amlodipine			
		Essential Tremor		Besilate)	C		
		Gait Disturbance		Clopidogrel Sulfate			

Hemiparesis	(Clopidogrel	
Hypoaesthesia	Sulfate)	C
Intervertebral Disc	Verapamil	
Protrusion	Hydrochloride	
Malaise	(Verapamil	
Memory Impairment	Hydrochloride)	C
Muscle Twitching	Alprazolam	
Paraesthesia	(Alprazolam)	C
Pineal Neoplasm	Sulbutamol	
Refusal Of Treatment By	(Sulbutamol)	C
Patient	Isosorbide Dinitrate	
Sensation Of Heaviness	(Isosorbide	
Transient Ischaemic	Dinitrate)	C
Attack	Azelastine	
Treatment Noncompliance	Hydrochloride	
Tremor	(Azelastine	
Urine Abnormality	Hydrochloride)	C
Urine Odour Abnormal		
Visual Acuity Reduced		

Freedom Of Information (FOI) Report

Date:11/17/04ISR Number: 4506902-XXReport Type:Expedited (15-DaCompany Report #2004088109
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Asthenia Blood Pressure Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
	Dizziness Feeling Cold Hypoaesthesia Pain In Extremity		All Other Therapeutic Products (All Other Therapeutic Products) Dyazide (Hydrochlorothiazide , Triamterene) Warfarin Sodium (Warfarin Sodium) Omeprazole (Omeprazole)	C C C C C		

Date:11/17/04ISR Number: 4524577-0Report Type:Periodic Company Report #US-JNJFOC-20040500628
 Age:45 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Disability ORAL Other	Aplastic Anaemia Herpetic Stomatitis Pyrexia Sepsis	Health Professional	Levaquin (Levofloxacin) Tablets Temodar (Temozolomide) Capsules	PS SS		ORAL ORAL
, 1 IN 1 DAY, ORAL 360 MG, 1 IN 1 DAY, ORAL			Temodar (Temozolomide) Capsules Temodar (Temozolomide)	SS SS		ORAL ORAL

300 MG, 1 IN	Capsules	SS	ORAL
1 DAY, ORAL			
	Temodar (Temozolomide)		
100 MG, ORAL	Capsules	SS	ORAL
	Temodar (Temozolomide)		
80 MG, ORAL	Capsules	SS	ORAL
	Neurontin (Gabapentin)		
1200 MG, 3 IN	Capsules	SS	ORAL
1 DAY, ORAL			
ORAL	Septra (Bactrim)	SS	ORAL
	Thalidomide (Thalidomide)	SS	
1200 MG, 1 IN			
1 DAY	Heparin (Heparin)	SS	
INJECTION	Famciclovir (Famciclovir)	C	
	Codeine (Codeine)	C	
	Phenobarbital (Phenobarbital)	C	
	Aspirin (Acetylsalicylic Acid) Tablets	C	

Freedom Of Information (FOI) Report

Fragmin	
(Heparin-Fraction,	
Sodium Salt)	C
Cpt-11 (Irinotecan)	C
Bcnu (Carmustine)	C
Taxol (Paclitaxel)	C
Tamiflu	
(Oseltamivir)	C
Ceftriaxone	
(Ceftriaxone)	C

Date:11/18/04ISR Number: 4507310-8Report Type:Expedited (15-DaCompany Report #2004010780
 Age:69 YR Gender:Male I/FU:F

Outcome	PT
Disability	Actinic Keratosis
Other	Aldolase Increased
	Arthralgia
	Blood Cholesterol
	Increased
	Blood Creatinine
	Increased
	Blood Glucose Increased
	Blood Potassium Increased
	Blood Sodium Increased
	Blood Urine Present
	Bundle Branch Block Left
	C-Reactive Protein
	Increased
	Difficulty In Walking
	Diverticulum
	Drug Hypersensitivity
	Emotional Distress
	Fatigue
	Feeling Abnormal
	Feeling Cold
	Flushing
	Gait Disturbance
	Head Injury
	Hyperhidrosis
	Hypertension
	Impaired Work Ability
	Injury
	Insomnia
	Irritability

Joint Crepitation
Joint Range Of Motion
Decreased
Kyphosis
Limb Injury
Low Density Lipoprotein
Increased
Muscle Atrophy
Muscle Spasms
Muscular Weakness
Myalgia
Myopathy
Myositis
Neck Injury
Nervousness

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Neuralgia Neuropathy Pain					
10 MG (DAILY)		Pain In Extremity Polymyositis	Consumer	Lipitor (Atorvastatin)	PS		
300 MG, ORAL		Prostatic Specific Antigen Increased		Neurontin (Gabapentin)	SS		ORAL
DAILY, ORAL		Protein Urine Rectal Polyp Red Blood Cell Sedimentation Rate		Hyzaar (Hydrocholorthiazide , Losartan Potassium)	SS		ORAL
		Increased Sinus Headache Skin Irritation Spinal Column Stenosis Tenderness Urinary Tract Infection		Influenza Vaccine (Influenza Vaccine) All Other Therapeutic Products (All Other Therapeutic Products) Vicodin (Hydrocodine Bitartrate, Paracetamol) Acetylsalicylic Acid (Acetylsalicylic Acid) Amlodipine Besilate (Amlodipine Besilate)	C C C C		

Date:11/18/04ISR Number: 4507394-7Report Type:Expedited (15-DaCompany Report #2004065245
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Glucose Increased Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG (800 MG, 3 IN 1 D), ORAL		Diabetic Neuropathy False Positive Laboratory Result		Venlafaxine			

Insomnia

Hydrochloride
(Venlafaxine
Hydrochloride)

SS

ORAL

150 MG (150

MG, 1 IN 1

D), ORAL

Metformin
(Metformin)

C

Salbutamol
(Salbutamol)

C

Pantoprazole
(Pantoprazole)

C

Phazyme (Diastase,
Pancreatin, Pepsin,
Simeticone)

C

Celecoxib
(Celecoxib)

C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/18/04ISR Number: 4507396-0Report Type:Expedited (15-DaCompany Report #2004068968

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Migraine	Consumer	Neurontin (Tablets) (Gabapentin)	PS		
2400 MG		Osteoarthritis Road Traffic Accident		Muscle Relaxants (Muscle Relaxants)	SS		
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		
				Ibuprofen (Ibuprofen)	SS		
				Paracetamol (Paracetamol)	SS		
				Excedrin P.M. (Diphenhydramine Citrate, Paracetamol)	SS		

Date:11/18/04ISR Number: 4507399-6Report Type:Expedited (15-DaCompany Report #2004061141

Age:72 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Failure Cough	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (1 IN		Dyspnoea	Professional				
1 D), ORAL		Gastric Banding		Co-Diovan (Hydrochlorothiazide , Valsartan)	C		
				Rofecoxib (Rofecoxib)	C		
				Premarin (Estrogens Conjugated)	C		

Date:11/18/04ISR Number: 4512879-3Report Type:Direct

Age: Gender:Female I/FU:I

Company Report #CTU # 232450

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased Inadequate Analgesia		Gabapentin Teva	400mg	PS Teva	ORAL
ONE CAPSULE		Middle Insomnia					
3 TIMES		Neuralgia					
PER DAY		Pharmaceutical Product					
ORAL		Complaint					

Date:11/19/04ISR Number: 4505970-9Report Type:Expedited (15-DaCompany Report #GB-BRISTOL-MYERS SQUIBB COMPANY-12732475
Age:80 YR Gender:Male I/FU:F

Outcome	PT
Death	Atrial Fibrillation
Hospitalization - Initial or Prolonged	Blood Creatinine Increased
Other	Blood Ph Increased Blood Potassium Decreased Blood Sodium Decreased Cardiomegaly Diarrhoea Myocardial Infarction

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Neutrophil Count Increased Oliguria	Report Source	Product	Role	Manufacturer	Route
		Pulmonary Oedema Respiratory Alkalosis Troponin Increased Ventricular Fibrillation Vomiting White Blood Cell Count Increased	Health Professional	Irbesartan Morphine Feldene Celebrex Neurontin Prozac	PS SS SS SS SS SS	Bristol-Myers Squibb Company	

Date:11/19/04ISR Number: 4506900-6Report Type:Expedited (15-DaCompany Report #2004088110
Age: Gender:Female I/FU:I

Outcome Dose Other	Duration	PT Gallbladder Disorder	Report Source Consumer	Product Neurontin (Gabapentin)	Role PS	Manufacturer	Route

Date:11/19/04ISR Number: 4509462-2Report Type:Expedited (15-DaCompany Report #2004064324
Age: Gender:Male I/FU:F

Outcome Dose Death	Duration	PT Completed Suicide	Report Source Consumer	Product Neurontin (Gabapentin)	Role PS	Manufacturer	Route

Date:11/19/04ISR Number: 4509466-XReport Type:Expedited (15-DaCompany Report #001-0945-M0200657
Age:44 YR Gender:Female I/FU:F

Outcome Hospitalization - Initial or Prolonged Disability Other	PT Abdominal Pain Upper Affect Lability Aggression Agitation Anger Arthralgia Arthritis Arthropod Bite Asthenia

Balance Disorder
Cardiovascular Disorder
Carpal Tunnel Syndrome
Change Of Bowel Habit
Chest Pain
Chromaturia
Cognitive Disorder
Constipation
Convulsion
Coordination Abnormal
Deafness
Death Of Friend
Decreased Appetite
Depression
Diarrhoea
Dissociation
Disturbance In Attention
Dizziness
Dry Mouth
Dysarthria

Freedom Of Information (FOI) Report

Dyskinesia
Dyspepsia
Dysphagia
Dyspnoea
Dysuria
Ear Pain
Electrocardiogram
Abnormal
Essential Tremor
Eustachian Tube
Dysfunction
Fall
Fatigue
Fibrocystic Breast
Disease
Flat Affect
Gastrooesophageal Reflux
Disease
Goitre
Granuloma
Hallucination, Visual
Hand Fracture
Head Injury
Headache
Heart Rate Increased
Heart Rate Irregular
Hordeolum
Hot Flush
Hypoaesthesia
Hypothyroidism
Impaired Healing
Imprisonment
Increased Tendency To
Bruise
Increased Upper Airway
Secretion
Insomnia
Intentional Self-Injury
Intervertebral Disc
Degeneration
Intervertebral Disc
Disorder
Irritability
Joint Injury
Joint Sprain
Joint Swelling
Laryngitis

Lumbar Radiculopathy
Memory Impairment
Mouth Breathing
Nasal Discomfort
Nasal Ulcer
Nausea
Neck Injury
Neck Pain
Nystagmus
Onychomadesis
Open Wound
Otitis Media
Pain In Extremity
Paraesthesia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1800 MG (DAILY), ORAL		Pharyngitis Polyneuropathy Rash Generalised				
		Rhinitis Allergic Road Traffic Accident	Consumer Health	Neurontin (Gabapentin)	PS	ORAL
		Sedation				
		Sinus Disorder	Lamotrigine			
		Skin Ulcer	(Lamotrigine)	C		
		Sneezing	Lithium Carbonate			
		Somnolence	(Lithium Carbonate)	C		
		Staring	Clonazepam			
		Stress	(Clonazepam)	C		
		Suicidal Ideation	Methylphenidate			
		Syncope	Hydrochloride			
		Tardive Dyskinesia	(Methylphenidate			
		Tendonitis	Hydrochloride)	C		
		Thirst	Levothyroxine Sodium			
		Tinnitus	(Levothyroxine			
		Tremor	Sodium)	C		
		Urinary Tract Infection	Liothyronine Sodium			
		Vaginal Candidiasis	(Liothyronine			
		Vertigo Positional	Sodium)	C		
		Vestibular Neuronitis	Sertraline			
		Vestibulitis	Hydrochloride			
		Viral Labyrinthitis	(Sertraline			
		Vision Blurred	Hydrochloride)	C		
		Weight Decreased	Pilocarpine			
		Weight Increased	Hydrochloride			
		Wound Infection	(Pilocarpine			
			Hydrochloride)	C		
			Metoprolol Succinate			
			(Metoprolol			
			Succinate)	C		
			Lansoprazole			
			(Lansoprazole)	C		
			Hyoscyamine Sulfate			
			(Hyoscyamine			
			Sulfate)	C		
			Diltiazem			
			Hydrochloride			
			(Diltiazem			
			Hydrochloride)	C		
			Bupropion			
			Hydrochloride			

(Bupropion Hydrochloride)	C
Celebrex (Celebrex)	C
Quetiapine Fumarate (Quetiapine Fumarate)	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C
Olanzapine	

Freedom Of Information (FOI) Report

(Olanzapine)	C
Omeprazole	
(Omeprazole)	C
Nasal Preparations	
(Nasal Preparations)	C
Nefazodone	
Hydrochloride	
(Nefazodone	
Hydrochloride)	C
Totolin	
(Guaifenesin,	
Phenylpropanolamine	
Hydrochloride)	C
Meclozine	
(Meclozine)	C
Fluticasone	
Propionate	
(Fluticasone	
Propionate)	C
Trimethobenzamide	
Hydrochloride	
(Trimethobenzamide	
Hydrochloride)	C
Respaire-Sr-120	
(Guaifenesin,	
Pseudoephedrine	
Hydrochloride)	C
Risperidone	
(Risperidone)	C

Date:11/19/04ISR Number: 4509481-6Report Type:Expedited (15-DaCompany Report #2004075222
 Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Amnesia	Health	Neurontin			
Initial or Prolonged	Depression	Professional	(Gabapentin)	PS		ORAL
600 MG, ORAL						
Other	Drug Effect Decreased		Prednisone			
	Excitability		(Prednisone)	SS		ORAL
60 MG (15 MG,						
4 IN 1 D),	Foot Operation					
	Gastritis					
ORAL	Headache		Vicodin (Hydrocodoe			

Mania	Bitartrate,	
Neuralgia	Paracetamol)	SS
Sleep Disorder	Escitalopram	
Somnolence	(Escitalopram)	SS
Suicidal Ideation	Provella-14	
White Blood Cell Count	(Estrogens	
Increased	Conjugated,	
	Medroxyprogesterone	
	Acetate)	C
	Nifedipine	
	(Nifedipine)	C
	Propranolol	
	Hydrochloride	
	(Propranolol	
	Hydrochloride)	C
	Cetirizine	
	Hydrochloride	
	(Cetirizine	
	Hydrochloride)	C
	Lansoprazole	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Lansoprazole) C

Date:11/19/04ISR Number: 4509487-7Report Type:Expedited (15-DaCompany Report #2004091367

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Gastric Haemorrhage	Consumer	Neurontin (Gabapentin)	PS		
2700 MG (900 MG, 3 IN 1 D),				Oxaprozin (Oxaprozin) All Other Therapeutic Products (All Other Therapeutic Products)	SS		

Date:11/19/04ISR Number: 4512801-XReport Type:Direct

Age: Gender:Male I/FU:I

Company Report #CTU 232556

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Refusal Of Treatment By Patient		Generic Neurontin (Gabapentin)	PS		
300 MG BID							

Date:11/19/04ISR Number: 4512809-4Report Type:Direct

Age: Gender:Female I/FU:I

Company Report #CTU 232485 E

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 25 MG PO HS		Idiopathic		Olanzapine	PS		ORAL
Initial or Prolonged 400 MG PO AM		Thrombocytopenic Purpura		Gabapentin	SS		ORAL
800 MG HS							

Date:11/19/04ISR Number: 4512861-6Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 232499

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Choking		Neurontin 300 Mg Po			
		Pharmaceutical Product		Qid	PS		
QID		Complaint					

Date:11/22/04ISR Number: 4507923-3Report Type:Direct
Age:50 YR Gender:Female I/FU:I

Company Report #CTU 232688

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dyspnoea		Neurontin Generic	PS		
100 MG QID		Fatigue					
		Vision Blurred					

Date:11/22/04ISR Number: 4507935-XReport Type:Direct
Age:51 YR Gender:Female I/FU:I

Company Report #CTU 232685

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
300 MG 2 PO		Headache		Gabapentin	PS		ORAL
TID		Nausea					
		Pharmaceutical Product Complaint Stomach Discomfort					

Date:11/22/04ISR Number: 4508008-2Report Type:Direct Company Report #CTU 232651
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	5 MG ONCE/PO		Blood Pressure Increased		Metalozone 5 Mg	PS		ORAL
400 MG THREE			Pharmaceutical Product Complaint		Neurontin 400 Mg	SS		
TIME			Therapeutic Response Unexpected With Drug Substitution		Gabapentin	SS		

Date:11/22/04ISR Number: 4510257-4Report Type:Expedited (15-DaCompany Report #2004061114
Age:60 YR Gender:Female I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	20 MG (20 MG,		Burning Sensation Cataract	Consumer Health	Lipitor (Atorvastatin)	PS		ORAL
Other	1 IN 1 D),		Coronary Artery Occlusion	Professional				
ORAL			Drug Hypersensitivity					
300 MG (300			Drug Ineffective Glaucoma		Neurontin (Gabapentin)	SS		ORAL

MG, 1 IN 1	Gout		
	Hypoaesthesia		
D), ORAL	Infusion Related Reaction	Vancomycin	
	Mobility Decreased	(Vancomycin)	SS
INTRAVENOUS	INTRAVENOUS		
	Myalgia	Cefazolin Sodiuium	
	Pain	(Cefazolin Sodiuium)	SS
INTRAVENOUS	INTRAVENOUS		
	Pain In Extremity	Metoprolol Tartrate	
	Paraesthesia	(Metoprolol	
	Postoperative Infection	Tartrate)	C
	Rash Pruritic	Metformin	
	Red Man Syndrome	Hydrochloride	
	Wound Complication	(Metformin	
	Wound Infection	Hydrochloride)	C
		Novolin 20/80	
		(Insulin Human,	
		Insulin Isophane,	
		Human Biosynthetic)	C
		Furosemide	
		(Furosemide)	C
		Potassium Chloride	
		(Potassium Chloride)	C
		Nicotinic Acid	
		(Nicotinic Acid)	C
		Acetylsalicylic Acid	
		(Acetylsalicylic	
		Acid)	C
		Lisinopril	
		(Lisinopril)	C
		Colchicine	

Freedom Of Information (FOI) Report

(Colchicine) C
 Human 70/30 (Insulin
 Human Injection,
 Isophane, Insulin
 Human Zinc
 Suspension) C

Date:11/22/04ISR Number: 4510268-9Report Type:Expedited (15-DaCompany Report #2004067531
 Age:42 YR Gender:Male I/FU:F

Outcome	PT
Death	Accidental Death
Hospitalization -	Agitation
Initial or Prolonged	Anger
Disability	Anxiety
Other	Arthralgia
	Atherosclerosis
	Back Pain
	Brain Oedema
	Burning Sensation
	Cardio-Respiratory Arrest
	Cardiomegaly
	Coma
	Coronary Artery
	Atherosclerosis
	Drug Ineffective
	Drug Interaction
	Erectile Dysfunction
	Excoriation
	Fall
	Feeling Abnormal
	General Physical
	Condition Abnormal
	Haematemesis
	Hepatic Steatosis
	Hypertensive Heart
	Disease
	Hypoaesthesia
	Intentional Self-Injury
	Intervertebral Disc
	Operation
	Joint Injury
	Laceration
	Limb Discomfort
	Lumbar Radiculopathy
	Major Depression

Mood Swings
Muscle Spasms
Muscle Spasticity
Nerve Root Injury
Neuropathic Pain
Pain
Pain In Extremity
Paraesthesia
Post Procedural
Complication
Pulmonary Oedema
Radiculitis
Radiculopathy
Renal Cyst
Respiratory Arrest

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Respiratory Depression Snoring Tearfulness Tenderness Ventricular Hypertrophy	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, 2 IN 1 D), ORAL			Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Benadryl (Diphenhydramine)	SS		ORAL
1-3 TABLETS Q12H, ORAL				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL
1-2 EVERY 4-6 HOURS (10 MG), ORAL				Hydrocodone (Hydrocodone)	SS		ORAL
10 MG (10 MG, 1 IN 1 D), ORAL				Zolpidem Tartrate (Zolpidem Tartrate)	SS		ORAL
80 MG (80 MG, 1 IN 1 D), ORAL				Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		ORAL
				Amitriptyline Hydrochloride (Amitriptyline			

Hydrochloride)	C
Fentanyl (Fentanyl)	C
Cyclobenzaprine	
Hydrochloride	
(Cyclobenzaprine	
Hydrochloride)	C
Bupropion	
Hydrochloride	
(Bupropion	
Hydrochloride)	C
Quinine Sulfate	
(Quinine Sulfate)	C
Sertraline	
Hydrochloride	
(Sertraline	
Hydrochloride)	C
Trazodone	
(Trazodone)	C
Hydroxyzine Embonate	
(Hydroxyzine	
Embonate)	C

Date:11/22/04ISR Number: 4510298-7Report Type:Expedited (15-DaCompany Report #2004058557
Age:45 YR Gender:Female I/FU:F

Outcome	PT
Other	Economic Problem
	Multiple Sclerosis
	Tooth Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Outcome	Report Source	Product	Role	Manufacturer	Route
400 MG (400 MG), ORAL		Tooth Fracture Treatment Noncompliance Weight Decreased	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Cetirizine Hydrochloride (Cetirizine Hydrochloride)	C		
				Ranitidine Hydrochloride (Ranitidine Hydrochloride)	C		
				Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride)	C		
				Morphine Sulfate (Morphine Sulfate)	C		
				Interferon Beta (Interferon Beta)	C		
				Morphine Sulfate (Morphine Sulfate)	C		
				Loperamide Hydrochloride (Loperamide Hydrochloride)	C		
				Oxybutynin (Oxybutynin)	C		
				Tizanidine Hydrochloride (Tizanidine Hydrochloride)	C		

Date:11/22/04ISR Number: 4510308-7Report Type:Expedited (15-DaCompany Report #2004045177
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Death		Health	Neurontin			

300 MG, ORAL

Professional

(Gabapentin)

PS

ORAL

Estradiol	
(Estradiol)	C
All Other	
Therapeutic Products	
(All Other	
Therapeutic	
Products)	C
Amlodipine Besilate	
(Amlodipine	
Besilate)	C
Levothyroxine	
(Levothyroxine)	C
Oxcarbazepine	
(Oxcarbazepine)	C
Olanzapine	
(Olanzapine)	C
Venlafaxine	
Hydrochloride	
(Venlafaxine	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C

Date:11/22/04ISR Number: 4510436-6Report Type:Expedited (15-DaCompany Report #KII-2004-0013986
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Acidosis Blood Alkaline Phosphatase Increased Blood Bicarbonate Decreased	Study Health Professional Other	Oxycodone Hydrochloride (Similar To Nda 20-553)(Oxycodone Hydrochloride)	PS		ORAL
ORAL		Blood Chloride Increased Blood Magnesium Increased		Olanzapine(Olanzapin e)	SS		ORAL
ORAL		Blood Pressure Systolic Increased		Gabapentin(Gabapenti n)	SS		ORAL
ORAL		Chest Pain Cough		Sertraline (Sertraline)	SS		ORAL
ORAL		Diabetes Insipidus Hypokalaemia		Benzodiazepine Derivatives()	SS		ORAL
ORAL		Multiple Drug Overdose		Lithium (Lithium)	SS		ORAL
		Pco2 Decreased Po2 Decreased Somnolence White Blood Cell Count Increased					

Date:11/22/04ISR Number: 4510736-XReport Type:Expedited (15-DaCompany Report #2004091185
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Phlebothrombosis	Foreign Health Professional Company	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1							

Representative

D), ORAL

Date:11/22/04ISR Number: 4511754-8Report Type:Expedited (15-DaCompany Report #2004089776
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decreased Activity	Consumer	Neurontin			
Disability		Diabetes Mellitus		(Gabapentin)	PS		
Other		Drug Ineffective		Gabapentin			
		Medication Error		(Gabapentin)	SS		
		Pain In Extremity		Oxycodone			
		Peroneal Muscular Atrophy		Hydrochloride			
		Therapeutic Response		(Oxycodone			
		Unexpected		Hydrochloride)	C		

Date:11/22/04ISR Number: 4511797-4Report Type:Expedited (15-DaCompany Report #2004090054
 Age:66 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Amnesia
Initial or Prolonged	Convulsion
Other	Coronary Artery Occlusion
	Crying
	Hypertension

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hypotension Post Procedural Complication	Report Source	Product	Role	Manufacturer	Route
25 MG (25 MG, 1 IN 1 D), ORAL		Ruptured Cerebral Aneurysm	Consumer	Zoloft (Sertraline)	PS		ORAL
1200 MG (300 MG, 2 IN 2 D)				All Other Therapeutic Products (All Other Therapeutic Products) Neurontin (Gabapentin)	SS SS		
80 MG (40 MG, 2 IN 1 D), ORAL				Accupril (Quinapril Hydrochloride)	SS		ORAL
10 MG (10 MG, 1 IN 1 D), ORAL	6 MON			Norvasc (Amlodipine)	SS		ORAL
				Clonidine (Clonidine)	C		

Date:11/22/04ISR Number: 4678797-5Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 241147

Outcome Dose Other THREE TID	Duration	PT Pharmaceutical Product Complaint Pyrexia	Report Source	Product	Role	Manufacturer	Route
				Neurontin 300 Mg	PS		

Urticaria
Vomiting

Date:11/23/04ISR Number: 4510658-4Report Type:Expedited (15-DaCompany Report #2004-UK-01033UK
Age:80 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Atrial Fibrillation	Foreign	Morphine			
Hospitalization - Initial or Prolonged		Blood Creatinine Increased	Health Professional	(0015/0122/0208/0157) (Morphine)	PS		
INTRAVENOUS	IV	Blood Ph Increased		Feldene (Piroxicam)	SS		
		Blood Potassium Decreased		Celebrex (Celecoxib)	SS		
		Blood Sodium Decreased		Neurontin			
		Cardiomegaly		(Gabapentin)	SS		
		Diarrhoea		Irbesartan			
		Electrolyte Imbalance		(Irbesartan)	SS		
		Myocardial Infarction		Prozac (Fluoxetine			
		Neutrophil Count Increased		Hydrochloride)	SS		
		Oliguria					
		Pulmonary Oedema					
		Respiratory Alkalosis					
		Ventricular Fibrillation					
		Vomiting					
		White Blood Cell Count Increased					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/23/04
 Age:55 YR
 Gender:Male
 I/FU:I

Company Report #CTU 232777

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Vision Blurred		Gabapentin	PS		
				Fosinopril Na	C		
				Meloxicam	C		
				Acetaminophen	C		
				Clotrimazole 1%	C		
				Cyclobenzaprine Hcl	C		
				Omeprazole	C		
				Diazepam	C		
				Irbesartan	C		
				Diltiazem	C		
				Gabapentin	C		
				Paroxetine Hcl	C		

Date:11/23/04
 Age:62 YR
 Gender:Male
 I/FU:F

Company Report #2004083706

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Apathy	Consumer	Neurontin			
		Erythema	Health	(Gabapentin)	PS		ORAL
600 MG (300		Fatigue	Professional				
MG, 2 IN 1		Nervousness					
D), ORAL		Personality Change		Lithium (Lithium)	C		
		Self Esteem Decreased		Esomeprazole			
				(Esomeprazole)	C		

Date:11/23/04
 Age:
 Gender:Female
 I/FU:F

Company Report #2004080553

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Accidental Overdose	Health	Neurontin			
Initial or Prolonged		Creatinine Renal	Professional	(Gabapentin)	PS		ORAL
1800 MG (600							

Other	Clearance Decreased		
MG, 3 IN 1			
	Dialysis		
D), ORAL			
	Disease Progression	Fentanyl (Fentanyl)	SS
TRANSDERMAL	50 MCG, 1 IN		
	Disorientation		
3 D,			
TRANSDERMAL	Extraocular Muscle		
	Disorder	Mycophenolate	
	Memory Impairment	Mofetil	
	Mental Status Changes	(Mycophenolate	
	Psychotic Disorder	Mofetil)	C
		Tacrolimus	C
		Acetylsalicylic Acid	
		(Acetylsalicylic	
		Acid)	C
		Prednisone	
		(Prednisone)	C
		Calcium Acetate	
		(Calcium Acetate)	C
		Labetalol	
		(Labetalol)	C
		Isosorbide Dinitrate	
		(Isosorbide	
		Dinitrate)	C
		Enalapril Maleate	
		(Enalapril Maleate)	C
		Paroxetine	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride
 (Paroxetine
 Hydrochloride) C
 Clonazepam
 (Clonazepam) C
 Valproate Semisodium
 (Valproate
 Semisodium) C

Date:11/23/04ISR Number: 4512085-2Report Type:Expedited (15-DaCompany Report #2004091369
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Circulatory Collapse	Health Professional	Neurontin (Gabapentin)	PS		
				Interferon Beta (Interferon Beta)	SS		

Date:11/23/04ISR Number: 4512091-8Report Type:Expedited (15-DaCompany Report #2004091215
 Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Upper Aphasia	Consumer	Lipitor (Atorvastatin)	PS		
Other		Cerebrovascular Accident		Neurontin (Gabapentin)	SS		ORAL
2400 MG (400 MG, 2 IN 1 D), ORAL		Coronary Artery Occlusion					
		Dyspepsia					
		Haematochezia					
1600 MG (400 MG, 2 IN 1 D), ORAL		Muscle Spasms		Gabapentin (Gabapentin)	SS		ORAL
		Pharmaceutical Product					
		Complaint					
		Treatment Noncompliance					
				Rofexoxib (Rofecoxib)	SS		
				Simvastatin (Simvastatin)	SS		

Minoxidil
(Minoxidil) C

Date:11/23/04ISR Number: 4527959-6Report Type:Periodic Company Report #2003121740
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression Drug Ineffective	Consumer Health	Lipitor (Atorvastatin)	PS		ORAL
10 MG		Facial Pain	Professional				
(DAILY), ORAL		Feeling Abnormal Glaucoma		Neurontin (Gabapentin)	SS		
300 MG		Pollakiuria					
(DAILY)		Tooth Disorder		Detrol La (Tolterodine L-Tartrate)	SS		ORAL
ORAL				Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
				Metoprolol (Metoprolol)	C		
				Glibenclamide (Glibenclamide)	C		

Freedom Of Information (FOI) Report

Metformin Hydrochloride (Metformin Hydrochloride)	C
Lisinopril (Lisinopril)	C
Glyceryl Trinitrate (Glyceryl Trinitrate)	C
Pantoprazole (Pantoprazole)	C
Gemfibrozil (Gemfibrozil)	C
Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)	C
Fenofibrate (Fenofibrate)	C
Triobe (Folic Acid, Pyridoxine, Cyanocobalamin)	C
Aspirin (Acetylsalicylic Acid)	C
Trazodone (Trazodone)	C

Date:11/24/04ISR Number: 4511321-6Report Type:Direct
Age:39 YR Gender:Male I/FU:I

Company Report #CTU 232987

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nausea		Gabapentin 300mg	PS		ORAL
900MG PO TID		Pharmaceutical Product Complaint		Epivir	C		
				Videx Ec	C		
				Sustiva	C		
				Azithromycin	C		
				Amitriptyline	C		
				Morphine Sr	C		
				Remeron	C		
				Rifabutin	C		
				Ethambutol	C		

Ativan C
Diflucan C

Date:11/24/04ISR Number: 4512966-XReport Type:Direct Company Report #CTU 232951
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disturbance In Attention		Neurontin	PS		ORAL
200 MG 2 BID		Eye Swelling					
PO		Memory Impairment					
		Sleep Disorder					
		Thinking Abnormal					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

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
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		(Paracetamol)	SS		
		Pneumonia					
		Pneumonia Aspiration					
		Polysubstance Abuse					
		Pulmonary Congestion					
		Pulmonary Oedema					
		Respiratory Depression					
		Resuscitation					

Date:11/26/04ISR Number: 4514668-2Report Type:Expedited (15-DaCompany Report #2004092446

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Atrioventricular Block	Foreign	Neurontin			
		First Degree	Health	(Gabapentin)	PS		ORAL
ORAL		Atrioventricular Block	Professional				
		Second Degree					

Date:11/26/04ISR Number: 4514882-6Report Type:Expedited (15-DaCompany Report #2004092362

Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Diabetes Insipidus Pharmaceutical Product	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL		Complaint	Professional	Neurontin (Tablets) (Gabapentin)	SS		
UNKNOWN							

Date:11/26/04ISR Number: 4515017-6Report Type:Expedited (15-DaCompany Report #2004085873
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Breast Pain Hypertrophy Breast	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 2 IN 1 D), ORAL		Mastitis	Professional Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/26/04ISR Number: 4515021-8Report Type:Expedited (15-DaCompany Report #2004011179
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (3 IN 1 D), ORAL		Condition Aggravated Diarrhoea Erectile Dysfunction	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:11/26/04ISR Number: 4515317-XReport Type:Expedited (15-DaCompany Report #2004077798
Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability ORAL		Crying Fear Hallucination	Foreign Consumer	Neurontin	PS		ORAL

Date:11/26/04ISR Number: 4515324-7Report Type:Expedited (15-DaCompany Report #2004081142
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG (600 Other MG), ORAL		Anaemia Drug Ineffective Drug Interaction	Foreign Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
ORAL		Dyslipidaemia Epilepsy	Company Representative	Oxcarbazepine (Oxcarbazepine)	SS		ORAL
2 IN 1 D, ORAL		Hyponatraemia Leukopenia Neutropenia		Risperidone (Risperidone) Noctran 10 (Acepromazine,	SS		ORAL

100 MG

Aceprometazine,
 Clorazepate
 Dipotassium) SS
 Phenobarbital Sodium
 (Phenobarbital
 Sodium) SS

 Pravastatin Sodium
 (Pravastatin Sodium) C
 Ferrous Sulfate
 (Ferrous Sulfate) C

Date:11/26/04ISR Number: 4515331-4Report Type:Expedited (15-DaCompany Report #HQWYE561219NOV04
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Difficulty In Walking Drug Interaction Dysarthria Feeling Cold Joint Stiffness	Consumer	Effexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		
300 MG		Mobility Decreased Musculoskeletal Stiffness		Ambien (Zolpidem Tartrate)	SS		
100 MG		Nightmare Pain Speech Disorder		Endocet (Oxycodone Hydrochloride/Parace tamol)	SS		
7.5/3.25		Tenderness					
EVERY 4 HOURS							
THROUGHOUT							

Freedom Of Information (FOI) Report

THE DAY

Hydroxyzine
(Hydroxyzine) SS

25 MG IN THE

EVENING

Magnesium
(Magnesium) SS

400 MG

Neurontin
(Gabapentin) SS

ONE IN AM,

TWO IN

EVENING

Premarin (Conjugated
Estrogens) SS
Trazodone
(Trazodone) SS

100 MG IN THE

EVENING

Unspecified Vitamins
(Unspecified
Vitamins) SS

Date:11/26/04ISR Number: 4515340-5Report Type:Expedited (15-DaCompany Report #2004091487

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		
Other		Convulsion		Tiagabine Hydrochloride (Tiababine Hydrochloride)	SS		
		Drug Interaction		Alendronate Sodium (Alendronate Sodium)	C		ORAL

70 MG (70 MG,

1 IN 1 WK),

ORAL

Date:11/26/04ISR Number: 4515432-0Report Type:Expedited (15-DaCompany Report #2004096020
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety Convulsion Depression Drug Ineffective Hallucination	Consumer	Neurontin (Gabapentin)	PS		

Date:11/26/04ISR Number: 4515440-XReport Type:Expedited (15-DaCompany Report #2004093246
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide Polytraumatism	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:11/26/04ISR Number: 4515449-6Report Type:Expedited (15-DaCompany Report #2004093208
Age: Gender:I/FU:I

Outcome	Duration	PT	Report Source
Death		Completed Suicide	Health Professional Company

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative

Dose	Duration	Product	Role	Manufacturer	Route
		Neurontin (Gabapentin)	PS		

Date:11/26/04ISR Number: 4515452-6Report Type:Expedited (15-DaCompany Report #2004094919
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Gabapentin (Gabapentin)	PS		

Date:11/26/04ISR Number: 4515467-8Report Type:Expedited (15-DaCompany Report #2004084796
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (200 Other MG, 1 IN 1 D)		Confusional State Drug Ineffective Feeling Abnormal Grand Mal Convulsion Paraesthesia Oral	Consumer Health Professional	Neurontin (Gabapentin) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	PS C		

Date:11/26/04ISR Number: 4515548-9Report Type:Expedited (15-DaCompany Report #2004067531
Age:42 YR Gender:Male I/FU:F

Outcome	PT
Death	Accidental Death
Hospitalization - Initial or Prolonged	Agitation Anger
Disability	Anxiety
Other	Arthralgia Aspiration Atherosclerosis

Back Pain
Brain Oedema
Burning Sensation
Cardiomegaly
Coma
Computerised Tomogram
Abnormal
Coronary Artery
Atherosclerosis
Drug Ineffective
Drug Interaction
Drug Level Above
Therapeutic
Erectile Dysfunction
Excoriation
Fall
Haematemesis
Hepatic Steatosis
Hypertensive Heart
Disease
Hypoaesthesia
Insomnia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Intervertebral Disc Protrusion Lumbar Radiculopathy			
600 MG (300 MG, 2 IN 1 D), ORAL		Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Professional				
			Muscle Spasms			
			Muscle Tightness			
			Neck Pain Pain			
			Benadryl (Diphenhydramine)	SS		ORAL
			Pain In Extremity Paraesthesia Post Procedural Complication			
1-3 TABLETS Q12H, ORAL			Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL
			Pseudarthrosis			
1-2 EVERY 4-6 HOURS (10 MG), ORAL			Pulmonary Oedema Radiculitis			
			Hydrocodone (Hydrocodone)	SS		ORAL
			Renal Cyst			
			Respiratory Arrest			
10 MG (10 MG, 1 IN 1 D),ORAL			Respiratory Depression Snoring			
			Zolpidem Tartrate (Zolpidem Tartrate)	SS		ORAL
			Spinal Disorder			
			Spinal Myelogram Abnormal			
80 MG (80 MG, 1 IN 1 D), ORAL			Tenderness Ventricular Hypertrophy Vomiting			
			Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		ORAL
			Amitriptyline Hydrochloride (Amitriptyline			

Hydrochloride)	C
Fentanyl (Fentanyl)	C
Cyclobenzaprine	
Hydrochloride	
(Cyclobenzaprine	
Hydrochloride)	C
Bupropion	
Hydrochloride	
(Bupropion	
Hydrochloride)	C
Quinine Sulfate	
(Quinine Sulfate)	C
Sertraline	
Hydrochloride	
(Sertraline	
Hydrochloride)	C
Trazodone	
(Trazodone)	C
Hydroxyzine Embonate	
(Hydroxyzine	
Embonate)	C

Date:11/26/04ISR Number: 4515553-2Report Type:Expedited (15-DaCompany Report #2004084794
Age: Gender:Female I/FU:F

Outcome	PT
Other	Arterial Bypass Operation
	Diabetes Mellitus
	Diabetic Neuropathy

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hypertension Muscle Spasms Neuropathy Peripheral	Report Source	Product	Role	Manufacturer	Route
2400 MG (600 MG, 4 IN 1 D), ORAL			Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
50 MG (50 MG, 1 IN 1 D), ORAL				Zoloft (Sertraline)	SS		ORAL
30 MG (30 MG, 1 IN 1 D), ORAL				Procardia (Nifedipine)	SS		ORAL

Date:11/26/04ISR Number: 4515555-6Report Type:Expedited (15-DaCompany Report #2004089776
Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other		Bone Disorder Diabetes Mellitus Drug Ineffective Pain In Extremity Pharmaceutical Product Complaint Therapeutic Response Unexpected	Consumer	Neurontin (Tablets) (Gabapentin) Gabapentin (Gabapentin) Oxycodone Hydrochloride (Oxycodone Hydrochloride)	PS SS C		

Date:11/26/04ISR Number: 4515568-4Report Type:Expedited (15-DaCompany Report #2004091687
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction Drug Toxicity	Health Professional	Neurontin (Gabapentin)	PS		
(3 IN 1 D)		Oophorectomy Palpitations		Estradiol (Estradiol) Ibuprofen (Ibuprofen)	SS SS		
PRN (800 MG)				Progesterone (Progesterone)	SS		
(1 IN 1 WK)				Testosterone (Testosterone)	SS		
1 DROP (1 DROP, 1 IN 1 D)							

Date:11/26/04ISR Number: 4515572-6Report Type:Expedited (15-DaCompany Report #2004093703
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis Gout Osteoporosis Surgery	Consumer	Neurontin (Gabapentin) Metaxalone (Metaxalone) All Other Therapeutic Products (All Other Therapeutic Products)	PS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:11/26/04ISR Number: 4515605-7Report Type:Expedited (15-DaCompany Report #2004021476
 Age:84 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (400 Other MG, 3 IN 1 D), ORAL	Blood Cholesterol Increased	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
	Blood Creatinine Increased	Professional Company				
	Blood Glucose Increased Blood Pressure Decreased Brain Scan Abnormal Bundle Branch Block Left Cardio-Respiratory Arrest Circulatory Collapse Ear Pain Fatigue Fibrin D Dimer Increased Haematocrit Increased Haemoglobin Increased Hypertension Loss Of Consciousness Lymphopenia Malaise Oxygen Saturation Decreased Pallor Paraesthesia Treatment Noncompliance Vomiting	Representative	Acetylsalicylate Lysine (Acetylsalicylate Lysine) Paracetamol (Paracetamol)	C C		

Date:11/26/04ISR Number: 4515612-4Report Type:Expedited (15-DaCompany Report #2004091667
 Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other 1500 MG (300 MG, 5 IN 1	Abdominal Discomfort Amnesia	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
	Anxiety	Professional				

D), ORAL

Confusional State

Thinking Abnormal

Ibuprofen
(Ibuprofen)

C

Date:11/26/04ISR Number: 4515623-9Report Type:Expedited (15-DaCompany Report #2004093707

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (100 MG, 3 IN 1		Coagulation Time Prolonged	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Drug Interaction Haematoma	Professional				
		Haemodilution Prothrombin Time Abnormal		Oxycodone Hydrochloride (Oxycodone Hydrochloride) Diclofenac (Dicloenac) Pandeine Co (Codeine Phosphate, Paracetamol) Alendornate Sodium (Alendronate Sodium)	C C C C		

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Freedom Of Information (FOI) Report

Diazepam 9diazepam) C
 Paroxetine
 (Paroxetine) C
 Betahistine
 (Betahistine) C
 Nulytely (Macrogol,
 Potassium Chloide,
 Sodium Bicarbonate,
 Sodium Chloride) C
 Flunitrazepam9flunit
 razepam) C
 Nifedipine
 (Nifedipine) C
 Furosemide
 (Furosemide) C
 Ferrous Fumarate
 (Ferrous Fumarate) C

Date:11/26/04ISR Number: 4515625-2Report Type:Expedited (15-DaCompany Report #2004095235

Age:72 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (1 D), Other ORAL		Eosinophil Percentage Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Epilepsy	Professional				
		Haemoglobin Decreased Pruritus Toxic Skin Eruption					

Date:11/26/04ISR Number: 4515636-7Report Type:Expedited (15-DaCompany Report #2004093336

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other (1800 MG), ORAL		Gingival Atrophy Tooth Loss	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
			Professional				
				Lansoprazole			

(Lansoprazole)	C
Fluoxetine	
Hydrochloride	
(Fluoxetine	
Hydrochloride)	C
Aporex	
(Dextropropoxyphene	
Hydrochloride,	
Paracetamol)	C
Zopiclone	
(Zopiclone)	C
Diclofenac	
9diclofenac0	C
Nefopam	
Hydrochloride	
Nefopam	
Hydrochloride)	C
Buprenorphine	
Hydrochloride	
(Buprenorphine	
Hydrochloride)	C
Amitriptyline	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Amitriptyline) C

Date:11/29/04ISR Number: 4514799-7Report Type:Direct
Age:55 YR Gender:Female I/FU:I

Company Report #CTU 233068

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain		Gabapentin By Teva	PS	Teva	ORAL
400 MG QID		Cerebrovascular Accident					
PO/ 2		Chest Pain					
OCCASIONS		Coordination Abnormal		Effexor	C		
		Palpitations		Zanaflex	C		
		Pharmaceutical Product		Darvocet N	C		
		Complaint		Prevacid	C		
		Speech Disorder		Sonata	C		
		Visual Disturbance		Clonidine	C		

Date:11/30/04ISR Number: 4515161-3Report Type:Direct
Age:51 YR Gender:Female I/FU:I

Company Report #CTU 233229

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Headache		Gabapentin	PS		ORAL
300 MG 2 PO		Nausea					
Intervention to		Pharmaceutical Product					
TID		Complaint					
Prevent Permanent		Stomach Discomfort					
Impairment/Damage							

Date:11/30/04ISR Number: 4515169-8Report Type:Direct
Age: Gender: I/FU:I

Company Report #CTU 233249

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
300 MG PO BID		Convulsion		Neurontin	PS		ORAL
		Pharmaceutical Product					
		Complaint					

Date:11/30/04ISR Number: 4517905-3Report Type:Expedited (15-DaCompany Report #2004085873
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Mastitis	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 2 IN 1 D)			Professional Company				
ORAL			Representative				

Date:11/30/04ISR Number: 4518036-9Report Type:Expedited (15-DaCompany Report #9042
Age:80 YR Gender:Male I/FU:F

Outcome	PT
Death	Alkalosis Atrial Fibrillation Blood Creatinine Increased Blood Ph Increased Blood Potassium Decreased Blood Pressure Increased Blood Sodium Decreased Cardiomegaly

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
INTRAVENOUS	2.5 MG	Foreign	Morphine	PS		
	IV					
	Increased		Piroxicam	SS		
	Oliguria		Celecoxib	SS		
	Oxygen Saturation		Gabapentin	SS		
	Decreased		Irbesartan	SS		
	Pulmonary Oedema		Fluoxetine	SS		
	Respiratory Alkalosis		Furosemide	C		
	Ventricular Fibrillation		Amiodarone	C		
	Vomiting					
	White Blood Cell Count					
	Increased					

Date:11/30/04ISR Number: 4518092-8Report Type:Expedited (15-DaCompany Report #2004095236

Age: Gender: I/FU:I

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route
Dose						
Other		Consumer	Neurontin (Gabapentin)	PS		
900 MG (300						
MG, 3 IN 1 D)						
			Zolpidem Tartrate (Zolpidem Tartrate0	SS		
(10 MG)						
			Estrogens Conjugted (Estrogens			
(0.625 MG)			Conjugated)	SS		
			Venlafaxine Hydrochloride			
(300 MG)			9venlafaxine Hydrochloride)	SS		
			Oxycocet (Oxycodone Hydrochloride, Paracetamol)	SS		
(6 IN 1 D)						
			Hydroyzine Hydrochloride			

(25 MG, 1 IN

1 D)

(100 MG, 1 IN

1D)

(400 MG)

(Hydroxyzine
Hydrochloride) SS

Trazadone
(Trazadone) SS

Magnesium
(Magnesium) SS

Vitamins (Vitamins) SS

Date:12/01/04ISR Number: 4517910-7Report Type:Direct
Age:47 YR Gender:Female I/FU:I

Company Report #CTU 233350

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disturbance In Attention		Neurontin	PS		ORAL
200 MG 2 BID		Eye Swelling					
PO		Memory Impairment					
		Sleep Disorder					
		Thinking Abnormal					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/04ISR Number: 4518532-4Report Type:Expedited (15-DaCompany Report #2004037837

Age:73 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG (600 Other MG, 3 IN 1 D), ORAL	Ankle Fracture Blood Cholesterol Increased Flat Feet	Consumer	Neurontin (Gabapentin)	PS		ORAL
10 MG (10 MG, 1 IN 1 D), ORAL	Foot Fracture Herpes Zoster Osteoporosis Osteoporotic Fracture Pain		Lipitor (Atorvastatin)	SS		ORAL
			Rosiglitazone Maleate (Rosiglitazone Maleate)	C		
			Insulin Human (Insulin Human)	C		
			Insulin Glargine (Insulin Glargine)	C		
			Furosemide (Furosemide)	C		
			Amlodipine Besilate (Amlodipine Besilate)	C		
			Folic Acid (Folic Acid)	C		
			Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
			Carisoprodol (Carisoprodol)	C		
			Norgesic Forte (Acetylsalicylic Acid, Caffeine, Orphenadrine Citrate)	C		
			Tocopherol (Tocopherol)	C		

Date:12/01/04ISR Number: 4518535-XReport Type:Expedited (15-DaCompany Report #2004082915

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Arthralgia	Health	Neurontin			
Initial or Prolonged	Blood Cholesterol	Professional	(Gabapentin)	PS		ORAL
900 MG (300	Increased					
Other	Body Height Decreased					
MG, 3 IN 1						
D), ORAL	Bursitis		Lipitor			
	Cyst		(Atorvastatin)	SS		ORAL
ORAL	Road Traffic Accident		Calcium Carbonate			
	Scoliosis		(Calcium Carbonate)	C		
	Spinal Cord Disorder		Centrum Silver			
	Spinal Osteoarthritis		(Ascorbic Acid,			
	Wheelchair User		Calcium, Minerals			
			Nos, Retinol,			
			Tocopheryl Acetate,	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/01/04ISR Number: 4518949-8Report Type:Expedited (15-DaCompany Report #2004095233

Age:0 DY Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Duration Drug Exposure During Pregnancy	Foreign Health	Neurontin (Gabapentin)	PS		
TRANSPLACENTAL Congenital Anomaly PLACENTAL	1.8 GRAM, Ventricular Septal Defect	Professional				
TRANSPLACENTAL PLACENTAL	2 MG,		Lorazepam (Lorazepam)	SS		
TRANSPLACENTAL PLACENTAL	100 MG,		Topiramate (Topiramate)	SS		

Date:12/01/04ISR Number: 4518964-4Report Type:Expedited (15-DaCompany Report #2004096022

Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other (1500 MG), ORAL	Duration Hallucination, Auditory	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
			Beclometasone (Beclometasone)	C		
			Paracetamol (Paracetamol)	C		
			C		
			Salmeterol (Salmeterol)	C		
			Tramadol (Tramadol)	C		
			Sumatriptan (Sumatriptan)	C		
			Cyclizine (Cyclizine)	C		
			Dipipanone			

(Dipipanone) C
Fluoxetine
(Flooxetine) C

Date:12/01/04ISR Number: 4519286-8Report Type:Expedited (15-DaCompany Report #2004096018
Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin			
Other		Completed Suicide		(Gabapentin)	PS		
		Polytraumatism					

Date:12/01/04ISR Number: 4519289-3Report Type:Expedited (15-DaCompany Report #2004096021
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Convulsion	Consumer	Neurontin			
Other		Hallucination		(Gabapentin)	PS		
		Headache					
		Loss Of Consciousness					
		Memory Impairment					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/02/04ISR Number: 4518328-3Report Type:Expedited (15-DaCompany Report #2004044965

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		No Adverse Drug Effect	Consumer	Neurontin			
			Health	(Gabapentin)	PS		
			Professional	Paracetamol			
				(Paracetamol)	C		

Date:12/02/04ISR Number: 4518381-7Report Type:Expedited (15-DaCompany Report #2004096313

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Circulatory Collapse	Consumer	Neurontin			
Other				(Gabapentin)	PS		
				Interferon Beta			
				(Interferon Beta)	SS		

Date:12/02/04ISR Number: 4519415-6Report Type:Expedited (15-DaCompany Report #2004097851

Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pancreatitis	Foreign	Neurontin (Tablets)			
Other		Urinary Tract Infection	Health	(Gabapentin)	PS		ORAL
300 MG (100			Professional				
MG, 3 IN 1				Sinemet (Carbidopa,			
				Levodopa)	C		
				Clopidogrel Sulfate			
				(Clopidogrel			
				Sulfate)	C		
				Donepezil			
				Hydrochloride			
				(Donepezil			
				Hydrochloride)	C		
				Paracetamol			
				(Paracetamol)	C		

D), ORAL

Calcium (Calcium) C
 Citalopram
 Hydrobromide
 (Citalopram
 Hydrobromide) C
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C
 Morphine (Morphine) C
 Pantoprazole
 (Pantoprazole) C

Date:12/02/04ISR Number: 4519422-3Report Type:Expedited (15-DaCompany Report #2004096163
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Loss Of Consciousness	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/02/04ISR Number: 4519427-2Report Type:Expedited (15-DaCompany Report #2004092446
 Age:64 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1800 MG (300 MG, 6 IN 1 D), ORAL		Atrioventricular Block First Degree Atrioventricular Block Second Degree Vertigo	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Levothyroxine (Levothyroxine)	C		
				Amitriptyline (Amitriptyline)	C		
				Benzyloenicillin Sodium (Benzyloenicillin Sodium)	C		
				Diclofenac (Diclofenac)	C		

Date:12/02/04ISR Number: 4519924-XReport Type:Direct Company Report #CTU 233458
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 600 MG BID Initial or Prolonged ORAL		Asthenia General Physical Health		Gabapentin 300 Mg	PS		ORAL
10 MG BID ORAL		Deterioration Weight Decreased		Methadone 10 Mg	SS		ORAL
				Atenolol	C		
				Buspirone	C		
				Cyanocobalamin	C		
				Docusate	C		
				Furosemide	C		
				Metolazone	C		
				Omeprazole	C		
				Kcl	C		

Date:12/03/04ISR Number: 4518918-8Report Type:Direct
Age:69 YR Gender:Female I/FU:I

Company Report #CTU 233615

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product		Gabapentin	PS		
300 MG TID		Complaint Rash Erythematous					

Date:12/03/04ISR Number: 4519747-1Report Type:Direct
Age:29 YR Gender:Female I/FU:I

Company Report #CTU 233542

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Gabapentin	PS		ORAL
300 MG TID PO		Drug Intolerance Pharmaceutical Product Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/03/04ISR Number: 4520359-4Report Type:Direct
Age: Gender: I/FU:I

Company Report #CTU 233629

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated		Neurontin	PS		ORAL
300 MG PO TID		Muscle Spasticity Pain Pharmaceutical Product Complaint					

Date:12/03/04ISR Number: 4520410-1Report Type:Expedited (15-DaCompany Report #2004097842
Age:2 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Body Height Below Normal Drug Exposure During Pregnancy Late Developer Premature Baby Small For Dates Baby Weight Gain Poor	Foreign Health Professional	Neurontin (Neurontin) Phenobarbital (Phenobarbital) Clobazam (Clobazam)	PS SS SS		

Date:12/03/04ISR Number: 4520413-7Report Type:Expedited (15-DaCompany Report #2004081134
Age:80 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death Hospitalization - ORAL Initial or Prolonged Other		Alkalosis Atrial Fibrillation Blood Creatinine Increased Blood Potassium Decreased Blood Sodium Decreased Cardiomegaly Diarrhoea Myocardial Infarction Neutrophil Count Increased	Foreign Health Professional	Neurontin (Gabapentin) Feldene (Piroxicam) Celecoxib (Celecoxib) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Irbesartan (Irbesartan)	PS SS SS SS SS		ORAL ORAL

Oliguria
Pulmonary Oedema
Ventricular Fibrillation
Vomiting
White Blood Cell Count
Increased

Morphine (Morphine) SS
Prochlorperazine
(Prochlorperazine) C

Date:12/03/04ISR Number: 4520416-2Report Type:Expedited (15-DaCompany Report #2004097200
Age:81 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (900 MG, 3 IN 1 D), ORAL	Jaundice Cholestatic	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
			Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
			Nifedipine (Nifedipine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/03/04ISR Number: 4520418-6Report Type:Expedited (15-DaCompany Report #2004081132

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bone Marrow Depression	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:12/03/04ISR Number: 4520477-0Report Type:Expedited (15-DaCompany Report #2004097960

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged ORAL		Activities Of Daily Living Impaired	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Confusional State		Antidepressants (Antidepressants)	C		
		Coordination Abnormal		Anxiolytic (Anxyolytics)	C		
		Mental Disorder		Antihypertensives	C	C	
		Motor Dysfunction					
		Overdose					

Date:12/03/04ISR Number: 4520506-4Report Type:Expedited (15-DaCompany Report #2004051029

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Brachial Plexus Injury	Consumer	Neurontin (Gabapentin)	PS		
Other		Limb Injury	Health Professional				
		Multiple Fractures					
		Muscle Spasms					
		Neck Injury					
		Pain					
		Polytraumatism					
		Road Traffic Accident					
		Scapula Fracture					
		Skeletal Injury					

Date:12/03/04ISR Number: 4520507-6Report Type:Expedited (15-DaCompany Report #2004074283
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Balance Disorder	Consumer	Neurontin			
Initial or Prolonged	Deafness	Health	(Gabapentin)	PS		
Other	Tinnitus	Professional				

Date:12/03/04ISR Number: 4520508-8Report Type:Expedited (15-DaCompany Report #2004009355
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Cardiac Disorder
Other	Chest Pain
	Chills
	Condition Aggravated
	Drug Tolerance Decreased
	Erectile Dysfunction
	Facial Pain
	Feeling Cold

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	General Physical Health Deterioration	Report Source	Product	Role	Manufacturer	Route
1800 MG	(TID), ORAL	Gingival Pain Glossodynia Neck Pain	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
20 MG	(DAILY), ORAL	Oesophageal Spasm Pain In Jaw Sleep Disorder	Professional	Lipitor (Atorvastatin)	SS		ORAL
1000 MG	(BID), ORAL	Tooth Abscess Tremor		Lithium (Lithium) (Lithium) Naproxen (Naproxen) Rofecoxib (Rofecoxib) Amoxicillin (Amoxicillin)	SS SS SS SS		ORAL
				All Other Therapeutic Products (All Other Therapeutic Products) Levothyroxine Sodium (Levothyroxine Sodium) Vitamins (Vitamins) Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C C C C		

Date:12/06/04ISR Number: 4520426-5Report Type:Direct
Age:35 YR Gender:Male I/FU:I

Company Report #CTU 233732

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG 2 BID	35 DAY	Condition Aggravated		Neurontin	PS		

Convulsion
Drug Effect Decreased
Pharmaceutical Product
Complaint

Date:12/06/04ISR Number: 4520866-4Report Type:Direct
Age:75 YR Gender:Male I/FU:I

Company Report #CTU 233708

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abnormal Behaviour Anger		Neurontin 300 Mg Pfizer	PS	Prizer	ORAL
300 MG BID PO		Balance Disorder Gait Disturbance Impatience		Enalapril Triamptereine/Hctz	C C		

Date:12/06/04ISR Number: 4520912-8Report Type:Expedited (15-DaCompany Report #L04-USA-07403-12
Age:35 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Completed Suicide Multiple Drug Overdose	Literature Health Professional	Citalopram Tramadol Gabapentin	PS SS SS		
Death							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/06/04ISR Number: 4521149-9Report Type:Expedited (15-DaCompany Report #2004097068
Age:53 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Abdominal Pain Upper Agranulocytosis Decreased Immune Responsiveness Lymphadenopathy Neuropathic Pain Neutropenic Sepsis Upper Respiratory Tract Infection White Blood Cell Count Decreased	Foreign Literature Health Professional	Gabapentin (Gabapentin) Therapeutic Radiopharmaceuticals (Therapeutic Radiopharmaceuticals) Lansoprazole (Lansoprazole) Morphine Sulfate (Morphine Sulfate) Senna (Senna)	PS C C C C		

Date:12/06/04ISR Number: 4521154-2Report Type:Expedited (15-DaCompany Report #2004098594
Age: Gender: I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Renal Failure	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:12/06/04ISR Number: 4521169-4Report Type:Expedited (15-DaCompany Report #2004090318
Age:59 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2400 MG (800 Other MG, 3 IN 1 D), ORAL	Dyspepsia Gamma-Glutamyltransferase Increased Multiple Sclerosis Oedema Peripheral Pain In Extremity	Foreign Health Professional Company Representative	Neurontin (Tablets) (Gabapentin) Propofan (Caffeine, Carbasalate Calcium, Chlorphenamine Maleate,	PS		ORAL

Dextropoxyphene, C
Pinaverium Bromide C
(Pinaverium Bromide)
Atorvastatin Calcium C
(Atorvastatin
Calcium)

Date:12/06/04ISR Number: 4521683-1Report Type:Expedited (15-DaCompany Report #L04-USA-07403-20
Age:44 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Levothyroxine Valproic Acid Gabapentin	PS SS SS		

Date:12/06/04ISR Number: 4521685-5Report Type:Expedited (15-DaCompany Report #L04-USA-07403-02
Age:53 YR Gender: I/FU:I

Outcome	PT	Report Source
Death	Completed Suicide Multiple Drug Overdose	Literature Health

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Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
		Levothyroxine	PS		
		Ethylene Glycol (Polyethylene Glycol)	SS		
		Gabapentin	SS		

Date:12/06/04ISR Number: 4521961-6Report Type:Expedited (15-DaCompany Report #2004082916
Age:44 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2400 MG (1200 Other MG, 2 IN 1 D), ORAL	Brachial Plexus Lesion Cervical Rib Excision Monoplegia Pain In Extremity	Foreign Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Morphine (Morphine)	C		

Date:12/06/04ISR Number: 4522197-5Report Type:Expedited (15-DaCompany Report #HQWYE644223NOV04
Age:38 YR Gender:Female I/FU:I

Outcome	PT
Death	Apnoea Arthralgia Arthropathy Aspiration Asthenia Blood Urine Present Cardio-Respiratory Arrest Chills Coma Completed Suicide Dizziness Drug Toxicity Fall Gastric Perforation Haematemesis

Headache
Hypotension
Injury
Insomnia
Intentional Misuse
Intervertebral Disc
Disorder
Lividity
Medication Error
Muscle Spasms
Neck Pain
Overdose
Pain In Extremity
Peripheral Coldness
Petechiae
Pharmaceutical Product
Complaint
Postoperative Infection
Precancerous Cells
Present
Pulmonary Congestion

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pulse Absent Rash Smear Cervix Abnormal	Consumer	Effexor (Venlafaxine Hydrochloride, Tablet)	PS		
		Ulnar Nerve Palsy	Consumer	Duragesic (Fentanyl)	SS		
				Neurontin (Gabapentin)	SS		
				Atenolol	C		
				Lorazepam	C		
				Flexeril (Cyclobenzaprine Hydrochloride)	C		
				Bextra (Valdecoxib)	C		
				Temazepam	C		
				Topamax (Topiramate)	C		
				Ambien (Zolpidem Tartrate)	C		
				Wellbutrin XL (Bupropion Hydrochloride)	C		
				Diazepam	C		
				Oxycodone Hydrochloride	C		
				Clonazepam	C		
				Morphine	C		

Date:12/07/04ISR Number: 4521855-6Report Type:Expedited (15-DaCompany Report #2004090054
Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 25 MG (25 MG, Initial or Prolonged 1 IN 1 D), Other ORAL		Amnesia	Consumer	Zoloft (Sertraline)	PS		ORAL
		Convulsion	Health				
		Coronary Artery Occlusion	Professional				
		Crying Headache Hypertension Hypotension Ruptured Cerebral		All Other Therapeutic Products (All Other Therapeutic Products)	SS		

	Aneurysm Therapeutic Response	Neurontin (Gabapentin)	SS	
1200 MG (300 MG, 2 IN 2 D)	Increased			
OPHTHALMIC	80 MG (40 MG, 2 IN 1 D),	Accupril (Quinapril Hydrochloride)	SS	
ORAL				
10 MG (10 MG, 1 IN 1 D),		Norvasc (Amlodipine)	SS	ORAL
ORAL	6 MON	Clonidine (Clonidine)	C	

Date:12/07/04ISR Number: 4521944-6Report Type:Direct
Age:36 YR Gender:Male I/FU:I

Company Report #CTU 233776

Outcome
Disability
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
ONCE A DAY		Headache Medication Error		Mfg: Teva Generic For Neurontin 400mg	PS	Teva	
		Pharmaceutical Product Complaint					

Date:12/07/04ISR Number: 4521945-8Report Type:Direct Company Report #CTU 233774
 Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Drug Effect Decreased		Teva Generic For Neurontin 400mg	PS	Teva	
THREE TIMES A DAY		Pharmaceutical Product Complaint					

Date:12/07/04ISR Number: 4522175-6Report Type:Expedited (15-DaCompany Report #2004100272
 Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Retinal Haemorrhage	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
48 00 MG (1200 MG, 4 IN 1 D), ORAL			Company Representative				

Date:12/07/04ISR Number: 4522179-3Report Type:Expedited (15-DaCompany Report #2004098814
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Anxiety	Consumer	Lipitor			

Initial or Prolonged 80 MG (80 MG, Other 1 IN 1 D)	Burning Sensation	(Atorvastatin)	PS
	Chest Pain		
	Difficulty In Walking	Neurontin	
900 MG (300 MG, 3 IN 1 D)	Drug Ineffective	(Gabapentin)	SS
	Dry Mouth		
20 MG (20 MG, 1 IN 1 D)	Feeling Abnormal	Zoloft (Sertraline)	SS
	Hypoaesthesia		
	Muscle Twitching	Clopidogrel Sulfate	
	Myocardial Infarction	(Clopidogrel	
	Nerve Injury	Sulfate)	C
	Paraesthesia	Omeprazole	
	Psychomotor Hyperactivity	(Omeprazole)	C
	Sleep Disorder	Metoprolol	
	Stent Occlusion	(Metoprolol)	C
		Temazepam	
		(Temazepam)	C
		Acetylsalicylic Acid	
		(Acetylsalicylic	
		Acid)	C
		Potassium Chloride	
		(Potassium Chloride)	C
		Methocarbamol	
		(Methocarbamol)	C
		Levothyroxine Sodium	
		(Levothyroxine	
		Sodium)	C
		Dyazide	
		(Hydrochlorothiazide	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

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Date:12/07/04ISR Number: 4522280-4Report Type:Expedited (15-DaCompany Report #2004081132

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		No Adverse Effect	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:12/07/04ISR Number: 4522603-6Report Type:Expedited (15-DaCompany Report #2004075203

Age:38 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 600 MG (300 MG, 2 IN 1 D), ORAL		Pelvic Venous Thrombosis Pulmonary Embolism	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
5 MG (1D), ORAL				Risperidone (Risperidone)	SS		ORAL
40 DROP (20 DROP, 2 IN 1 D), ORAL				Diazepam (Diazepam)	SS		ORAL
4 MG (2 MG, 2 IN 1 D), ORAL				Hydromorphone Hydrochloride (Hydromorphone Hydrochloride)	SS		ORAL

Date:12/07/04ISR Number: 4522604-8Report Type:Expedited (15-DaCompany Report #2004048216
Age:12 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation	Foreign	Neurontin			
		Condition Aggravated	Health	(Gabapentin)	PS		ORAL
900 MG (300		Drug Ineffective	Professional				
MG, 3 IN 1		Encephalopathy	Company				
D), ORAL		Grand Mal Convulsion	Representative	Carbamazepine			
		Psychomotor Retardation		(Carbamazepine)	C		

Date:12/07/04ISR Number: 4523147-8Report Type:Direct Company Report #CTU 233735
Age:35 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Condition Aggravated		Neurontin	PS		
400 MG 2 TID 35 DAY		Convulsion					
		Pharmaceutical Product					
		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/08/04ISR Number: 4522762-5Report Type:Expedited (15-DaCompany Report #2004098100
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	2400 MG (800 MG, 3 IN 1 D)	Drug Ineffective Weight Increased	Foreign Health Professional	Neurontin (Gabapentin)	PS		
				Clomipramine Hydrochloride (Clomipramine Hydrochloride)	C		
				Pygeum Africanum (Pygeum Africanum)	C		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		
				Levocarnitine (Levocarnitine)	C		

Date:12/08/04ISR Number: 4523015-1Report Type:Expedited (15-DaCompany Report #2004099600
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1200 MG, ORAL	Anxiety Blood Creatine	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Phosphokinase Increased	Professional	Diclofenac (Diclofenac Sodium)	SS		
		Chorea	Other	Doxepin Hydrochloride (Doxepin Hydrochloride)	C		
		Clonus		Fentanyl (Fentanyl)	C		
		Coordination Abnormal		Ibuprofen (Ibuprofen)	C		
		Depressed Level Of Consciousness		Aciclovir (Aciclovir)	C		
		Encephalitis Viral					
		Headache					
		Psychomotor Hyperactivity					
		Renal Failure					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Dizziness	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL		Grand Mal Convulsion Nausea	Professional Company Representative	Nitrazepam (Nitrazepam)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - TRANSPLACENTAL	DF Q DAY	Drug Exposure During Pregnancy	Foreign Health	Temesta	PS		
Initial or Prolonged TRAN-P		Ventricular Septal Defect	Professional	Neurontin	SS		
TRANSPLACENTAL	DF Q DAY		Other	Topimax	SS		
TRAN-P							
TRANSPLACENTAL	DF Q DAY						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

TRAN-P

Date:12/08/04ISR Number: 4524999-8Report Type:Expedited (15-DaCompany Report #2004099657
 Age:90 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Abasia	Consumer	Neurontin			
Other		Delusion		(Gabapentin)	PS		ORAL
ORAL							
		Dementia		Celebrex (Celecoxib)	SS		ORAL
ORAL							
		Illusion		Quinine	C		
		Mental Impairment		Centrum Silver			
				(Ascorbic Acid,			
				Calcium, Minerals			
				Nos, Retinol,			
				Tocopheryl Acetate,	C		
				Calcium	C		
				Ascorbic Acid	C		

Date:12/09/04ISR Number: 4523341-6Report Type:Expedited (15-DaCompany Report #2004100388
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hip Arthroplasty	Consumer	Neurontin			
		Osteoporosis		(Gabapentin)	PS		
900 MG (300							
MG, 3 IN 1		Shoulder Arthroplasty					
D),							
				Prednisone			
				(Prednisone)	C		

Date:12/09/04ISR Number: 4523358-1Report Type:Direct Company Report #CTU 234003
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other
300MG TID

Chest Pain

Gabapentin 300mg PS

ORAL

ORAL

Hydrochlorothiazide C
Ranitidine C
Atenolol C
Lisinopril C

Date:12/09/04ISR Number: 4523551-8Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 233992

Outcome

PT
Contusion
Convulsion
Dyskinesia
Electroencephalogram
Abnormal
Eye Rolling
Fall
Limb Injury
Mental Retardation
Severity Unspecified
Movement Disorder
Muscle Twitching
Pharmaceutical Product
Complaint

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Tremor Urinary Tract Infection					
400 MG T.I.D				Neurontin Generic & Brand Names	PS		
5 MG T.I.D.				Valium	SS		

Date:12/09/04ISR Number: 4523848-1Report Type:Direct Company Report #CTU 234005
Age:50 YR Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	400 MG TID		Affective Disorder		Gabapentin 400 Mg	PS		
			Drug Effect Decreased Dysphoria Pharmaceutical Product Complaint					

Date:12/09/04ISR Number: 4524003-1Report Type:Expedited (15-DaCompany Report #2004102771
Age: Gender:Female I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other			Haemorrhage Loss Of Consciousness Mental Disorder Overdose Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:12/09/04ISR Number: 4524079-1Report Type:Expedited (15-DaCompany Report #2004049712
Age:77 YR Gender:Male I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -			Confusional State	Foreign	Neurontin			

Initial or Prolonged 1200 MG (1 D)	Parkinson'S Disease	Company	(Gabapentin)	PS
		Representative	Lamaline (Belladonna Extract, Caffeine, Opium Tincture, Paracetamol)	C
			Ketoprofen (Ketoprofen)	C
			All Other Therapeutic Products (All Other Therapeutic Products)	C

Date:12/09/04ISR Number: 4524098-5Report Type:Expedited (15-DaCompany Report #2004102431
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cholestasis Condition Aggravated	Foreign Health	Neurontin (Gabapentin)	PS		
UNKNOWN	UNKNOWN	Microlithiasis Trigeminal Neuralgia	Professional Company Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/09/04ISR Number: 4524204-2Report Type:Expedited (15-DaCompany Report #2004097851
 Age:76 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Constipation Diarrhoea	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
300 MG (100 MG, 3 IN 1 D), ORAL		Infection Pancreatitis Urinary Tract Infection Weight Decreased	Professional	Sinemet (Marbidopa, Levodopa) Clopidogrel Sulfate (Clopidogrel Sulfate) Donepezil Hydrochloride (Donepezil Hydrochloride) Paracetamol (Paracetamol) Calcium (Calcium) Citalopram Hydrobromide) (Citalopram Hydrobromide) All Other Therrapeutic Products (All Other Therapeutic Products) Morphine (Morphine) Pantoprazole (Pantoprazole)	SS C C C C C C C		

Date:12/10/04ISR Number: 4523726-8Report Type:Direct
 Age: Gender:Female I/FU:I Company Report #USP 234097

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Effect Decreased		Gabapentin	PS		
300 MG AM 600							

Headache

Pharmaceutical Product
Complaint

Date:12/10/04ISR Number: 4525059-2Report Type:Expedited (15-DaCompany Report #2004076262

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abdominal Pain Upper Drug Ineffective Insomnia Loss Of Consciousness Malaise Pain Pain In Extremity Retching Vomiting	Consumer	Neurontin (Gabapentin) Hydromorphone Hydrochloride (Hydromorphone Hydrochloride) Morphine (Morphine) Methocarbamol (Methocarbamol) Rofecoxib (Refecoxib)	PS SS SS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/10/04ISR Number: 4525067-1Report Type:Expedited (15-DaCompany Report #2004101771
Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sudden Cardiac Death	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG (300 MG, 1 IN 1 D), ORAL			Company Representative				
				Carbamazepine (Carbamazepine)	C		

Date:12/10/04ISR Number: 4525342-0Report Type:Expedited (15-DaCompany Report #2004100270
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Delusion Illusion Impaired Driving Ability	Consumer	Neurontin (Gabapentin)	PS		

Date:12/10/04ISR Number: 4526546-3Report Type:Expedited (15-DaCompany Report #2004090950
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 20 MG (20 MG, Initial or Prolonged 1 IN 1 D), ORAL		Dermatitis Exfoliative	Foreign	Bextra (Valdecoxib)	PS		ORAL
		Drug Hypersensitivity	Health Professional				
		Rash Erythematous					
300 MG (300 MG, 1 IN 1 D), ORAL		Skin Exfoliation Stevens-Johnson Syndrome	Company Representative	Neurontin (Gabapentin)	SS		ORAL
				Colchine (Colchine)	C		

Date:12/13/04ISR Number: 4525964-7Report Type:Direct
Age:44 YR Gender:Female I/FU:I

Company Report #CTU 234204

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Convulsion		Gabapentin 400mg Five Times Daily	PS		ORAL
400MG	5X/D						
ORAL		Drug Effect Decreased					
		Pharmaceutical Product Complaint					

Date:12/13/04ISR Number: 4526095-2Report Type:Direct
Age:40 YR Gender:Male I/FU:I

Company Report #CTU 234183

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Disturbance In Attention		Neurontin	PS		
SEE IMAGE	6 MON						
Intervention to Prevent Permanent Impairment/Damage		Visual Disturbance					

Date:12/13/04ISR Number: 4526810-8Report Type:Expedited (15-DaCompany Report #2004059295
Age: Gender:Female I/FU:F

Outcome	PT
Other	Anxiety Asthma

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Dose	Duration	Condition	Report Source	Product	Role	Manufacturer	Route
		Balance Disorder Chronic Obstructive Pulmonary Disease					
1800 MG (1 D), ORAL		Condition Aggravated Hypersensitivity	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
		Memory Impairment					
2 IN 1 D		Polyneuropathy Sinus Disorder Tooth Extraction		Zyrtec-D 12 Hour (Pseudoephedrine, Cetirizine)	SS		
2 IN 1 D				Mirapex (Pramipexole) (Pramipexole)	SS		
2 IN 1 D				Clonazepam (Clonazepam)	SS		
				Mirtazapine (Mirtazapine)	SS		
				Amitriptyline (Amitriptyline)	SS		
				Montelukast Sodium (Montelukast Sodium)	C		
				Combivent (Ipratropium Bromide, Salbutamol Sulfate)	C		
				Mirtazapine (Mirtazapine)	C		
				Esomeprazole (Esomeprazole)	C		
				Combivent (Ipratropium Bromide, Salbutamol Sulfate)	C		
				Salbutamol (Salbutamol)	C		
				Calamine/Camphor/Dip henhydramine (Calamine, Camphor, Diphenhydramine)	C		
				Bupropion Hydrochloride (Bupropion			

Date:12/13/04ISR Number: 4526811-XReport Type:Expedited (15-DaCompany Report #2004102854
Age:38 YR Gender:Female I/FU:I

Outcome	PT
Death	Aspiration
Other	Blood Pressure
	Immeasurable
	Blood Urine Present
	Cell Death
	Chills
	Coma
	Completed Suicide
	Device Failure
	Dizziness
	Drug Toxicity
	Epistaxis
	Gastric Perforation

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Cholesterol Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 1 IN 1 D), ORAL		Flatulence Myocardial Infarction					
ORAL		Paraesthesia Tinnitus		Lipitor (Atovastatin)	SS		ORAL
				Lisinopril (Lisinopril)	SS		
				Rosuvastatin (Rosuvastatin)	SS		
				Simvastatin (Simvastatin)	SS		
10 MG (10 MG, 1 IN 1 D)				Potassium (Potassium)	SS		
				Pyridoxine Hydrochloride (Pyridoxine)			

Freedom Of Information (FOI) Report

Hydrochloride)	SS
Ezetimibe	
(Ezetimibe)	C
All Other	
Therapeutic Products	
(All Other	
Therapeutic	
Products)	C
Doxepin (Doxepin)	C
Acetylsalicylic Acid	
(Acetylsalicylic	
Acid)	C
Calcium Citrate	
(Calcium Citrate)	C
Estrogens	
(Estrogens)	C
Fish Oil (Fish Oil)	C
Vitamins (Vitamins)	C
Minerals Nos	
(Minerals Nos)	C
Vicon Forte (Folic	
Acid, Minerals Nos,	
Vitamins Nos)	C
Retinol (Retinol)	C
Tocopherol	
(Tocopherol)	C

Date:12/13/04ISR Number: 4526994-1Report Type:Expedited (15-DaCompany Report #2004087791
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Foreign	Gabapentin			
Other		Orthostatic Hypotension	Health Professional	(Gabapentin)	PS		
				Ciclosporin			
				(Ciclosporin)	C		
				Digoxin (Digoxin)	C		
				Omeprazole			
				(Omeprazole)	C		
				Amitriptyline			
				(Amitriptyline)	C		
				Bumetanide			
				(Bumetanide0	C		
				Lisinopril			
				(Lisinopril)	C		
				Atorvastatin			

(Atorvastatin)	C
Aciclovir	
(Aciclovir0	C
Prednisolone	
(Prednisolone)	C
Allopurinol	
(Allopurinol)	C
Mycophenolate	
Mofetil	
(Mycophenolate	
Mofetil)	C
Spirolactone	
(Spirolactone)	C
Bisoprolol	
(Bisoprolol)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/13/04ISR Number: 4527002-9Report Type:Expedited (15-DaCompany Report #2004103970

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Benign Gastrointestinal Neoplasm	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
	Haemorrhoids		Bromazepam (Bromazepam)	C		

Date:12/13/04ISR Number: 4527006-6Report Type:Expedited (15-DaCompany Report #2004103919

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Dislocation Of Vertebra Road Traffic Accident	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
			Bromazepam (Bromazepam)	C		

Date:12/14/04ISR Number: 4528915-4Report Type:Expedited (15-DaCompany Report #2004104252

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (300 MG, 1 IN 1 D), ORAL ORAL	Asthenia Clonic Convulsion Condition Aggravated Coordination Abnormal Difficulty In Walking Drug Ineffective Dyskinesia Lumbar Radiculopathy Memory Impairment Muscle Spasms Overdose Sensory Disturbance	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Methadone (Methadone)	SS		ORAL
			Atorvastatin (Atorvastatin)	C		
			Celecoxib (Celecoxib)	C		

Speech Disorder
Ulcer

Date:12/14/04ISR Number: 4528976-2Report Type:Expedited (15-DaCompany Report #2004103580
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Herpes Zoster	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Professional	All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:12/14/04ISR Number: 4529313-XReport Type:Expedited (15-DaCompany Report #2004104224
Age:63 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Antinuclear Antibody Positive
Other	Epstein-Barr Virus Infection
	Erythema Infectiosum

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Dose	Duration	Haematoma Parvovirus B19 Serology Positive	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1 D), ORAL		Platelet Disorder Thrombocytopenic Purpura	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Troxerutin (Troxerutin)	SS		ORAL
10 MG (10 MG, DAILY), ORAL				Zolpidem (Zolpidem)	SS		ORAL
ORAL				Irbesartan (Irbesartan)	SS		ORAL
300 MG (150 MG, 2 IN 1 D), ORAL				Tramadol Hydrochloride (Tramadol Hydrochloride)	SS		ORAL
ORAL				Caffeine / Opium/ Paracetamol (Caffeine, Opium Tincture, Paracetamol)	SS		ORAL

Date:12/14/04ISR Number: 4529912-5Report Type:Expedited (15-DaCompany Report #2004103044

Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Completed Suicide	Consumer	Neurontin (Gabapentin)	PS		

Date:12/15/04ISR Number: 4529926-5Report Type:Direct Company Report #CTU 234362
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
300 MG TID PO		Drug Effect Decreased		Neurontin (Generic	PS		ORAL

Date:12/15/04ISR Number: 4530218-9Report Type:Expedited (15-DaCompany Report #2004108052
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
1200 MG (300 MG, 4 IN 1 D)		Shoulder Operation Spinal Operation	Consumer	Neurontin (Gabapentin)	PS		

Date:12/15/04ISR Number: 4530219-0Report Type:Expedited (15-DaCompany Report #2004104706
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other							
900 MG (300 MG, 3 IN 1 D), ORAL		Blood Creatinine Increased Hypothyroidism Nephrectomy Off Label Use	Health Professional	Neurontin (Gabapentin) Atorvastatin (Atorvastatin) Acetylsalicylic	PS C		ORAL

Freedom Of Information (FOI) Report

Acid(Acetylsalicylic Acid) C
 Diazepam (Diazepam) C
 Fluoxetine Hydrochloride (Floxetine Hydrochloride) C
 Valproate Semisodium (Valproate Semisodium) C
 Metoprolol (Metoprolol) C
 Tolterodine L-Tartrate (Tolterodine L-Tartrate) C
 Lamotrigine (Lamotrigine) C
 All Other Therapeutic Products (All Other Therapeutic Products) C

Date:12/15/04ISR Number: 4530224-4Report Type:Expedited (15-DaCompany Report #2004068315
 Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Disorder Drug Ineffective	Consumer Health	Neurontin (Gabapentin)	PS		
1600 MG (800 MG, 2 IN 1 D)			Professional	Insulin (Insulin)	C		

Date:12/15/04ISR Number: 4530251-7Report Type:Expedited (15-DaCompany Report #2004097851
 Age:76 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Constipation Diarrhoea	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
300 MG (100							

Pancreatitis

Professional

Urinary Tract Infection

D), ORAL

Weight Decreased

Sinemet (Carbidopa, Levodopa)	SS
Clopidogrel Sulfate (Clopidogrel Sulfate)	C
Donepezil Hydrochloride (Donepezil Hydrochloride)	C
Paracetamol (Paracetamol)	C
Calcium (Calcium)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
All Other Therapeutic Products (All Other	

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Therapeutic
 Products) C
 Morphine (Morphine) C
 Pantoprazole
 (Pantoprazole) C

Date:12/15/04ISR Number: 4530266-9Report Type:Expedited (15-DaCompany Report #2004089442

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	800 MG (400 MG, 2 IN 1 D), ORAL	Breast Cancer Cardiac Arrest	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other	MG, 2 IN 1 D), ORAL	Cataract	Professional				
	10 MG (10 MG, 2 IN 1 D), ORAL	Chest Pain		Norvasc (Amlodipine)	SS		ORAL
	400 MG (200 MG, 2 IN 1 D)	Dementia Alzheimer'S Type					
		Dyspepsia					
		Feeding Disorder					
		Feeling Abnormal		Celebrex (Celecoxib)	SS		
		Kidney Infection					
	10 MG (10 MG, 2 IN 1 D), ORAL	Knee Arthroplasty Rotator Cuff Syndrome Urinary Tract Infection Vascular Bypass Graft		Donepezil Hydrochloride (Donepezil Hydrochloride)	SS		ORAL
		Weight Decreased					

Date:12/15/04ISR Number: 4530272-4Report Type:Expedited (15-DaCompany Report #2004104710

Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Grand Mal Convulsion	Foreign	Neurontin			

ORAL	Incontinence	Health	(Gabapentin)	PS	ORAL
		Professional	Carbamazepine (Carbamazepine)	C	
			Valproate Sodium (Valproate Sodium)	C	

Date:12/15/04ISR Number: 4530276-1Report Type:Expedited (15-DaCompany Report #2004103665
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Herpes Zoster	Foreign Health Professional	Neurontin (Tablets) (Gabapentin)	PS		
Other				All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:12/15/04ISR Number: 4530338-9Report Type:Expedited (15-DaCompany Report #2004104699
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Circulatory Collapse	Foreign Company Representative	Neurontin (Gabapentin)	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/15/04ISR Number: 4530340-7Report Type:Expedited (15-DaCompany Report #2004104608

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Post Procedural	Foreign	Neurontin			
ORAL		Haemorrhage	Health	(Gabapentin)	PS		ORAL
			Professional				

Date:12/15/04ISR Number: 4530399-7Report Type:Expedited (15-DaCompany Report #RENA-10959

Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Condition Aggravated	Foreign	Renagel	PS		ORAL
0.8 G TID PO							
Initial or Prolonged		Hyperkeratosis	Health	Beloc Zok	SS		
300 MG QD		Morphoea	Professional	Neurontin	SS		
			Company	Phos-Ex	C		
			Representative	Einsalpha	C		
			Other	Vigantoletten	C		
				Restex	C		
				Rohypnol	C		
				Marcumar	C		

Date:12/16/04ISR Number: 4530578-9Report Type:Expedited (15-DaCompany Report #2004081549

Age:84 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Balance Disorder	Foreign	Neurontin (Tablets)			
1500 MG (300		Gait Disturbance	Health	(Gabapentin)	PS		ORAL
MG, 5 IN 1		Speech Disorder	Professional				
D), ORAL		Sudden Hearing Loss					

All Other
Therapeutic Products
(All Other
Therapeutic

Products) C
 Flecainide Acetate C
 (Flecainide Acetate) C
 Atorvastatin Calcium
 (Atorvastatin
 Calcium) C
 Clopidogrel Sulfate
 (Clopidogrel
 Sulfate) C

Date:12/16/04ISR Number: 4531152-0Report Type:Expedited (15-DaCompany Report #2004105931
 Age:79 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Confusional State Coordination Abnormal	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
		Depressed Level Of Consciousness Muscle Twitching	Professional	Cyclizine (Cyclizine)	C		
				Diclofenac (Diclofenac)	C		
				Furosemide (Furosemide)	C		
				Omeprazole (Omeprazole)	C		
				Bisoprolol			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Bisoprolol) C
 Lacidipine
 (Lacidipine) C
 Morphine Sulfate
 (Morphine Sulfate) C

Date:12/16/04ISR Number: 4531647-XReport Type:Direct
 Age:69 YR Gender:Female I/FU:I

Company Report #CTU 234503

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypersensitivity Pharmaceutical Product Complaint		Neurontin	PS		

Date:12/16/04ISR Number: 4531839-XReport Type:Expedited (15-DaCompany Report #2004088763
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Economic Problem Pain	Consumer	Neurontin (Gabapentin) All Other Therapeutic Products (All Other Therapeutic Products)	PS SS		

Date:12/16/04ISR Number: 4677943-7Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 243029

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nausea		Gabapentin	PS		ORAL
100 MG PO		Pharmaceutical Product Complaint					

Date:12/17/04ISR Number: 4530642-4Report Type:Expedited (15-DaCompany Report #FR-ABBOTT-04P-056-0283040-00
 Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Encephalopathy		Kaletra Soft Gelatin			
Hospitalization -		Hepatic Failure		Capsules	PS		
Initial or Prolonged				Zidovudine			
				W/Lamivudine	SS		ORAL
				Ribavirin	SS		ORAL
				Perinterferon	SS		
SUBCUTANEOUS							
				Buprenorphine	SS		ORAL
				Gabapentin	SS		ORAL

Date:12/17/04ISR Number: 4532221-1Report Type:Expedited (15-DaCompany Report #2004-10-0038
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		General Physical Health	Foreign	Viraferonpeg (Peg			
Hospitalization -		Deterioration	Health	Interferon Alfa-2b			
Initial or Prolonged		Hepatic Encephalopathy	Professional	Recombinant) Redipen	PS		
SUBCUTANEOUS	120 MCG						
SUBCUTANEOUS							
		Hepatic Failure	Company				
			Representative	Rebetol (Ribavirin)			
			Other	Capsules	SS		ORAL
1000 MG QD							

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Freedom Of Information (FOI) Report

ORAL			Neurontin Capsules	SS		ORAL
400 MG TID						
ORAL			Kaletra (Lopinavir/Ritonavir) Capsules	SS		ORAL
3 DOSES BID						
ORAL			Combivir (Lamivudine/Zidovudine) Capsules	SS		ORAL
150 MG BID						
ORAL			Subutex (Buprenorphine Hcl) Tablets	SS		ORAL
8 MG QD ORAL						
			Zeclar (Clarithromycin)	C		
			Minocycline	C		
			Aldactone	C		

Date:12/17/04ISR Number: 4532326-5Report Type:Expedited (15-DaCompany Report #2004083497
Age:69 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	200 MG (100 MG, 2 IN 1	Atrial Fibrillation Cerebrovascular Accident	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other		Dizziness	Professional				
D), ORAL		Drug Ineffective					
		Dyspnoea		Omeprazole			
		Dyspnoea Exertional		(Omeprazole)	C		
		Economic Problem		Furosemide			
		Transient Ischaemic Attack		(Furosemide)	C		

Date:12/17/04ISR Number: 4532351-4Report Type:Expedited (15-DaCompany Report #2004104252
Age:40 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Clonic Convulsion Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (300 MG, 1 IN 1 D), ORAL		Coordination Abnormal Difficulty In Walking					
ORAL		Drug Ineffective Dyskinesia		Methadone (Methadone)	SS		ORAL
		Dysphasia Feeling Abnormal Memory Impairment Mental Disorder Muscle Spasms Overdose Photosensitivity Reaction Ulcer Verbigeration		Atorvastatin (Atorvastatin) Celecoxib (Celecoxib)	C C		

Date:12/17/04ISR Number: 4532352-6Report Type:Expedited (15-DaCompany Report #2004106658
Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death		Completed Suicide Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:12/17/04ISR Number: 4532353-8Report Type:Expedited (15-DaCompany Report #2004106719
Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin			
Life-Threatening		Drug Ineffective		(Gabapentin)	PS		
Other							

Date:12/19/04ISR Number: 4679576-5Report Type:Direct Company Report #CTU 243019
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Muscle Spasms		Gabapentin	PS		ORAL
300 MG ONE							
BID PO							

Date:12/20/04ISR Number: 4531775-9Report Type:Expedited (15-DaCompany Report #GB-ROCHE-389208
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous		Clonazepam	PS	Roche	ORAL
TAKEN UNTIL		Drug Exposure During		Detrusitol	SS		ORAL
THE 7TH WEEK		Pregnancy					
OF PREGNANCY.		Pregnancy		Zopiclone	SS		ORAL
				Gabapentin	SS		ORAL
				Propranolol	SS	Roche	ORAL
UNKNOWN	TAKEN EVERY			Sertraline	SS		

DAY.

Date:12/20/04ISR Number: 4534032-XReport Type:Direct Company Report #CTU 234662
Age:9.5 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Gabapentin	300 Mg		
		Pharmaceutical Product		Teva, Usa		PS	Teva, Usa
300 MG	ONCE						ORAL
		Complaint					
AT BEDTIM							
		Tremor					
ORAL							

Date:12/20/04ISR Number: 4536117-0Report Type:Expedited (15-DaCompany Report #2004010780
Age:69 YR Gender:Male I/FU:F

Outcome	PT
Disability	Actinic Keratosis
Other	Aldolase Increased
	Arthralgia
	Blood Cholesterol Increased
	Blood Glucose Increased
	Blood Potassium Increased
	Blood Sodium Increased
	Blood Urine Present
	Bundle Branch Block Left
	Diverticulum
	Drug Hypersensitivity
	Emotional Disorder
	Fatigue
	Feeling Cold
	Head Injury

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hyperhidrosis Impaired Work Ability Injury	Consumer	Lipitor (Atorvastatin)	PS		
10 MG (DAILY)		Insomnia Joint Crepitation	Consumer				
300 MG, ORAL		Joint Range Of Motion Decreased		Neurontin (Gabapentin)	SS		ORAL
(DAILY), ORAL		Kyphosis Low Density Lipoprotein Increased Mental Disorder		Hyzaar (Hydrochlorothiazide , Losartan Potassium)	SS		ORAL
		Muscle Atrophy Myopathy Neck Injury Nervous System Disorder Nervousness Neuralgia Ph Urine Decreased Polymyositis Prostatic Specific Antigen Increased Protein Urine Present Rectal Polyp Sinus Headache Spinal Column Stenosis Urinary Tract Infection Visual Acuity Reduced Weight Decreased		Influenza Vaccine (Influenza Vaccine) All Other Therapeutic Products (All Other Therapeutic Products) Vicodin (Hydrocodone Bitartrate, Paracetamol) Acetylsalicylic Acid (Acetylsalicylic Acid) Amlodipine Besilate (Amlodipine Besilate)	C C C C		

Date:12/20/04ISR Number: 4536148-0Report Type:Expedited (15-DaCompany Report #2004106702

Age:35 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		

Date:12/20/04ISR Number: 4536154-6Report Type:Expedited (15-DaCompany Report #2004107123

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Withdrawal Syndrome Surgery	Consumer	Neurontin (Gabapentin)	PS		ORAL
4000 MG, ORAL							

Date:12/20/04ISR Number: 4536167-4Report Type:Expedited (15-DaCompany Report #2004112247
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ankle Fracture Balance Disorder	Consumer Company	Neurontin (Gabapentin)	PS		
900 MG (1 D)							
		Fall	Representative	Oxcarbazepine (Oxcarbazepine)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/20/04ISR Number: 4536175-3Report Type:Expedited (15-DaCompany Report #2004107078
 Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200 MG (400 MG, 3 IN 1 D), ORAL	Cardiac Failure Sudden Death	Consumer	Neurontin (Gabapentin)	PS		ORAL
				All Other Therapeutic Products (All Other Therapeutic Products)	SS		

Date:12/20/04ISR Number: 4536333-8Report Type:Expedited (15-DaCompany Report #2004080275
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	200 MG (100 MG, 2 IN 1 D)	Renal Cyst	Consumer	Neurontin (Gabapentin)	PS		
				Aricept (Donepezil)	SS		

Date:12/20/04ISR Number: 4536449-6Report Type:Expedited (15-DaCompany Report #2004106809
 Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury	Consumer	Neurontin (Gabapentin)	PS		

Date:12/20/04ISR Number: 4536659-8Report Type:Expedited (15-DaCompany Report #2004105620
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Aggression Agitation	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other MG, 3 IN 1		Anxiety	Professional				
D), ORAL		Condition Aggravated					
20 MG (20 MG, 1 IN 1 D), ORAL		Crying Delirium Histrionic Personality Disorder Mental Disorder Nightmare		Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		ORAL
30 MG (30 MG, 1 IN 1 D), ORAL		Suicidal Ideation Trismus Weight Decreased		Mianserin Hydrochloride (Mianserin Hydrochloride)	SS		ORAL
1 MG (1 MG, 1 IN 1 D), ORAL				Zolpidem (Zolpidem)	SS		ORAL
SUBCUTANEOUS 1 IN 1 D), SUBCUTANEOUS	25 MG (25 MG,			Fentanyl (Fentanyl)	SS		
60 MG (10 MG, 6 IN 1 D), ORAL				Morphine Sulfate (Morphine Sulfate)	SS		ORAL

Freedom Of Information (FOI) Report

Macrogol (Macrogol) C
 Eugynon
 (Ethinylestradiol,
 Levonorgestrel) C

Date:12/20/04ISR Number: 4536670-7Report Type:Expedited (15-DaCompany Report #2004087791
 Age:61 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Loss Of Consciousness	Foreign	Gabapentin			
Disability		Orthostatic Hypotension	Health	(Gabapentin)	PS		
Other			Professional	Ciclosporin			
				(Ciclosporin)	C		
				Digoxin (Digoxin)	C		
				Omeprazole			
				(Omeprazole)	C		
				Amitriptyline			
				(Amitriptyline)	C		
				Bumetanide			
				(Bumetanide)	C		
				Lisinopril			
				(Lisinopril)	C		
				Atorvastatin			
				(Atorvastatin)	C		
				Aciclovir			
				(Aciclovir)	C		
				Prednisolone			
				(Prednisolone)	C		
				Allopurinol			
				(Allopurinol)	C		
				Mycophenolate			
				Mofetil			
				(Mycophenolate			
				Mofetil)	C		
				Spironolactone			
				(Spironolactone)	C		
				Bisoprolol			
				(Bisoprolol)	C		

Date:12/20/04ISR Number: 4536677-XReport Type:Expedited (15-DaCompany Report #2004108386
 Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	100 MG (100 MG, 1 IN 1 D), ORAL	Agitation Amnesia Mental Disorder Syncope Tremor Urinary Tract Disorder	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
				Zolpidem (Zolpidem) Clonazepam (Clonazepam)	C C		

Date:12/20/04ISR Number: 4536984-0Report Type:Expedited (15-DaCompany Report #2004110883
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Acute Respiratory Distress Syndrome	Foreign Health Professional	Neurontin (Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:12/20/04ISR Number: 4536992-XReport Type:Expedited (15-DaCompany Report #2004110436
Age:88 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Fall	Foreign	Gabapentin			
Life-Threatening		Hypotension	Health	(Gabapentin)	PS		ORAL
Hospitalization -		Renal Failure Acute	Professional				
MG, 1 IN 1		Respiratory Failure	Company				
Initial or Prolonged			Representative	Amiodarone			
D), ORAL				(Amiodarone)	C		
Other				Furosemide			
				(Furosemide)	C		
				Tiotropium			
				(Tiotropium)	C		
				Salmeterol			
				(Salmeterol)	C		
				Salbutamol			
				(Salbutamol)	C		
				Beclometasone			
				(Beclometasone)	C		

Date:12/20/04ISR Number: 4679569-8Report Type:Direct Company Report #CTU 243016
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Pruritus		Gabapentin Purepac -			
		Rash		Elizabeth, N.J.	PS	Purepac	ORAL
300 MG PO BID							

Date:12/21/04ISR Number: 4533160-2Report Type:Expedited (15-DaCompany Report #FR-RB-1183-2004
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hepatic Encephalopathy		Peginterferon			
Hospitalization -		Hepatic Failure		Alfa-2b	PS		
SUBCUTANEOUS							
Initial or Prolonged				Ribavirin	SS		ORAL

Gabapentin	SS	ORAL
Kaletra	SS	ORAL
Combivir	SS	ORAL
Buprenorphine		
Hydrochloride	SS	ORAL
Clarithromycin	C	
Minocycline	C	
Spiroinolactone	C	

UNKNOWN UNKNOWN
 UNKNOWN UNKNOWN
 UNKNOWN UNKNOWN

Date:12/21/04ISR Number: 4534136-1Report Type:Direct Company Report #CTU 234756
 Age:22 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nausea		Neurontin 300mg	PS		
4POAM,							
3PONOON &							
4POHS							

Date:12/21/04ISR Number: 4534475-4Report Type:Direct Company Report #CTU 234760
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pruritus Rash		Gabapentin Mfr Purepac - Elizabeth,			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

300 MG PO BID N.J. PS Purepac ORAL

Date:12/21/04ISR Number: 4537331-0Report Type:Expedited (15-DaCompany Report #MK200411-0251-1
Age:45 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Hydrocodone/Apap Tablets (Strength Unk) Gabapentin	PS SS		

Date:12/21/04ISR Number: 4537334-6Report Type:Expedited (15-DaCompany Report #MK200411-0250-1
Age:23 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse	Literature Health Professional	Hydrocodone/Apap Tablets (Strength Unk) Gabapentin	PS SS		

Date:12/21/04ISR Number: 4537896-9Report Type:Expedited (15-DaCompany Report #MK200412-0140-1
Age:35 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Multiple Drug Overdose	Literature Health Professional	Tramadol Hydrochloride Citalopram Gabapentin	PS SS SS		

Date:12/21/04ISR Number: 4538106-9Report Type:Expedited (15-DaCompany Report #2004081549
Age:84 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Balance Disorder	Foreign	Neurontin (Tablets			

1500 MG (300	Dysphasia	Health)(Gabapentin)	PS	ORAL
MG, 5 IN 1	Gait Disturbance	Professional			
D), ORAL	Hypoacusis				
	Memory Impairment		All Other Therapeutic Products (All Other Therapeutic Products)	C	
			Flecainide Acetate (Flecainide Acetate)	C	
			Atorvastatin Calcium (Atorvastatin Calcium)	C	
			Clopidogrel Sulfate (Clopidogrel Sulfate)	C	

Date:12/21/04ISR Number: 4538118-5Report Type:Expedited (15-DaCompany Report #2004096313
Age: Gender: I/FU:F

Outcome	PT	Report Source
Other	Circulatory Collapse	Consumer Health

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
		Neurontin (Gabapentin)	PS		
		Interferon Beta (Interferon Beta)	SS		

Date:12/21/04ISR Number: 4538156-2Report Type:Expedited (15-DaCompany Report #2004111689
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG (600		Bone Disorder					
MG, 4 IN 1		Convulsion					
D), ORAL		Feeling Abnormal					
		Hypersensitivity		Gabapentin (Gabapentin)	SS		ORAL
2400 MEQ (600		Malaise					
MG, 4 IN 1		Pain					
D), ORAL		Pharmaceutical Product					
		Complaint		Clonazepam (Clonazepam)	C		
		Skin Disorder					

Date:12/21/04ISR Number: 4538214-2Report Type:Expedited (15-DaCompany Report #2004076256
Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Haemodialysis	Health	Neurontin (Gabapentin)	PS		
Hospitalization -		Renal Failure Acute	Professional				
1200 MG (600							
Initial or Prolonged		Renal Tubular Necrosis	Company				
MG, 2 IN 1 D)							
Other		Urinary Tract Infection	Representative	Gatifloxacin			

(400 MG),

ORAL

(Gatifloxacin)

SS

ORAL

Pravastatin Sodium (Pravastatin Sodium)	C
Rosiglitazone Maleate (Rosiglitazone Maleate)	C
Gezor (Hydrochlorothiazide , Quinapril Hydrochloride)	C
Risedronate Sodium (Rosiglitazone Maleate)	C
Nortriptyline (Nortriptyline)	C
Raloxifene Hydrochloride (Raloxifene Hydrochloride)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/21/04ISR Number: 4538271-3Report Type:Expedited (15-DaCompany Report #2004107080
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dyspnoea Hypersensitivity	Consumer	Celebrex (Celecoxib)	PS		ORAL
400 MG (200 MG, 2 IN 1 D), ORAL		Sensation Of Foreign Body Swelling Face					
300 MG (300 MG, 1 IN 1 D), ORAL				Neurontin (Gabapentin)	SS		ORAL
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		

Date:12/21/04ISR Number: 4538415-3Report Type:Expedited (15-DaCompany Report #2004CG02461
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required Intervention to Prevent Permanent 1 DF QD PO Impairment/Damage		Cerebrovascular Accident Facial Pain Facial Palsy	Foreign Health Professional	Zomig Neurontin Zoloft	PS SS SS		ORAL
			Other	Melodia Ibuprofen Lexomil	SS C C		

Date:12/21/04ISR Number: 4538437-2Report Type:Expedited (15-DaCompany Report #MK200412-0029-1
Age:49 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Literature	Methadose Methadone			

Intentional Misuse

Health
Professional

Hy Oral Con Usp
Gabapentin
Methocarbamol

PS
SS
SS

Date:12/21/04ISR Number: 4538460-8Report Type:Expedited (15-DaCompany Report #MK200412-0027-1
Age:38 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Intentional Misuse	Literature Health Professional	Methadose@ Methadone Hy Oral Con Usp Fluoxetine Gabapentin			
					PS		
					SS		
					SS		

Date:12/22/04ISR Number: 4535632-3Report Type:Direct Company Report #CTU 234960
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ONE PO TID		Abdominal Pain Constipation Insomnia		Gabapentin 300 Mg	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/22/04ISR Number: 4539058-8Report Type:Expedited (15-DaCompany Report #2004009355

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG (600 Other MG, 3 IN 1 D), ORAL	Abdominal Pain Back Pain Chest Pain Erectile Dysfunction	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
20 MG (DAILY), ORAL	Facial Pain Feeling Cold Gastrooesophageal Reflux		Lipitor (Atorvastatin)	SS		ORAL
1000 MG (BID), ORAL	Disease Gingival Pain Glossodynia Hypersomnia Irritability Nasopharyngitis Neck Pain Nervous System Disorder		Lithium (Lithium) (Lithium) Naproxen (Naproxen) Rofecoxib (Rofecoxib) Amoxicillin (Amoxicillin)	SS SS SS SS		ORAL
	Oesophageal Spasm Pain In Jaw Sleep Disorder Somnolence Tooth Abscess Treatment Noncompliance Tremor		Levothyroxine Sodium (Levothyroxine Sodium) Vitamins (Vitamins) Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C C		

Date:12/22/04ISR Number: 4539237-XReport Type:Expedited (15-DaCompany Report #2004087791

Age:61 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other	Fall Loss Of Consciousness Orthostatic Hypotension	Foreign Health Professional	Gabapentin (Gabapentin) Ciclosporin (Ciclosporin)	PS C		

Digoxin (Digoxin)	C
Omeprazole	
(Omeprazole)	C
Amitriptyline	
(Amitriptyline)	C
Bumetanide	
(Bumetanide)	C
Lisinopril	
(Lisinopril)	C
Atorvastatin	
(Atorvastatin)	C
Aciclovir	
(Aciclovir)	C
Prednisolone	
(Prednisolone)	C
Allopurinol	
(Allopurinol)	C
Mycophenolate	
Mofetil	
(Mycophenolate	
Mofetil)	C
Spironolactone	
(Spironolactone)	C
Bisoprolol	
(Bisoprolol)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/22/04ISR Number: 4539289-7Report Type:Expedited (15-DaCompany Report #2004077798
Age:62 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Crying	Foreign	Neurontin (Tablets)			
ORAL		Fear	Consumer	(Gabapentin)	PS		ORAL
		Hallucination	Health Professional				

Date:12/22/04ISR Number: 4539766-9Report Type:Expedited (15-DaCompany Report #2004064324
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Akathisia	Consumer	Neurontin			
Other		Anger		(Gabapentin)	PS		
1200 MG (600		Anxiety					
MG, 2 IN 1 D)		Asthenia		Paroxetine			
		Completed Suicide		Hydrochloride			
		Decreased Interest		(Paroxetine			
		Disturbance In Attention		Hydrochloride)	SS		
		Drug Ineffective		Paroxetine			
		Feeling Abnormal		Hydrochloride			
		Hallucination, Auditory		(Paroxetine			
		Insomnia		Hydrochloride)	C		
		Irritability		Risperidone			
		Memory Impairment		(Risperidone)	C		
		Mood Swings		Hydroxyzine Embonate			
		Obsessive-Compulsive		(Hydroxyzine			
		Disorder		Embonate)	C		
		Ochlophobia		Venlafaxine			
		Panic Attack		Hydrochloride			
		Self-Medication		(Venlafaxine			
		Stress		Hydrochloride)	C		
		Treatment Noncompliance		Quetiapine Fumarate			
				(Quetiapine			
				Fumarate)	C		
				Famotidine			
				(Famotidine)	C		
				Narine Repetabs			
				(Loratadine,			

Pseudoephedrine Sulfate) C
 Lansoprazole (Lansoprazole) C
 Clarithromycin (Clarithromycin) C

Date:12/23/04ISR Number: 4536202-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0360023A
 Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2 YR		Hepatic Encephalopathy	Combivir	PS	Glaxosmithkline	ORAL
Hospitalization -	2 YR		Liver Disorder	Kaletra	SS		ORAL
Initial or Prolonged				Rebetol	SS		ORAL
1000MG per							
day				Alpha 2b Interferon	SS		
SUBCUTANEOUS	120MCG per						
day				Subutex	SS		ORAL
8MG Per day				Neurontin	SS		ORAL
400MG Three							
times per day	2 YR						

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Freedom Of Information (FOI) Report

UNKNOWN	Zeclar	SS
UNKNOWN	Minocycline	SS
UNKNOWN	Aldactone	SS

Date:12/23/04ISR Number: 4537277-8Report Type:Expedited (15-DaCompany Report #PHNU2004DE04127
 Age: Gender: I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 80 mg, QD 44640MIN	Alanine Aminotransferase Increased		Locol	PS	Novartis Sector: Pharma	
4DF/week	Aspartate Aminotransferase Increased Blood Creatine		Sertraline Piracetam Gabapentin Testosteron	SS SS SS SS		
	Phosphokinase Increased Blood Creatine Phosphokinase Mb Dyspnoea Muscular Weakness Myalgia Rhabdomyolysis					

Date:12/23/04ISR Number: 4539327-1Report Type:Expedited (15-DaCompany Report #001-0945-M0200657
 Age:44 YR Gender: I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability Other	Abdominal Pain Upper Affect Lability Agitation Anger Arthralgia Arthropod Bite Asthenia Balance Disorder Blood Thyroid Stimulating Hormone Decreased Brief Psychotic Disorder

With Marked Stressors
Carpal Tunnel Syndrome
Change Of Bowel Habit
Chest Pain
Chromaturia
Constipation
Convulsion
Coordination Abnormal
Deafness
Death Of Friend
Decreased Appetite
Depression
Diarrhoea
Dissociative Disorder
Dizziness
Dry Mouth
Dysarthria
Dyskinesia
Dyspepsia
Dysphagia
Dyspnoea
Dysuria

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Freedom Of Information (FOI) Report

Ear Pain
Electrocardiogram
Abnormal
Eustachian Tube
Dysfunction
Fall
Fatigue
Feeling Jittery
Fibrocystic Breast
Disease
Flat Affect
Gastrooesophageal Reflux
Disease
Goitre
Granuloma
Hallucination, Visual
Hand Fracture
Head Injury
Headache
Heart Rate Increased
Heart Rate Irregular
Hordeolum
Hot Flush
Hypoaesthesia
Hypothyroidism
Impaired Healing
Imprisonment
Increased Tendency To
Bruise
Increased Upper Airway
Secretion
Insomnia
Intervertebral Disc
Degeneration
Intervertebral Disc
Disorder
Irritability
Joint Injury
Joint Sprain
Joint Swelling
Laboratory Test Abnormal
Laryngitis
Lumbar Radiculopathy
Memory Impairment
Meniere'S Disease
Mental Disorder
Mouth Breathing

Nasal Discomfort
Nasal Ulcer
Nausea
Nervous System Disorder
Neuropathy
Nystagmus
Osteoarthritis
Otitis Media
Paraesthesia
Pharyngitis
Poor Peripheral
Circulation
Rash
Rash Erythematous

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration		Report Source	Product	Role	Manufacturer	Route
		Rhinitis Allergic Road Traffic Accident Sedation					
1800 MG		Sinus Disorder Sneezing	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
(DAILY), ORAL		Somnolence	Professional				
		Staring		Lamotrigine			
		Suicidal Ideation		(Lamotrigine0	C		
		Syncope		Lithium Carbonate			
		Tardive Dyskinesia		(Lithium Carbonate0	C		
		Tendonitis		Clonazepam(Clonazepa			
		Thirst		m)	C		
		Tinnitus		Methyphenidate			
		Tremor		Hydrochloride			
		Unresponsive To Verbal Stimuli		(Methylphenidate			
		Urinary Tract Infection		Hydrochloride)	C		
		Vaginal Candidiasis		Levothyroxine Sodium			
		Vertigo		(Levothyroxine			
		Vertigo Positional		Sodium)	C		
		Vestibular Neuronitis		Setraline			
		Vestibulitis		Hydrochloride			
		Viral Labyrinthitis		(Sertraline			
		Vision Blurred		Hydrochloride0	C		
		Weight Decreased		Pilocarpine			
		Weight Increased		Hydrochloride			
		White Blood Cell Count Decreased		(Pilocarpine			
		Wound Infection		Hydrochloride)	C		
				Metoprolo Succinate			
				(Metoprolol			
				Succinate0	C		
				Lansoprazole			
				(Lansoprazole)	C		
				Hyoscyamine Sulfate			
				(Hyoscyamine			
				Sulfate)	C		
				Diltiazem			
				Hydrochloride			
				(Diltiazem			
				Hydrochloride)	C		
				Bupropion			
				Hydrochloride			
				(Bupropion			
				Hydrochloride)	C		
				Celebrex (Celecoxib0	C		

Quetiapine Fumarate (Quetiapine Fumarate0	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide0	C
Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C
Olanzapine (Olanzapine)	C
Omeprazole (Omeprazole0	C

Freedom Of Information (FOI) Report

Nasal Preparations
 (Nasal Preparations0 C
 Nefazodone
 Hydrochloride
 (Nefazodone
 Hydrochloride) C
 Totolin
 (Guaifenesin,
 Phenylpropanolamine
 Hydrochloride) C
 Meclozine
 (Meclozine) C
 Fluticasone
 Propionate
 (Fluticasone
 Propionate) C
 Trimethobenzamide
 Hydrochloride
 (Trimethobenzamide
 Hydrochloride) C
 Respaire -Sr -120
 (Guaifenesin,
 Pseudoephedrine
 Hydrochloride) C
 Risperidone
 (Risperidone) C

Date:12/23/04ISR Number: 4539745-1Report Type:Expedited (15-DaCompany Report #2004114461

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Guillain-Barre Syndrome	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional				

Date:12/23/04ISR Number: 4539943-7Report Type:Expedited (15-DaCompany Report #2004113561

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disorientation	Consumer	Neurontin			

(400 MG),	Dysstasia	(Gabapentin)	PS	ORAL
ORAL	Incoherent			
ORAL		Escitalopram (Escitalopram)	SS	ORAL
		Alprazolam (Alprazolam)	C	
		Meloxicam (Meloxicam)	C	
		Tizanidine Hydrochloride (Tizanidine Hydrochloride)	C	
		Quetiapine Fumarate (Quetiapine Fumarate)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/23/04ISR Number: 4539945-0Report Type:Expedited (15-DaCompany Report #2004114433

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG, 3 IN 1 D), ORAL		Laceration Self Injurious Behaviour	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Lamotrigine (Lamotrigine)	C		
				Benzatropine Mesilate (Benzatropine Mesilate)	C		
				Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		
				Clonazepam (Clonazepam)	C		
				Risperidone (Risperidone)	C		

Date:12/23/04ISR Number: 4539955-3Report Type:Expedited (15-DaCompany Report #2004110716

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other (2 IN 1 D), ORAL		Abnormal Behaviour Amnesia Anger Papilloma Viral Infection Vulval Cancer	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:12/23/04ISR Number: 4539956-5Report Type:Expedited (15-DaCompany Report #2004111537

Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2700 MG (900 Other MG, 3 IN 1 D), ORAL	Balance Disorder Condition Aggravated Confusional State Mobility Decreased Myoclonus Neurological Symptom Paraesthesia Paraneoplastic Syndrome Peripheral Coldness Polyneuropathy	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:12/23/04ISR Number: 4539957-7Report Type:Expedited (15-DaCompany Report #2004111569
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Other	Abasia Amnesia Joint Dislocation Pain Post Procedural Complication Somnolence

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Stupor

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
400 MG (100 MG, 4 IN 1 D), ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
			Hytrin (Terazosin Hydrochloride)	C		
			Potassium (Potassium)	C		
			Synthroid (Levothyroxine Sodium)	C		

Date:12/23/04ISR Number: 4539958-9Report Type:Expedited (15-DaCompany Report #2004114437
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	1200 MG (1 D)	Condition Aggravated Fibromyalgia	Consumer	Neurontin (Gabapentin)	PS		
		Macular Degeneration Pain Treatment Noncompliance		Dyazide (Hydrochlorothiazide , Triamterene)	C		

Date:12/23/04ISR Number: 4540046-6Report Type:Expedited (15-DaCompany Report #2004114450
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Hepatorenal Failure	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		

Methadone
(Methadone) SS
All Other
Therapeutic Products
(All Other
Therapeutic
Products) SS

Date:12/27/04ISR Number: 4541537-4Report Type:Expedited (15-DaCompany Report #2004113535
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anticonvulsant Drug Level	Foreign	Neurontin			
Other		Decreased	Health	(Gabapentin)	PS		
		Condition Aggravated	Professional	Valproate Sodium			
		Grand Mal Convulsion	Company	(Valproate Sodium)	SS		
			Representative				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/27/04ISR Number: 4541712-9Report Type:Expedited (15-DaCompany Report #DEU-2004-0001150

Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction International Normalised Ratio Decreased	Foreign Health Professional Other	Oxygesic 10 Mg (Oxycodone Hydrochloride) Cr Tablet	PS		ORAL
SEE IMAGE							
				Marcumar (Phenprocoumon)	SS		ORAL
TABLET, ORAL							
				Amitriptyline (Amitriptyline)	SS		ORAL
10 MG, BID,							
ORAL							
				Gabapentin (Gabapentin)	SS		ORAL
1200 MG,							
DAILY, ORAL							
				Isosorbide Mononitrate	C		
				Pantoprazol (Pantoprazole Sodium)	C		
				Beloc-Zoc Forte (Metoprolol Succinate)	C		
				Atorvastatin Calcium (Atorvastatin Calcium)	C		

Date:12/27/04ISR Number: 4542485-6Report Type:Expedited (15-DaCompany Report #2004099657

Age:90 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Abasia	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Blindness					
ORAL							
		Delusion		Celebrex (Celecoxib)	SS		ORAL
ORAL							

Illusion
Thinking Abnormal

Quinine (Quinine) C
Centrum Silver (See
Image) C
Calcium(Calcium) C
Ascorbic Acid
(Ascorbic Acid) C

Date:12/27/04ISR Number: 4542491-1Report Type:Expedited (15-DaCompany Report #2004113565

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Consumer	Neurontin			
Other		Drug Ineffective		(Gabapentin)	PS		
1200 MG (400		Drug Toxicity					
MG 3 IN 1 D)		Fatigue		Dilantin Suspension			
		Gait Disturbance		(Phenytoin Sodium)	SS		
		Grand Mal Convulsion		Carbamazepine			
		Pain In Extremity		(Carbamazepine)	SS		
		Restless Legs Syndrome		Ranitidine			
		Sluggishness		Hydrochloride			
		Vision Blurred		(Ranitidine			
				Hydrochloride)	C		
				Dilantin Suspension			
				(Phenytoin Sodium)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/28/04ISR Number: 4539453-7Report Type:Direct
Age: Gender: I/FU:I

Company Report #USP 57013

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Medication Error		Neurontin	PS	Pfizer Pharmaceuticals Ltd/Parke-Davis	
CAPSULES							

Date:12/28/04ISR Number: 4540344-6Report Type:Direct
Age:75 YR Gender:Female I/FU:I

Company Report #CTU 235189

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dyspnoea		Neurontin	PS		
Death							

Date:12/28/04ISR Number: 4540589-5Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 235237

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Gabapentin	PS		ORAL
Other		Pain		Gliduride	C		
300 MG PO QHS 1 MON							
Pharmaceutical Product							
Complaint							
Nexium							
Fosamox							

Date:12/28/04ISR Number: 4540591-3Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 235238

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Hypersensitivity		Neurontin 400 One Po	PS		ORAL
Other		Pharmaceutical Product		Qd			
ONE PO QD							
Complaint							
[LIFETIME]							
Rash							

Date:12/29/04ISR Number: 4542912-4Report Type:Expedited (15-DaCompany Report #2004117393
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Drug Ineffective	Consumer	Neurontin			
Other		Pain		(Gabapentin)	PS		
		Suicide Attempt					

Date:12/29/04ISR Number: 4542915-XReport Type:Expedited (15-DaCompany Report #2004117310
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Consumer	Neurontin			
		Drug Ineffective		(Gabapentin)	PS		

Date:12/29/04ISR Number: 4542916-1Report Type:Expedited (15-DaCompany Report #2004117391
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Consumer	Neurontin			
Other		Gun Shot Wound		(Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:12/29/04ISR Number: 4543719-4Report Type:Direct
Age:41 YR Gender:Male I/FU:I

Company Report #CTU 235362

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria		Gabapentin 400 Mg Cap-Alpha	PS	Alpha	
800 MG BY							
MOUTH TID							

Date:12/29/04ISR Number: 4543923-5Report Type:Direct
Age:52 YR Gender:Female I/FU:I

Company Report #CTU 235262

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Required		Anger		Gabapentin 300 Mg			
Intervention to		Fall		Tablets Ivax	PS	Ivax	
900 MG QID							
Prevent Permanent		Hair Plucking		Seroquel	C		
Impairment/Damage		Mood Swings		Prevacid	C		
		Pharmaceutical Product		Zetia	C		
		Complaint		Plavix	C		
		Psychomotor Hyperactivity		Loratidine	C		
		Restless Legs Syndrome		Tramadol Hcl	C		
		Scratch		Bextra	C		
		Self Injurious Behaviour		Norvasc	C		
		Somnolence		Neurontin	C		

Date:12/30/04ISR Number: 4544532-4Report Type:Expedited (15-DaCompany Report #GXKR2004GB00603
Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Foreign	Gabapentin (Ngx)			
		Drug Exposure During	Health	(Gabapentin) Unknown	PS		ORAL
ORAL							
		Pregnancy	Professional	Propranolol (Ngx)			
			Other	(Propranolol)	SS		ORAL
ORAL							
				Detrusitol (Tolterodine L-Tartrate)Tablet,			

2 MG, QID,		2mg	SS	ORAL
ORAL				
BID, ORAL		Zopiclone (Zopiclone)	SS	ORAL
ORAL		Clonazepam (Clonazepam)	SS	ORAL
QID		Sertraline (Sertraline)	SS	

Date:12/30/04ISR Number: 4544790-6Report Type:Expedited (15-DaCompany Report #2004117398
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Depression Suicidal Ideation	Foreign Health	Pregabalin (Pregabalin)	PS		ORAL
ORAL			Professional	Neurontin (Gabapentin)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/03/05ISR Number: 4543862-XReport Type:Expedited (15-DaCompany Report #GXKR2004GB00603

Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Foreign Health	Gabapentin (Ngx) (Gabapentin)	PS		ORAL
ORAL							
		Pregnancy Haemorrhage	Professional Other	Propranolol (Ngx) (Propranolol)	SS		ORAL
ORAL							
				Detrusitol (Tolterodine L-Tartrate) Tablet, 2mg	SS		ORAL
2 MG,							
QID,ORAL							
				Zopiclone (Zopiclone)	SS		ORAL
BID,ORAL							
				Clonazepam (Clonazepam)	SS		ORAL
ORAL							
				Sertraline (Sertraline)	SS		
QID							

Date:01/03/05ISR Number: 4545194-2Report Type:Expedited (15-DaCompany Report #2004119096

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatic Failure	Foreign Health Professional	Gabapentin (Gabapentin)	PS		
				Carbamazepine	C		
				Antiepileptics	C		
				Amitriptyline	C		

Date:01/03/05ISR Number: 4545318-7Report Type:Expedited (15-DaCompany Report #2004096163

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Loss Of Consciousness	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL			Health Professional				

Date:01/03/05ISR Number: 4545485-5Report Type:Expedited (15-DaCompany Report #2004116627
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased	Health Professional	Neurontin (Gabapentin)	PS		
				All Other Therapeutic Products (All Other Therapeutic Products)	SS		

Date:01/03/05ISR Number: 4545827-0Report Type:Expedited (15-DaCompany Report #2004088109
Age:76 YR Gender:Female I/FU:F

Outcome	Duration	PT
Hospitalization - Initial or Prolonged		Blood Pressure Increased
Other		Dizziness Feeling Cold

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, 3 IN 1 D), ORAL		Hypoaesthesia Labyrinthitis Movement Disorder Pain In Extremity	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				All Other Therapeutic Products(All Other Therapeutic Products) Dyazide (Hydrochlorothiazide , Triamterene) Warfarin Sodium(Warfarin Sodium) Omeprazole (Omeprazole) Fentanyl (Fentanyl) Pantoprazole (Pantoprazole) Flurazepam Hydrochloride(Flurazepam Hydrochloride_	C C C C C C		

Date:01/03/05ISR Number: 4545834-8Report Type:Expedited (15-DaCompany Report #2004077846

Age:43 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other SEE IMAGE		Blood Pressure Increased Blood Sodium Decreased	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Convulsion Drug Ineffective Treatment Noncompliance	Professional	Bextra(Valdecoxib) Cyclobenzaprine (Cyclobenzaprine) Levetiracetam (Levetiracetam) Alprazolam	SS C C		

(Alprazolam) C
Dextropropoxyphene
(Dextropropoxyphene) C

Date:01/03/05ISR Number: 4545869-5Report Type:Expedited (15-DaCompany Report #2004119079
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other	Renal Failure Acute	Health Professional	Neurontin (Gabapentin)	PS		

Date:01/03/05ISR Number: 4545899-3Report Type:Expedited (15-DaCompany Report #2004076841
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other	Economic Problem Treatment Noncompliance	Consumer	Neurontin (Gabapentin)	PS		

1800 MG (600

MG,3 IN 1 D)

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/03/05ISR Number: 4545924-XReport Type:Expedited (15-DaCompany Report #2004117371
 Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Erythema Fibromyalgia	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Herpes Zoster Hot Flush Hyperhidrosis Medication Error Nausea Sleep Disorder Due To General Medical Condition, Insomnia Type Spinal Fusion Surgery Treatment Noncompliance		Oxycodone Hydrochloride (Oxycodone Hydrochloride) Oxycodone (Oxycodone) Amitriptyline (Amitriptyline)	C C C		

Date:01/03/05ISR Number: 4545928-7Report Type:Expedited (15-DaCompany Report #2004113565
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anticonvulsant Toxicity Fatigue	Consumer	Neurontin (Gabapentin)	PS		
1200 MG (400 MG , 3 IN 1 D)		Gait Disturbance Grand Mal Convulsion Pain Pain In Extremity Sluggishness Therapy Non-Responder Vision Blurred		Dilantin Suspension (Phenytoin Sodium) Carbamazepine (Carbamazepine) Rantidine Hydrochloride (Ranitidine Hydrochloride) Calcium (Calcium)	SS SS C C		

Date:01/03/05ISR Number: 4545935-4Report Type:Expedited (15-DaCompany Report #2004119995
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Surgery	Health Professional	Neurontin (Gabapentin)	PS		

Date:01/03/05ISR Number: 4545955-XReport Type:Expedited (15-DaCompany Report #2004010780
Age:69 YR Gender:Male I/FU:F

Outcome	PT
Disability	Actinic Keratosis
Other	Anhedonia
	Arthralgia
	Blood Urine Present
	Bundle Branch Block Left
	Difficulty In Walking
	Drug Hypersensitivity
	Emotional Disorder
	Fatigue
	Feeling Cold
	Flushing
	Hyperhidrosis
	Impaired Work Ability

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
10 MG (DAILY)		Insomnia Kyphosis Low Density Lipoprotein	Consumer	Lipitor (Atorvastatin)	PS		
300 MG, ORAL		Increased Mental Disorder	Consumer	Neurontin (Gabapentin)	SS		ORAL
(DAILY), ORAL		Muscle Atrophy Muscle Spasms	Consumer	Hyzaar (Hydrochlorothiazide , Losartan Potassium)	SS		ORAL
		Neck Injury Nervousness Neuropathy Pain	Consumer	Influenza Vaccine (Influenza Vaccine)	C		
		Pain In Extremity Ph Urine Decreased Polymyalgia Polymyositis Prostatic Specific Antigen Increased Protein Urine Present Spinal Column Stenosis Tenderness Visual Acuity Reduced Weight Decreased	Consumer	All Other Therapeutic Products (All Other Therapeutic Products)	C		
			Consumer	Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		
			Consumer	Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
			Consumer	Amlodipine Besilate (Amlodipine Besilate)	C		

Date:01/03/05ISR Number: 4545962-7Report Type:Expedited (15-DaCompany Report #2004048907

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anxiety Chest Pain	Consumer	Neurontin (Gabapentin)	PS		
200 MG (400		Condition Aggravated	Consumer	Cefalexin (Cefalexin)	C		
MG, 5 IN 1 D)		Cough Deformity Diaphragmatic Injury	Consumer	Warfarin Sodium	C		

Economic Problem	(Warfarin Sodium)	C
Gun Shot Wound	Phenylpropanolamine	
Hepatic Trauma	(Phenylpropanolamine	
Hyperventilation)	C
Mental Disorder	Cyproheptadine	
Pain	Hydrochloride	C
Pericardial Effusion	Yohimbine	
Pericardial Excision	(Yohimbine)	C
Pleural Effusion	Cetirizine	
Pneumonia	Hydrochloride	
Polytraumatism	(Cetirizine	
Suicide Attempt	Hydrochloride)	C
	Adapalene	
	(Adapalene)	C
	Meloxicam	
	(Meloxicam)	C
	Aripiprazole	
	(Aripiprazole)	C
	Ketorolac	
	Tromethamine	
	(Ketorolac	
	Tromethamine)	C

Freedom Of Information (FOI) Report

Iscom	
(Dichloralphenazone,	
Isometheptene	
Mucate, Paracetamol)	C
Clonazepam	
(Clonazepam)	C
Olanzapine	
(Olanzapine)	C
Zolpidem Tartrate	
(Zolpidem	
Tartrate)	C
Fluoxetine	
Hydrochloride	
(Fluoxetine	
Hydrochloride)	C
Raloxifene	
Hydrochloride	
(Raloxifene	
Hydrochloride)	C
Lithium Carbonate	
(Lithium Carbonate)	C
Tramadol	
Hydrochloride	
(Tramadol	
Hydrochloride)	C
Atenolol (Atenolol)	C
Trazodone	
9trazodone0	C
Propacet	
(Dextropropoxyphene	
Napsilate,	
Paracetamol)	C
Estratest H	
(Estrogens	
Esterified,	
Methyltestosterone0	C
Estradiol	
(Estradiol)	C
Venlafaxine	
Hydrochloride(Venlaf	
axine Hydrochloride)	C
Nefazodone	
Hydrochloride(Nefazo	
done Hydrochloride)	C
Zolmitriptan	
(Zolmitriptan)	C
Sodium Fluoride	

(Sodium Fluoride)	C
Amoxicillin	
(Amoxicillin)	C
Meperidine	
W/Promethazine	
(Pethidine,	
Promethazine)	C
Azithromycin	
(Azithromycin)	C
Aquatab C	
(Dextromethorphan,	
Guaifenesin,	
Phenylpropanolamine)	C
Prednisone	

Freedom Of Information (FOI) Report

(Prednisone)	C
Pravastain Sodium	
(Pravastatin Sodium)	C
Clindamycin	
(Clindamycin)	C
Chlorhexidine	
(Chlorhexidine)	C
Vicodin (Hydrocodone	
Bitartrate,	
Paracetamol)	C
Docusate Sodium	
(Docusate Sodium)	C
Risperidone	
(Risperidone)	C
Mirtazapine	
(Mirtazapine)	C
Bupropion	
Hydrochloride	
(Bupropion	
Hydrochloride)	C
Propranolol	
Hydrochloride	
(Propranolol	
Hydrochloride)	C
Sumatriptan	
Succinate	
(Sumatriptan	
Succinate)	C
Methocarbamol	
(Methocarbamol)	C
Naproxen (Naproxen)	C
.....	C
Narine Repetabs	
(Loratadine,	
Pseudoephedrine	
Sulfate)	C
Temezapam	
(Temazepam)	C
Quetiapine Fumarate	
(Quetiapine	
Fumarate0	C
Robutisin A-C /Old	
Form/ (Codeine	
Phosphate ,	
Guaifenesin,	
Pheniramine Maleate0	C
Flurazepam	
Hydrochloride	

(Flurazepam Hydrochloride)	C
Mometasone Furoate (Mometasone Furoate)	C
Adapalene (Adapalene)	C
Ronatic (Chlorphenamine Tannate, Mepyramine Tannate, Phenylephrine	C
Modafinil (Modafinil)	C
Tizanidine	

Freedom Of Information (FOI) Report

Hydrochloride(
 Tizanidine
 Hydrochloride) C
 Estrogens Conjugated
 (Estrogens
 Conjugated) C
 Estradiol
 (Estradiol) C
 Tiababine
 Hydrochloride
 9tiagabine
 Hydrochloride) C
 Sildenafil Citrate
 (Sildenafil
 Citrate) C
 Butorphanol Tartrate

 (Butorphanol
 Tartrate) C
 Trimethobenzamide
 Hydrochloride
 (Trimethobenzamide
 Hydrochloride) C
 Lithium Carbonate
 (Lithium Carbonate0 C
 Valproate Semisodium
 (Valproate
 Semisodium) C
 Beclometasone
 Dipropionate
 (Beclometasone
 Dipropionate) C
 Clavulin
 (Amoxicillin
 Trihydrate,
 Clavulanate
 Potassium) C

Date:01/03/05ISR Number: 4546041-5Report Type:Expedited (15-DaCompany Report #2004119074

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Brain Operation Therapy Non-Responder	Consumer	Neurontin (Gabapentin)	PS		

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Abdominal Pain Lower
Disability	Abdominal Pain Upper
Other	Abnormal Behaviour
	Acute Sinusitis
	Affect Lability
	Agitation
	Anger
	Angioneurotic Oedema
	Animal Bite
	Ankle Fracture

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Freedom Of Information (FOI) Report

Anorexia
Antinuclear Antibody
Positive
Anxiety
Arthralgia
Arthritis
Arthropod Bite
Asthenia
Asthma
Astigmatism
Atrial Flutter
Atrial Tachycardia
Atrioventricular Block
Back Pain
Bacteraemia
Balance Disorder
Blepharitis
Blepharospasm
Blood Albumin Decreased
Blood Glucose Increased
Blood Urea Decreased
Bronchitis
Bruxism
Burn Infection
Carpal Tunnel Syndrome
Cellulitis
Cervicitis
Change Of Bowel Habit
Chest Pain
Chromaturia
Cognitive Disorder
Condition Aggravated
Constipation
Convulsion
Coordination Abnormal
Crying
Deafness
Decreased Appetite
Depression
Diarrhoea
Diplopia
Dissociation
Disturbance In Attention
Dizziness
Drug Hypersensitivity
Drug Withdrawal Syndrome
Dry Mouth

Dysarthria
Dysgeusia
Dyskinesia
Dyspepsia
Dysphagia
Dysphasia
Dyspnoea
Dysuria
Ear Pain
Ecchymosis
Essential Tremor
Eustachian Tube
Dysfunction
Eyelid Ptosis

Freedom Of Information (FOI) Report

Facial Bones Fracture
Fall
Fibrocystic Breast
Disease
Flat Affect
Fungal Rash
Gastrointestinal Disorder
Gastrooesophageal Reflux
Disease
Gingival Recession
Goitre
Granuloma
Hallucination, Visual
Hand Fracture
Head Injury
Heart Rate Increased
Heart Rate Irregular
Hordeolum
Hot Flush
Hypersomnia
Hypoglycaemia
Hypothyroidism
Impaired Healing
Impaired Self-Care
Imprisonment
Impulsive Behaviour
Increased Tendency To
Bruise
Increased Viscosity Of
Bronchial Secretion
Insomnia
Intervertebral Disc
Degeneration
Intervertebral Disc
Disorder
Irritability
Irritable Bowel Syndrome
Joint Injury
Joint Sprain
Joint Swelling
Keratoconjunctivitis
Sicca
Laceration
Laryngitis
Limb Discomfort
Limb Injury
Loss Of Consciousness

Lumbar Radiculopathy
Macroglossia
Malnutrition
Mania
Mean Cell Haemoglobin
Concentration Increased
Medical Device
Complication
Medication Error
Memory Impairment
Meniere'S Disease
Menorrhagia
Mouth Breathing
Multiple Fractures

Freedom Of Information (FOI) Report

Muscle Spasms
Myopia
Nasal Discomfort
Nasal Ulcer
Nausea
Neck Injury
Neck Pain
Nervous System Disorder
Neuropathy
Nystagmus
Oedema Peripheral
Oesophageal Disorder
Onychomadesis
Otitis Media
Ovarian Cyst
Ovarian Cyst Ruptured
Ovarian Infection
Pain In Extremity
Pelvic Inflammatory
Disease
Pharyngitis
Photophobia
Photopsia
Pitting Oedema
Post Procedural
Complication
Premenstrual Syndrome
Propionibacterium
Infection
Pruritus
Psoriasis
Psychiatric Symptom
Pyrexia
Radiculitis
Rash
Rash Erythematous
Red Blood Cell Count
Decreased
Restless Legs Syndrome
Rhinitis Allergic
Road Traffic Accident
Sensory Disturbance
Sialoadenitis
Skin Ulcer
Sneezing
Somatisation Disorder
Somnolence

Speech Disorder
Staring
Stress
Suicidal Ideation
Swollen Tongue
Syncope
Tardive Dyskinesia
Tendonitis
Thermal Burn
Thirst
Thrombin Time Shortened
Tinnitus
Tongue Blistering
Tongue Disorder

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1800 MG	(DAILY), ORAL	Traumatic Brain Injury				
		Tremor				
		Tubo-Ovarian Abscess				
		Unresponsive To Verbal Stimuli	Neurontin (Gabapentin)	PS		ORAL
		Upper Respiratory Tract Infection	Lamotrigine (Lamotrigine)	C		
		Urinary Tract Infection	Lithium Carbonate (Lithium Carbonate)	C		
		Vaginal Candidiasis	Clonazepam (Clonazepam)	C		
		Vertigo Positional	Methylphenidate Hydrochloride (Methylphenidate Hydrochloride)	C		
		Vestibular Neuronitis	Levothyroxine Sodium (Levothyroxine Sodium)	C		
		Vestibulitis	Liothyronine Sodium (Liothyronine Sodium)	C		
		Viral Infection	Sertraline Hydrochloride (Sertraline Hydrochloride)	C		
		Viral Labyrinthitis	Pilocarpine Hydrochloride (Pilocarpine Hydrochloride)	C		
		Viral Sinusitis	Metoprolol Succinate (Metoprolol Succinate)	C		
		Vision Blurred	Lansoprazole (Lansoprazole)	C		
		Vomiting	Hyoscyamine Sulfate (Hyoscyamine Sulfate)	C		
		Weight Decreased	Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C		
		Weight Increased	Bupropion Hydrochloride			
		White Blood Cell Count Decreased				
		Wound Infection				
		Wrist Fracture				

(Bupropion Hydrochloride)	C
Celebrex (Celecoxib)	C
Quetiapine Fumarate (Quetiapine Fumarate)	C
Metoprolol Tartrate (Metoprolol Tartrate)	C
Citalopram Hydrobromide (Citalopram Hydrobromide)	C
Olanzapine (Olanzapine)	C
Omeprazole (Omeprazole)	C

Freedom Of Information (FOI) Report

Nasal Preparations
 (Nasal Preparations) C
 Nefazodone
 Hydrochloride
 (Nefazodone
 Hydrochloride) C
 Totolin
 (Guaifenesin,
 Phenylpropanolamine
 Hydrochloride) C
 Meclozine
 (Meclozine) C
 Fluticasone
 Propionate
 (Fluticasone
 Propionate) C
 Trimethobenzamide
 Hydrochloride
 (Trimethobenzamide
 Hydrochloride) C
 Respaire-Sr-120
 (Guaifenesin,
 Pseudoephedrine
 Hydrochloride) C
 Risperidone
 (Risperidone) C

Date:01/03/05ISR Number: 4546332-8Report Type:Expedited (15-DaCompany Report #2004117941
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arthritis	Consumer	Celebrex (Celecoxib)	PS		ORAL
Other		Bursitis		Neurontin			
ORAL		Condition Aggravated		(Gabapentin)	SS		ORAL
		Contusion		Sertraline			
		Coordination Abnormal		Hydrochloride	C		
		Drug Effect Decreased		Amlodipine Besilate	C		
		Excoriation		Paracetamol	C		
		Fall		Dyazide			
		Muscle Spasms		(Hydrochlorothiazide			
		Neuralgia		, Triamterene)	C		
		Post Procedural		Lotrel (Amlodipine,			

Complication
Spinal Fusion Surgery

Benazepril
Hydrochloride) C

Date:01/03/05ISR Number: 4546563-7Report Type:Expedited (15-DaCompany Report #CIP04002758
Age:81 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Cutaneous Vasculitis Inflammation Leukocytoclastic Vasculitis	Foreign Health Professional Other	Furadantine (Nitrofurantoin Macrocrystals) Capsule	PS		ORAL
1 CAPSUL		Pain Of Skin					
DAILY, ORAL		Staphylococcal Infection Urinary Tract Infection Vascular Purpura		Previscan (Fluindione) Tablet, 20 Mg	SS		ORAL
1 TABLET		Vasculitis Necrotising					
DAILY, ORAL				Zocor "Dieckmann"			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL			(Simvastatin)	SS	Dieckmann	ORAL
ORAL			Xatral/Fra/(Alfuzosin)	SS		ORAL
ORAL			Neurontin (Gabapentin)	SS		ORAL
			Renitec(Enalapril Maleate)	C		
			Vaccines	C		

Date:01/03/05ISR Number: 4548044-3Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 235446

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Increased		Metalozone 5 Mg	PS		ORAL
5 MG ONCE /PO		Drug Effect Decreased Pharmaceutical Product		Neurontin 400 Mg / Gabapentin	SS		
400 MG THREE TIMES		Complaint		Gabapentin	SS		

Date:01/04/05ISR Number: 4545715-XReport Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 235568

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Tremor		Gabapentin 600mg / 300mg	PS		ORAL
600MG PO BID;				Nefazadone	C		
300MG PO QHS				Tramazone	C		

Date:01/04/05ISR Number: 4545874-9Report Type:Direct
Age:55 YR Gender:Male I/FU:I

Company Report #CTU 235609

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation Dizziness		Gabapentin (Neurontin)	PS	(Pt Only Reactic To Generic	ORAL
300 MG PO TID							

Date:01/05/05ISR Number: 4546662-XReport Type:Expedited (15-DaCompany Report #2004AL000891
Age:58 YR Gender:Female I/FU:F

Outcome	PT
Death	Acidosis
Hospitalization - Initial or Prolonged	Acute Respiratory Distress Syndrome
	Alanine Aminotransferase Increased
	Arrhythmia
	Aspartate Aminotransferase Increased
	Blood Calcium Decreased
	Cardiac Arrest
	Coagulopathy
	Coma
	Compartment Syndrome
	Completed Suicide
	Haemodialysis
	Hypotension

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Intentional Misuse Labile Blood Pressure Procedural Complication Renal Failure	Report Source	Product	Role	Manufacturer	Route
PO			Literature Health Professional	Children'S Apap Elixir (Acetaminophen) (Alpharma)	PS	Alpharma	ORAL
PO				Infants' Apap Drops (Acetaminophen) (Alpharma)	SS	Alpharma	ORAL
PO				Olanzapine	SS		ORAL
PO				Buspirone	SS		ORAL
PO				Gabapentin Capsules, 100 Mg, 300 Mg & 400 Mg (Purepac)	SS	Purepac	ORAL
PO				Gabapentin Tablets, 600 Mg & 800 Mg (Purepac)	SS	Purepac	ORAL
PO				Paroxetine	SS		ORAL
PO				Levothyroxine	SS		ORAL
PO				Estrogens	SS		ORAL

Date:01/05/05ISR Number: 4548181-3Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 235746

Outcome Dose Other	Duration	PT Bladder Spasm Micturition Disorder	Report Source	Product	Role	Manufacturer	Route
1800 MG PO DAILY				Neurontin 300mg Purepac	PS	Purepac	ORAL
350 MG PO HS				Carisoprodol 350 Mg Watson	SS	Watson	ORAL

Outcome	PT
Death	Agitation
Hospitalization -	Alcohol Poisoning
Initial or Prolonged	Anger
Disability	Anxiety
Other	Arthralgia
	Aspiration
	Asthenia
	Brain Oedema
	Burning Sensation
	Cardio-Respiratory Arrest
	Cardiomegaly
	Coma
	Coronary Artery
	Atherosclerosis
	Drug Abuser
	Drug Ineffective
	Drug Interaction
	Drug Screen Positive
	Erectile Dysfunction
	Excoriation
	Fall
	Feeling Abnormal
	Haematemesis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Hepatic Steatosis Hypertensive Heart Disease				
600 MG (300 MG, 2 IN 1 D), ORAL		Hypoaesthesia Impaired Driving Ability	Neurontin (Gabapentin)	PS		ORAL
		Insomnia				
		Intentional Self-Injury				
ORAL		Intervertebral Disc Injury	Benadryl (Diphenhydramine)	SS		ORAL
1-3 TABLETS Q12H, ORAL		Intervertebral Disc Operation Intervertebral Disc Protrusion	Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL
		Laceration				
1-2 EVERY 4-6 HOURS (10 MG), ORAL		Lumbar Radiculopathy Major Depression	Hydrocodone (Hydrocodone)	SS		ORAL
		Mood Swings				
		Muscle Spasms				
10 MG (10 MG, 1 IN 1 D), ORAL		Muscle Spasticity Musculoskeletal Pain Pain Pain In Extremity	Zolpidem Tartrate (Zolpidem Tartrate)	SS		ORAL
		Paraesthesia Post Procedural Complication Post Procedural Pain	Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		ORAL
80 MG (80 MG, 1 IN 1 D), ORAL		Pseudarthrosis Pulmonary Oedema				
		Radiculitis Radiculopathy Renal Cyst	Amitriptyline Hydrochloride (Amitriptyline			

Snoring
Spinal Myelogram Abnormal
Suicide Attempt
Tearfulness
Ventricular Hypertrophy
Vomiting

Hydrochloride) C
Fentanyl (Fentanyl) C
Cyclobenzaprine
Hydrochloride
(Cyclobenzaprine
Hydrochloride) C
Bupropion
Hydrochloride
(Bupropion
Hydrochloride) C
Quinine Sulfate
(Quinine Sulfate) C
Sertraline
Hydrochloride
(Sertraline
Hydrochloride) C
Trazodone
(Trazodone) C
Hydroxyzine Embonate
(Hydroxyzine
Embonate) C
Carisoprodol
(Carisoprodol) C
Paroxetine
Hydrochloride
(Paroxetine
Hydrochloride) C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/07/05ISR Number: 4547651-1Report Type:Expedited (15-DaCompany Report #US-ABBOTT-05P-163-0285080-00
Age:27 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Imminent Abortion		Depakote 500mg	PS		
Initial or Prolonged	Threatened Labour		Gabapentin	SS		
	Vomiting		Levetiracetam	SS		
			Levetiracetam	SS		
			Zonisamide	SS		

Date:01/07/05ISR Number: 4548787-1Report Type:Direct Company Report #CTU 235907
Age:69 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Grimacing		Gabapentin 600 Mg	PS		ORAL
300 MG AT						
Initial or Prolonged	Movement Disorder					
NIGHT ORAL						
	Muscle Twitching					
	Tardive Dyskinesia					

Date:01/07/05ISR Number: 4551094-4Report Type:Expedited (15-DaCompany Report #2004121249
Age:41 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Other	Aggression	Consumer	Neurontin (Tablets)			
	Depression		(Gabapentin)	PS		ORAL
4 IN 1 D,						
ORAL	Drug Ineffective					
	Drug Withdrawal Syndrome		Acetylsalicylic Acid			
	Insomnia		(Acetylsalicylic			
	Restlessness		Acid)	C		
	Treatment Noncompliance		All Other			
			Therapeutic Products			
			(All Other			
			Therapeutic			
			Products)	C		
			All Other			
			Non-Therapeutic			

Date:01/07/05ISR Number: 4551098-1Report Type:Expedited (15-DaCompany Report #2004117371
 Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation	Consumer	Neurontin			
ORAL		Economic Problem		(Gabapentin)	PS		ORAL
		Erythema		Oxycodone			
		Feeling Hot		Hydrochloride			
		Fibromyalgia		(Oxycodone			
		Herpes Zoster		Hydrochloride)	C		
		Hyperhidrosis		Oxycodone			
		Insomnia		(Oxycodone)	C		
		Nausea		Amitriptyline			
		Pain		(Amitriptyline)	C		
		Surgery					
		Treatment Noncompliance					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/07/05ISR Number: 4551221-9Report Type:Expedited (15-DaCompany Report #2004067531
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Death	Consumer	Neurontin			
Hospitalization -		Anxiety	Health	(Gabapentin)	PS		ORAL
600 MG (300		Arthralgia	Professional				
Initial or Prolonged		Aspiration					
MG, 2 IN 1		Atherosclerosis		Benadryl			
Disability		Brain Oedema		(Diphenhydramine)	SS		ORAL
D), ORAL		Burning Sensation		Oxycodone			
Other		Cardio-Respiratory Arrest		Hydrochloride			
ORAL		Cardiomegaly		(Oxycodone			
		Coma		Hydrochloride)	SS		ORAL
1-3 TABLETS		Condition Aggravated					
Q12H, ORAL		Coronary Artery		Hydrocodone			
		Atherosclerosis		(Hydrocodone)	SS		
1-2 EVERY 4-6		Drug Interaction					
HOURS (10		Drug Screen Positive					
MG), ORAL		Excoriation		Zolpidem Tartrate			
		Faecal Occult Blood		(Zolpidem Tartrate)	SS		ORAL
10 MG (10 MG,		Positive					
1 IN 1 D) ,		Feeling Abnormal					
ORAL		Haematemesis		Citalopram			
		Hepatic Steatosis		Hydrobromide			
		Hypertensive Heart		(Citalopram			
80 MG (80 MG,		Disease		Hydrobromide)	SS		ORAL
		Hypoaesthesia					
1 IN 1 D),		Injury					
ORAL		Insomnia		Amitriptyline			
		Intentional Self-Injury		Hydrochloride			

Laceration	(Amitriptyline	
Major Depression	Hydrochloride)	C
Multiple Drug Overdose	Fentanyl (Fentanyl)	C
Accidental	Cyclobenzaprine	
Muscle Spasms	Hydrochloride	
Muscle Spasticity	(Cyclobenzaprine	
Neck Pain	Hydrochloride)	C
Pain	Bupropion	
Pain In Extremity	Hydrochloride	
Post Procedural	(Bupropion	
Complication	Hydrochloride)	C
Radiculopathy	Quinine Sulfate	
Renal Cyst	(Quinine Sulfate)	C
Respiratory Depression	Sertraline	
Vomiting	Hydrochloride	
	(Sertraline	
	Hydrochloride)	C
	Trazodone	
	(Trazodone)	C
	Hydroxyzine Embonate	
	(Hydroxyzine	
	Embonate)	C
	Carisoprodol	
	(Carisoprodol)	C
	Paroxetine	
	Hydrochloride	
	(Paroxetine	
	Hydrochloride)	C

Freedom Of Information (FOI) Report

Date:01/07/05ISR Number: 4562092-9Report Type:Expedited (15-DaCompany Report #2003018755
Age:37 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abasia
Initial or Prolonged	Abdominal Distension
Disability	Abdominal Pain
Other	Agitation
	Anxiety
	Apathy
	Arthralgia
	Blood Calcium Decreased
	Blood Creatinine
	Decreased
	Blood Electrolytes
	Decreased
	Blood Glucose Decreased
	Blood Lactate
	Dehydrogenase Decreased
	Blood Potassium Decreased
	Blood Pressure Increased
	Blood Triglycerides
	Increased
	Blood Urea Decreased
	Brain Damage
	Bursitis
	Cerebral Artery Occlusion
	Cerebrovascular Accident
	Cerebrovascular Disorder
	Constipation
	Convulsion
	Decreased Appetite
	Depression
	Diarrhoea
	Difficulty In Walking
	Dizziness
	Dysmenorrhoea
	Electrocardiogram St-T
	Change
	Electroencephalogram
	Abnormal
	Emotional Distress
	Eye Pain
	Facial Pain
	Facial Palsy
	Faecal Occult Blood
	Positive

Flatulence
Gliosis
Grand Mal Convulsion
Haemorrhoids
Hallucination, Auditory
Hemiparesis
Herpes Simplex
Insomnia
Intracranial Aneurysm
Irritable Bowel Syndrome
Joint Dislocation
Loss Of Consciousness
Mean Cell Haemoglobin
Increased
Mean Cell Volume Abnormal

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2400 MG (600, FOUR TIMES A DAY), ORAL		Menorrhagia	Neurontin (Tablets) (Gabapentin)	PS		ORAL
		Menstrual Disorder				
		Migraine				
20 MG (Q12H), ORAL		Mitral Valve Incompetence	Loestrin (Anovlar) (Norethindrone Acetate, Ethinyl Estradiol)	SS		
		Movement Disorder				
		Muscle Atrophy				
		Muscle Contracture				
		Muscle Rigidity				
		Muscle Spasms				
		Nervous System Disorder				
		Oral Pain				
		Osteopenia				
		Pain				
		Pco2 Decreased				
		Peroneal Nerve Palsy				
		Photopsia				
		Psychomotor Retardation				
		Psychotic Disorder				
Red Blood Cell Count Decreased						
Road Traffic Accident						
Schizophrenia, Disorganised Type						
Somatisation Disorder						
Suicidal Ideation						
Urinary Incontinence						
Vitamin B12 Increased						
Weight Increased						

Date:01/07/05ISR Number: 4625252-4Report Type:Periodic
Age:25 YR Gender:Female I/FU:I

Company Report #235144K04USA

Outcome	Duration	Report Source	Product	Role	Manufacturer	Route	
SUBCUTANEOUS	44 MCG, 3 IN	Consumer	Rebif (Interferon Beta)	PS			
							PT
							Genital Discharge
							Genital Pruritus Female
1 WEEKS,							

SUBCUTANEOUS Vaginal Discharge

Gabapentin SS

Date:01/07/05ISR Number: 4628915-XReport Type:Periodic Company Report #230219M04USA
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Circulatory Collapse	Health	Rebif (Interferon			
		Fall	Professional	Beta)	PS		
		Syncope		Neurontin			
				(Gabapentin)	SS		

Date:01/10/05ISR Number: 4548454-4Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0381214A
Age:42 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Agranulocytosis
Initial or Prolonged	Bipolar Disorder
	Bradycardia
	Hypotension
	Thrombocytopenia
	White Blood Cell Count

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Decreased

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1	YR		Lamictal	PS	Glaxosmithkline	ORAL
			Trileptal	SS		
			Anti-Psychotic Medication	SS		
2MG	Four times per day		Gabapril	SS		ORAL
			Neurontin	SS		
			Lithium	C	Glaxosmithkline	
			Claritin	C		
			Sonata	C		
			Colace	C		
			Multiple Vitamin	C		

Date:01/10/05ISR Number: 4548455-6Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0386384A
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agranulocytosis		Lamictal	PS	Glaxosmithkline	ORAL
		Bradycardia		Gabapril	SS		ORAL
2MG	Four times per day	Hypotension					
		Thrombocytopenia		Neurontin	SS		ORAL
				Lithium	C	Glaxosmithkline	
				Sonata	C		
				Claritin	C		
				Multivitamins	C		
				Colace	C		

Date:01/10/05ISR Number: 4549708-8Report Type:Expedited (15-DaCompany Report #2004066759
 Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Myocardial Infarction	Consumer	Lipitor			

80 MG (80 MG, 1 IN 1 D)	Health Professional	(Atorvastatin)	PS
900 MG (300 MG, 3 IN 1 D)		Neurontin (Gabapentin)	SS
20 MG (10 MG, 2 IN 1 D)		Glipizide (Glipizide)	SS
25 MG (25 MG, 1 IN 1 D)		Atenolol (Atenolol)	SS
60 MG (30 MG, 2 IN 1 D)		Lansoprazole (Lansoprazole)	SS
2.5 MG (2.5 MG, 1 IN 1 D)		Ramipril (Ramipril)	SS
		Clopidogrel Sulfate (Clopidogrel Sulfate)	C
		Glyceryl Trinitrate (Glyceryl Trinitrate)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/10/05ISR Number: 4549713-1Report Type:Expedited (15-DaCompany Report #2004058756

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Phantom Pain	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
3200 MG (800 MG, 4 IN 1 D), ORAL		Pharmaceutical Product Complaint					

Date:01/10/05ISR Number: 4549996-8Report Type:Expedited (15-DaCompany Report #2004081555

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety Somnolence	Consumer	Neurontin (Gabapentin)	PS		ORAL
SEE IMAGE							

Date:01/10/05ISR Number: 4550086-9Report Type:Expedited (15-DaCompany Report #2004074283

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Balance Disorder Deafness	Consumer Health	Neurontin (Gabapentin)	PS		
Other		Pain Tinnitus	Professional				

Date:01/10/05ISR Number: 4550090-0Report Type:Expedited (15-DaCompany Report #2004063257

Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Activities Of Daily Living Impaired	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
300 MG (100							

Aggression

Professional

D), ORAL

Agitation

Depression

Diabetes Mellitus

Inadequate Control

Gastric Disorder

Heart Rate Increased

Hypertension

Myocardial Infarction

Physical Assault

Celebrex (Celecoxib) SS

Amoxicillin

W/Clavulanate

Potassium

(Amoxicillin,

Clavulanate

C

Metoprolol

(Metoprolol)

C

Metformin

(Metformin)

C

Hydrochlorothiazide

(Hydrochlorothiazide

)

C

Repaglinide

(Repaglinide)

C

Moexipril

Hydrochloride

(Moexipril

Hydrochloride)

C

Glyceryl Trinitrate

(Glyceryl

Trinitrate)

C

Desloratadine

(Desloratadine)

C

Clavulin

(Amoxicillin

Freedom Of Information (FOI) Report

Trihydrate,
 Clavulanate
 Potassium) C
 Clopidogrel Sulfate
 (Clopidogrel
 Sulfate) C
 Lotrel (Amlodipine,
 Benazepril
 Hydrochloride) C
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C
 Lovastatin
 (Lovastatin) C

Date:01/10/05ISR Number: 4550106-1Report Type:Expedited (15-DaCompany Report #2004121824

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Limb Immobilisation Limb Injury	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 2 IN 1 D), ORAL				Oxybutynin (Oxybutynin)	C		

Date:01/10/05ISR Number: 4550390-4Report Type:Expedited (15-DaCompany Report #ZONI002085

Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG, 1 IN Required 1 D, ORAL Intervention to		Abortion Spontaneous Drug Exposure During Pregnancy Vomiting	Health Professional	Zonegran (Zonisamide)	PS		ORAL
				Depakote (Valproate			

Prevent Permanent
500 MG, ORAL
Impairment/Damage

1200 MG, ORAL

SEE IMAGE,

ORAL

Semisodium)	SS	ORAL
Neurontin (Gabapentin)	SS	ORAL
Keppra (Levetiracetam)	SS	ORAL

Date:01/10/05ISR Number: 4551283-9Report Type:Expedited (15-DaCompany Report #2004121923

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	600-900 MG (1	Condition Aggravated Drug Ineffective	Foreign Consumer	Neurontin (Gabapentin)	PS		
		Mood Swings					
		Retinal Disorder					
		Somnolence Visual Disturbance		All Other Therapeutic Products (All Other Therapeutic Products)	C		
				Karvea Hct (Hydrochlorothiazide , Irbesartan)	C		

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Freedom Of Information (FOI) Report

Salbutamol Sulfate
 (Salbutamol Sulfate) C
 Tiotropium Bromide
 (Tiotropium Bromide) C

Date:01/10/05ISR Number: 4551501-7Report Type:Expedited (15-DaCompany Report #2005000621
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Epilepsy	Foreign	Neurontin			
		Hepatic Function Abnormal	Health	(Gabapentin)	PS		
1200 MG (1 D)		Hyperammonaemia	Professional	Omeprazole			
		Liver Disorder		(Omeprazole)	C		

Date:01/10/05ISR Number: 4551814-9Report Type:Expedited (15-DaCompany Report #2004111564
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Cerebral Haemorrhage	Foreign	Neurontin			
Hospitalization -		Dizziness	Health	(Gabapentin)	PS		ORAL
800 MG (400		Headache	Professional				
Initial or Prolonged		Hypoaesthesia					
MG, 2 IN 1		Hyponatraemia		Diuretics			
D), ORAL		Treatment Noncompliance		(Diuretics)	SS		
		Vertigo		Ramipril (Ramipril)	C		
		Vomiting		Atorvastatin			
				(Atorvastatin)	C		
				Acetylsalicylic Acid			
				(Acetylsalicylic			
				Acid)	C		
				Metformin			
				Hydrochloride			
				(Metformin			
				Hydrochloride)	C		
				Bisoprolol Fumarate			
				(Bisoprolol			

Fumarate)

C

Date:01/10/05ISR Number: 4551914-3Report Type:Expedited (15-DaCompany Report #2004026162
Age:66 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Hepatitis Acute	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D)			Professional Company Representative Other	Aporex (Dextropoxyphene Hydrochloride, Paracematol) Hydroxyzine Hydrochloride (Hydroxylzine Hydrochloride) Thiocolchicoside (Thiocolchicoside)			
					C		
					C		
					C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/11/05ISR Number: 4549628-9Report Type:Expedited (15-DaCompany Report #US012745
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Diplopia	Health	Lamictal	PS	Glaxosmithkline	ORAL
Other		Dizziness	Professional	Gabitril	SS		ORAL
4MG Twice per day	15 DAY	Drug Interaction					
UNKNOWN		Dystonia		Celexa	SS		
300MG		Hallucination		Neurontin	SS		ORAL
Variable dose		Hypoglycaemia					
.5MG As required		Oedema Peripheral		Xanax	SS		ORAL
.5MG Twice per day		Vision Blurred					
		Visual Acuity Reduced		Klonopin	SS		ORAL
				Alcohol	C		
				Cocaine	C		

Date:01/11/05ISR Number: 4550014-6Report Type:Direct Company Report #CTU 236191
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ONE PO TID		Drug Ineffective		Neurontin 300 Mg	PS		ORAL
		Pharmaceutical Product Complaint		Gabapentin	C		

Date:01/11/05ISR Number: 4550018-3Report Type:Direct Company Report #CTU 236165
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Pain In Extremity		Gabapentin (Generic)			

400 MG 4 X

400 Qid

PS

ORAL

DAILY ; ORAL

Date:01/11/05ISR Number: 4550273-XReport Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 236134

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product Complaint		Gabapentin Er 200 Mg Bid	PS		ORAL
ORAL 200 MG		Rash Erythematous					
BID		Rash Pruritic					

Date:01/11/05ISR Number: 4550369-2Report Type:Direct
Age:32 YR Gender:Male I/FU:I

Company Report #CTU 236217

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Neurontin (Generic)	PS		
300 MG BID							

Date:01/11/05ISR Number: 4550538-1Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 236158

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Feeling Hot Pain In Extremity Pharmaceutical Product Complaint Skin Burning Sensation		Generic Gabapentin	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/11/05ISR Number: 4551176-7Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 236178

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anticonvulsant Drug Level		Gabapentin	PS		ORAL
1200 MG PO		Decreased					
TID		Condition Aggravated		Tegretol Xr	C		
		Convulsion		Keppra	C		
		Pharmaceutical Product		Diazepam	C		
		Complaint		Trazodone	C		

Date:01/11/05ISR Number: 4551178-0Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 236180

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
400 MG 2 TID		Drug Effect Decreased		Neurontin (Generic)	PS		
		Pharmaceutical Product					
		Complaint					

Date:01/11/05ISR Number: 4551202-5Report Type:Direct
 Age:41 YR Gender:Male I/FU:I

Company Report #CTU 236190

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cheilitis		Gabapentin 300 Mg			
300 MG PO QHS		Oral Mucosal Blistering		Capsules	PS		ORAL
		Pharmaceutical Product					
		Complaint					
		Stomatitis					
		Tongue Ulceration					

Date:01/11/05ISR Number: 4551248-7Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 236206

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Epistaxis		Gabapentin Generic	PS		ORAL
300 MG PO TID		Lethargy					
		Nausea					
		Pharmaceutical Product					
		Complaint					
		Vomiting					

Date:01/11/05ISR Number: 4551626-6Report Type:Expedited (15-DaCompany Report #2005004215
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin			
Other		Completed Suicide		(Gabapentin)	PS		
		Drug Ineffective					

Date:01/11/05ISR Number: 4551842-3Report Type:Expedited (15-DaCompany Report #CIP04002758
Age:81 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Antibody Test Positive					
Initial or Prolonged		Complement Factor					
		Increased					
		Inflammation					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Leukocytoclastic Vasculitis Pain Of Skin	Report Source	Product	Role	Manufacturer	Route
1 CAPSUL		Pulmonary Embolism Staphylococcal Infection Urinary Tract Infection Vascular Purpura	Foreign Health Professional Other	Furadantine (Nitrofurantoin Macrocrystals) Crystals	PS		ORAL
DAILY ORAL		Vasculitis Necrotising		Previscan (Fluidione) Tablet 20 Mg	SS		ORAL
1 TABLET				Zocor "Dieckmann" (Simvastatin)	SS		ORAL
DAILY ORAL				Xatral(Alfuzosin)	SS		ORAL
ORAL				Neurontin (Gabapentin)	SS		ORAL
ORAL				Renitec (Enalapril Maleate) Vaccines	C C		

Date:01/11/05ISR Number: 4552851-0Report Type:Direct
Age:29 YR Gender:Female I/FU:I

Company Report #CTU 236260

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300MG TID PO		Drug Ineffective		Gabapentin	PS		ORAL
		Drug Intolerance Pharmaceutical Product Complaint					

Date:01/12/05ISR Number: 4550575-7Report Type:Expedited (15-DaCompany Report #FR-ABBOTT-04P-056-0283040-00
Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Chills		Kaletra Soft Gelatin Capsules	PS		
Hospitalization - Initial or Prolonged		Coma		Kaletra Soft Gelatin Capsules	SS		
Other		Hepatic Encephalopathy		Zidovudine			
		Hepatic Failure		W/Lamivudine	SS		ORAL
		Oedema		Zidovudine			
		Pyrexia		W/Lamivudine	SS		
1	MON			Ribavirin	SS		ORAL
				Perinterferon	SS		
				Buprenorphine	SS		ORAL
				Gabapentin	SS		ORAL
				Minocycline	C		
				Clarithromycin	C		
				Pentamidine	C		
				Gabapentin	C		
2	YR						

Date:01/12/05ISR Number: 4552041-1Report Type:Expedited (15-DaCompany Report #2005001941
Age:27 YR Gender:Female I/FU:I

Outcome PT
Hospitalization - Abortion Threatened
Initial or Prolonged Drug Exposure During
Pregnancy

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pregnancy Vomiting	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin)	PS		
				Zonisamide (Zonisamide)	SS		
				Valproate Semisodium (Valproate Semisodium)	C		
				Levetiracetam (Levetiracetam)	C		

Date:01/12/05
 Age:49 YR
 Gender:Male
 I/FU:I

ISR Number: 4552091-5
 Report Type:Expedited (15-DaCompany Report #2005004121

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Blood Creatine Phosphokinase Increased	Consumer	Neurontin (Gabapentin)	PS		
Other		Blood Testosterone Decreased		Lipitor (Atorvastatin)	SS		
10 MG		Chest Pain		Bextra (Valdecoxib)	SS		
		Demyelination		Levothyroxine Sodium (Levothyroxine Sodium)	SS		
0.5 MG		Depression					
		Drug Ineffective					
		Drug Toxicity		Perindopril Erbumine (Perindopril Erbumine)	SS		
		Fatigue					
		Feeling Abnormal					
		Hypertension		Hydrochlorothiazide (Hydrochlorothiazide)	SS		
		Hypoaesthesia					
		Lethargy					
		Mental Disorder		Topiramate (Topiramate)	C		
		Muscle Disorder					
		Muscle Tightness		Naproxen Sodium (Naproxen Sodium)	C		
		Muscle Twitching					
		Muscular Weakness		Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
		Myalgia					
		Nervousness					
		Pain In Extremity					
		Sleep Apnoea Syndrome					
		Thyroid Function Test					

Abnormal
Viral Infection

Date:01/12/05ISR Number: 4552337-3Report Type:Direct
Age:81 YR Gender:Male I/FU:I

Company Report #CTU 236456

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Enuresis		Gabapentin 300mg	PS	Greenstone	ORAL
ONE BY MOUTH		Pharmaceutical Product					
AT BEDTIME		Complaint					

Date:01/13/05ISR Number: 4553093-5Report Type:Expedited (15-DaCompany Report #2005006045
Age: Gender:Male I/FU:I

Outcome
Hospitalization -
Initial or Prolonged

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ORAL		Neurontin Capsules	SS	ORAL
400 MG TID				
ORAL		Kaletra (Lopinavir/Ritonavir) Capsules	SS	ORAL
3 DOSES BID				
ORAL		Combivir (Lamivudine/Zidovudi ne) Capsules	SS	ORAL
150 MG BID				
ORAL		Subutex (Buprenorphine Hcl) Sublingual Tablets	SS	ORAL
8 MG QD ORAL				
		Zeclar (Clarithromycin)	C	
		Minocycline	C	
		Aldactone	C	
		Pentacarinat (Pentamidine Isethionate)	C	
		Lasilix	C	

Date:01/13/05ISR Number: 4553881-5Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 236674

Outcome PT
Other Hypersensitivity
Pharmaceutical Product

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Complaint Rash				
Dose	Duration		Report Source	Product	Role	Manufacturer
1 PO QD				Neurontin 400 1 Po Qd	PS	
						Route ORAL

Date:01/13/05ISR Number: 4553882-7Report Type:Direct
Age:55 YR Gender:Male I/FU:I

Company Report #CTU 236673

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aura Drug Effect Decreased		Gabapentin 300 Mg (Neurontin Generic)	PS		ORAL
300 MG PO BID	2	MON					
		Pharmaceutical Product Complaint		Primidone 50mg (Mysoline Generic)	SS		ORAL
50MG PO Q HS	2	MON					
		Simple Partial Seizures		Tegretol Depakote	C C		

Date:01/13/05ISR Number: 4553887-6Report Type:Direct
Age:55 YR Gender:Male I/FU:I

Company Report #CTU 236665

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation Pharmaceutical Product Complaint		Gabapentin (Generic For Neurontin)(300 Mg Tid)	PS		ORAL
300 MG PO TID							
		Rash					

Date:01/13/05ISR Number: 4554100-6Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 236683

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - ONCE DAILY Initial or Prolonged		Anhedonia Depression		Neurontin 600 Mg	PS		

Disability
 Fatigue
 Panic Attack
 Somnolence
 Suicidal Ideation

Date:01/14/05ISR Number: 4552640-7Report Type:Expedited (15-DaCompany Report #FR-RB-1183-2004
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Hepatic Encephalopathy		Peginterferon			
Hospitalization - SUBCUTANEOUS		Hepatic Failure		Alfa-2b	PS		
Initial or Prolonged		Oedema		Ribavirin	SS		ORAL
				Gabapentin	SS		ORAL
				Kaletra	SS		ORAL
				Combivir	SS		ORAL
				Buprenorphine Hydrochloride	SS		ORAL
				Clarithromycin	C		
UNKNOWN	UNKNOWN			Minocycline	C		
UNKNOWN	UNKNOWN			Spirolactone	C		
UNKNOWN	UNKNOWN			Pentamidine Isethionate	C		
DOSAGE				Furosemide	C		
UNKNOWN	UNKNOWN						
DOSAGE							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/05ISR Number: 4552650-XReport Type:Expedited (15-DaCompany Report #PHBS2005GB00599
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Arrest		Carbamazepine	PS	Novartis Sector:	
UNKNOWN		Cardiac Pacemaker				Pharma	
		Insertion		Gabapentin	SS		
UNKNOWN							
		Sinus Tachycardia		Levetiracetam	SS		
UNKNOWN							

Date:01/14/05ISR Number: 4552820-0Report Type:Expedited (15-DaCompany Report #US-ROCHE-376030
 Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Drug Interaction		Klonopin	PS	Roche	ORAL
REPORTED		Hypoglycaemia					
START DATE AS							
PRIOR TO 16							
JANUARY 2001.							
				Lamictal	I		ORAL
				Neurontin	I		ORAL
DOSING AMOUNT							
REPORTED AS:							
"2/1/1/2".							
				Xanax	I		ORAL
				Gabitril	I		ORAL
15 DAY							
				Celexa	I		ORAL

Date:01/14/05ISR Number: 4556084-3Report Type:Expedited (15-DaCompany Report #2005006201
 Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death Life-Threatening 1200 MG (400 Hospitalization - MG, 3 IN 1 Initial or Prolonged D), ORAL Other	Encephalopathy Hepatic Failure	Foreign Health Professional	Neurontin (Gabapentin)	PS	ORAL
300 MG (150 MG, 2 IN 1 D), ORAL			Zidovudine W/Lamivudine (Lamivudine, Zidovudine)	SS	ORAL
1 GRAM (1 IN 1 D), ORAL			Ribavirin (Ribavirin)	SS	ORAL
(2 IN 1 D), ORAL			Kaletra (Lopinavir, Ritonavir)	SS	ORAL
			Peginterferon Alfa-2b (Peginterferon Alfa-2b)	SS	
SUBCUTANEOUS	SUBCUTANEOUS		Buprenorphine Hydrochloride (Buprenorphine Hydrochloride)	SS	
SUBLINGUAL	8 MG (8 MG, 1 IN 1 D), SUBLINGUAL				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/05ISR Number: 4556429-4Report Type:Expedited (15-DaCompany Report #2005006035

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Cardiac Disorder	Consumer	Neurontin			
Other		Condition Aggravated		(Gabapentin)	PS		
600 MG (300		Drug Ineffective					
MG, 2 IN 1 D)		Emphysema		Sulfasalazine			
		Neuropathy		(Sulfasalazine)	C		
		Oxygen Saturation		Lisinopril			
		Abnormal		(Lisinopril)	C		
		Visual Disturbance		Glipizide			
				(Glipizide)	C		
				Pravastatin Sodium			
				(Pravastatin Sodium)	C		
				Ezetimibe			
				(Ezetimibe)	C		
				Salbutamol			
				(Salbutamol)	C		
				Furosemide			
				(Furosemide)	C		

Date:01/14/05ISR Number: 4556455-5Report Type:Expedited (15-DaCompany Report #2005002381

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiac Disorder	Consumer	Neurontin (Tablets)			
Other		Cerebrovascular Accident		(Gabapentin)	PS		ORAL
(300 MG),		Echocardiogram Abnormal					
ORAL		Gastric Ulcer Haemorrhage		Warfarin Sodium			
		Pain		(Warfarin Sodium)	SS		
		Treatment Noncompliance		Moracizine			
		Vascular Calcification		Hydrochloride			
		Vomiting		(Moracizine			
				Hydrochloride)	C		
				Atenolol (Atenolol)	C		
				Acetylsalicylic Acid			
				(Acetylsalicylic			
				Acid)	C		

Glyceryl Trinitrate	
(Glyceryl	
Trinitrate)	C
Atorvastatin	
(Atorvastatin)	C
Prinzide	
(Hydrochlorothiazide	
, Lisinopril)	C
Fenofibrate	
(Fenofibrate)	C
Lansoprazole	
(Lansoprazole)	C
Lidocaine	
Hydrochloride	
(Lidocaine	
Hydrochloride)	C
Fentanyl (Fentanyl)	C
Metoclopramide	
(Metoclopramide)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/14/05ISR Number: 4556717-1Report Type:Expedited (15-DaCompany Report #2004100399

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Mental Disorder	Consumer	Neurontin			
Initial or Prolonged		Nerve Injury		(Gabapentin)	PS		
Other		Pancreatitis		Risperidone			
		Sexual Dysfunction		(Risperidone)	SS		ORAL
ORAL		Weight Increased		All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:01/14/05ISR Number: 4556723-7Report Type:Expedited (15-DaCompany Report #2005002357

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abnormal Behaviour	Consumer	Neurontin			
		Disturbance In Attention		(Gabapentin)	PS		ORAL
1800 MG (300		Drug Ineffective For					
MG, 1 IN 4		Unapproved Indication					
HR), ORAL		Increased Appetite		Cannabis (Cannabis)	SS		
RESPIRATORY		Medical Device					
(INHALATION)	INHALATION	Complication		Losartan Potassium			
		Memory Impairment		(Losartan Potassium)	C		
		Unevaluable Event		Pilocarpine			
		Weight Increased		Hydrochloride			
				(Pilocarpine			
				Hydrochloride)	C		
				Levothyroxine Sodium			
				(Levothyroxine			
				Sodium)	C		
				Zolpidem Tartrate			
				(Zolpidem Tartrate)	C		

Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Chills		Combivir	PS	Glaxosmithkline	ORAL
2 YR							
Hospitalization -		Coma		Kaletra	SS		ORAL
2 YR							
Initial or Prolonged		Hepatic Cirrhosis		Rebetol	SS		ORAL
1000MG per							
day	DAY	Hepatic Encephalopathy					
		Hepatic Failure		Alpha 2b Interferon	SS		
SUBCUTANEOUS	120MCG	per					
day		Liver Disorder					
		Oedema		Subutex	SS		ORAL
8MG Per day							
		Pyrexia		Neurontin	SS		ORAL
400MG Three							
times per day	2	YR					
				Zeclar	SS		ORAL
				Minocycline	SS		ORAL
				Aldactone	SS		
UNKNOWN							
				Pentacarinat	SS		
RESPIRATORY							
(INHALATION)							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/18/05ISR Number: 4555346-3Report Type:Direct
Age:36 YR Gender:Female I/FU:I

Company Report #CTU 236921

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin 300mg			
ORAL		Drug Ineffective Pharmaceutical Product		Gabapentin	PS		ORAL
		Complaint					

Date:01/18/05ISR Number: 4555556-5Report Type:Direct
Age:56 YR Gender:Female I/FU:I

Company Report #CTU 237054

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin-Gabapentin			
Other		Feeling Abnormal Nausea		600 Mg Qid	PS		ORAL
600 MG QID PO		Pharmaceutical Product Complaint		Duragesic	C		
				Lidoderm Patches	C		
				Lortab	C		
				Flexeril	C		

Date:01/18/05ISR Number: 4555712-6Report Type:Periodic
Age:89 YR Gender:Female I/FU:I

Company Report #A001-002-005941

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Aricept (Donepezil			
Required		Condition Aggravated	Consumer	Hydrochloride)	PS		ORAL
Intervention to		Convulsion					
10 MG, 1 IN 1		Dizziness					
Prevent Permanent		Fatigue		Neurontin			
D, ORAL				(Gabapentin)	SS		
Impairment/Damage							
SEE IMAGE				Memantine			
				(Memantine)	SS		ORAL
ORAL				Antihypertensive			
				(Antihypertensive			
				(Antihypertensive)	C		
				Aggrenox (Asasantin)	C		

Date:01/18/05ISR Number: 4555843-0Report Type:Periodic
Age:51 YR Gender:Male I/FU:I

Company Report #A001-002-006183

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Required Intervention to SUBCUTANEOUS Prevent Permanent SUBCUTANEOUS Impairment/Damage	200 MCG,	Convulsion Mental Status Changes	Health Professional	Aricept (Donepezil Hydrochloride)	PS		ORAL
INTRA	VENOUS	INTRA	VENOUS	Aranesp (Darbepoetin Alfa)	SS		
				Marinol (Dronabinol)	SS		
				Kytril (Granisetron)	SS		
(NOT				Oxycontin (Oxycodone Hydrochloride)	SS		
20 MG, 1 IN							
12 HR				Neurontin (Gabapentin)	SS		
				Ambien (Zolpidem Tartrate)	SS		
5 MG, 1 IN 4				Oxy Ir (Oxycodone Hydrochloride)	SS		
HR				Morphine Sulfate	C		
				Chlorpromazine	C		

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Prograf (Tacrolimus)	C
Protonix (Pantoprazole)	C
Prandin (Repaglinide)	C
Zoloft (Sertraline Hydrochloride)	C
Glipizide	C
Procardia (Nifedipine)	C
Magnesium	C
Levaquin (Levofloxacin)	C
Pentamidine	C
Peri-Colace	C
Dulcolax (Bisacodyl)	C
Baby Aspirin (Acetylsalicylic Acid)	C
Dexamethasone	C
Adriamycin	C
Cisplatin	C
Senokot (Senna Fruit)	C
Inderal (Propranolol Hydrochloride)	C
Multivitamins	C
Insulin	C
Aldactone (Spironolactone)	C
Megace (Megestrol Acetate)	C

Date:01/18/05ISR Number: 4556552-4Report Type:Expedited (15-DaCompany Report #2004009355
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Accident
Other	Back Disorder
	Back Pain
	Blood Cholesterol
	Increased
	Chest Pain
	Chills

Condition Aggravated
Contusion
Disorientation
Drug Effect Decreased
Drug Ineffective
Drug Intolerance
Erectile Dysfunction
Facial Pain
Feeling Abnormal
Feeling Cold
Gastroesophageal Reflux
Disease
Gingival Disorder
Glossodynia
Hostility

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Adverse Event	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, 3 IN 1 D), ORAL		Hyporeflexia Impaired Healing Injury	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Oesophageal Spasm	Professional				
20 MG (1 IN 1 D), ORAL		Sedation Sleep Disorder Somnolence		Lipitor (Atorvastatin)	SS		ORAL
		Tooth Abscess					
1000 MG (1 IN 2 D), ORAL		Treatment Noncompliance Tremor Vertigo		Lithium (Lithium) (Lithium)	SS		
				Naproxen (Naproxen)	SS		
				Rofecoxib (Rofecoxib)	SS		
				Amoxicillin (Amoxicillin)	SS		ORAL
				All Other Therapeutic Products (All Other Therapeutic Products)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Vitamins (Vitamins)	C		
				Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Body Height Decreased Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL
SEE IMAGE							
		Nerve Injury Pain		Gabapentin (Gabapentin)	SS		
UNKNOWN	UNKNOWN						
		Pain In Extremity Pharmaceutical Product Complaint Pruritus		Levothyroxine Sodium (Levothyroxine Sodium) Losartan Potassium (Losartan Potassium) Famotidine (Famotidine) Verapamil (Verapamil) Trazodone (Trazodone) Tizanidine (Tizanidine) Cevimeline Hydrochloride (Cevimeline Hydrochloride) All Other Therapeutic Products	C C C C C C C C C C		

Freedom Of Information (FOI) Report

(All Other
Therapeutic
Products) C
Erythromycin
(Erythromycin) C
Estrogens Conjugated
(Estrogens
Conjugated) C
Multivitamins With
Minerals
(Multivitamins With
Minerals) C
B-Komplex "Leciva"
(Calcium
Pantothenate,
Nicotinamide,
Pyridoxine C
Fish Oil (Fish Oil) C
Magnesium
(Magnesium) C
Glucosamine
(Glucosamine) C
Calcium (Calcium) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C
Paracetamol
(Paracetamol) C
Tears Naturale
(Dextran 70,
Hyppromellose) C
Carmellose Sodium
(Carmellose Sodium) C

Date:01/18/05ISR Number: 4556571-8Report Type:Expedited (15-DaCompany Report #2005006149
Age:39 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Dysarthria Eating Disorder Feeling Abnormal Mental Disorder Pneumonia Staring	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:01/18/05ISR Number: 4556574-3Report Type:Expedited (15-DaCompany Report #2004230338US
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Dizziness Drug Interaction	Health Professional	Xanax Tablet (Alprazolam)	PS		ORAL
0.5 MG (AS NEEDED), ORAL		Hypoglycaemia Vision Blurred		Neurontin (Gabapentin) (Gabapentin)	SS		ORAL
(300 MG, UNKNOWN), ORAL				Tiagabine Hydrochloride (Tiagabine			

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20 MG (1 IN 1			Hydrochloride)	SS		ORAL
D), ORAL						
			Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		ORAL
20 MG						
(UNKNOWN),						
ORAL						
UNKNOWN	UNKNOWN		Clonazepam (Clonazepam)	SS		
UNKNOWN	UNK (1 IN 2		Lamotrigine (Lamotrigine)	SS		
D), UNKNOWN						

Date:01/18/05ISR Number: 4556596-2Report Type:Expedited (15-DaCompany Report #2005006105
 Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Gastrointestinal Haemorrhage	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (300							
MG, 1 IN 1							
D), ORAL							
				Balsalazide Sodium (Balsalazide Sodium)	C		
				Diazyde (Hydrochlorothiazide)	C		
				Nifedipine (Nifedipine)	C		
				Raloxifene Hydrochloride (Raloxifene Hydrochloride)	C		
				Multivitamins And			

Iron (Multivitamins
 And Iron) C
 Glucosamine
 (Glucosamine) C
 Nortriptyline
 (Nortriptyline) C
 Atorvastatin
 (Atorvastatin) C

Date:01/18/05ISR Number: 4556599-8Report Type:Expedited (15-DaCompany Report #2005004260

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional	Neurontin (Gabapentin)	PS		

Date:01/18/05ISR Number: 4556621-9Report Type:Expedited (15-DaCompany Report #2005006133

Age:76 YR Gender:Female I/FU:I

Outcome	PT
Other	Abasia Depression Dysstasia Eye Disorder Fall

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Glaucoma Surgery Hypotrichosis Lethargy	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Muscle Spasms Paraplegia					
		Skin Discolouration Skin Exfoliation Skin Nodule Spinal Cord Compression Viral Infection Vision Blurred		Metformin Hydrochloride (Metformin Hydrochloride) Furosemide (Furosemide) Clopidogrel Sulfate (Clopidogrel Sulfate) Metoprolol Tartrate (Metoprolol Tartrate) Desloratadine (Desloratadine) Amitriptyline (Amitriptyline)	C C C C C C		

Date:01/18/05ISR Number: 4556624-4Report Type:Expedited (15-DaCompany Report #2005007328
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Anxiety Completed Suicide Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		

Date:01/18/05ISR Number: 4556629-3Report Type:Expedited (15-DaCompany Report #2005002328
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other		Balance Disorder Constipation Hallucination Incoherent Weight Decreased	Consumer	Neurontin (Gabapentin) Alprazolam (Alprazolam) Analgesics	PS C C		

Lovastatin
 (Lovastatin) C
 Metoprolol Tartrate
 (Metoprolol
 Tartrate) C
 Levothyroxine Sodium
 (Levothyroxine
 Sodium) C
 Losartan Potassium
 (Losartan Potassium) C

Date:01/18/05ISR Number: 4556631-1Report Type:Expedited (15-DaCompany Report #2005007511
 Age:59 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/18/05ISR Number: 4556769-9Report Type:Expedited (15-DaCompany Report #S04-USA-05512-01
Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Dizziness	Health	Celexa (Citalopram			
Other		Drug Interaction	Professional	Hydrobromide)	PS		ORAL
40 MG QHS PO							
		Dystonia		Gabitril (Tiagabine			
4 MG BID		Hallucination, Auditory		Hydrochloride)	SS		
		Hypoglycaemia		Neurontin			
1800 MG QD		Oedema Peripheral		(Gabapentin)	SS		
		Vision Blurred		Xanax (Alprazolam)	SS		
0.5 MG PRN				Klonopin			
				(Clonazepam)	SS		
0.5 MG BID				Lamictal	SS		
100 MG QAM				Lamictal	SS		
50 MG QHS							

Date:01/18/05ISR Number: 4557219-9Report Type:Expedited (15-DaCompany Report #DEU-2004-0001150
Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Interaction	Foreign	Oxygesic 10 Mg			
		International Normalised	Health	(Oxycodone			
		Ratio Decreased	Professional	Hydrochloride) Cr			
SEE IMAGE		Prothrombin Time	Other	Tablet	PS		ORAL
		Shortened		Marcumar			
TABLET, ORAL				(Phenprocoumon)	SS		ORAL
				Amitriptyline			
10 MG, BID,				(Amitriptyline)	SS		ORAL
ORAL				Gabapentin			
				(Gabapentin)	SS		ORAL
1200 MG,							

DAILY, ORAL

Isosorbide
Mononitrate C
Pantozol
(Pantoprazole
Sodium) C
Beloc-Zok Forte
(Metoprolol
Succinate) C
Atorvastatin Calcium
(Atorvastatin
Calcium) C

Date:01/18/05ISR Number: 4557292-8Report Type:Expedited (15-DaCompany Report #2005006212
Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Amnesia Fatigue	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL		Hypercholesterolaemia Hypertriglyceridaemia Somnolence Weight Increased	Professional	Estradiol (Estradiol) Dydrogesterone (Dydrogesterone)	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/18/05ISR Number: 4557295-3Report Type:Expedited (15-DaCompany Report #2004110900
 Age:21 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Cough	Foreign	Neurontin			
Hospitalization -	Dysphagia	Health	(Gabapentin)	PS		ORAL
1500 MG (500						
Initial or Prolonged	Retching	Professional				
MG, 3 IN 1						
Other						
D), ORAL						
			Valproate Sodium			
			(Valproate Sodium)	C		
			Clobazam (Clobazam)	C		
			Anovlar			
			(Ethinylestradiol,			
			Norethisterone			
			Acetate)	C		
			Zuclopenthixol			
			(Zuclopenthixol)	C		

Date:01/19/05ISR Number: 4557116-9Report Type:Direct Company Report #CTU 237178
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Pain		Neurontin			
	Pharmaceutical Product		/Gabapentin	PS		ORAL
ORAL , TID						
300 -900 MG	Complaint					

Date:01/19/05ISR Number: 4558591-6Report Type:Expedited (15-DaCompany Report #2004056018
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Other	Abnormal Behaviour	Consumer	Neurontin			
	Adverse Event	Health	(Gabapentin)	PS		
	Amnesia	Professional	Morphine (Morphine)	SS		
	Economic Problem		All Other			
	Feeling Hot		Non-Therapeutic			

Granuloma
Heart Rate Increased
Hot Flush
Hyperhidrosis
Inadequate Analgesia
Insomnia
Loss Of Consciousness
Medical Device
Complication
Nerve Injury
Oedema
Pain
Speech Disorder
Spinal Disorder
Weight Increased

Products (All Other
Non-Therapeutic
Products) SS
Baclofen (Baclofen) SS

Date:01/19/05ISR Number: 4558623-5Report Type:Expedited (15-DaCompany Report #2004085287
Age: Gender:Female I/FU:I

Outcome PT
Other Cardiac Disorder
Depression
Muscle Spasms
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
600 MG (300 MG, 2 IN 1 CYCLICAL), ORAL		Oedema Peripheral Poor Peripheral Circulation Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:01/19/05
Age:83 YR
Gender:Male
ISR Number: 4558625-9
Report Type:Expedited (15-DaCompany Report #2004078403
I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1200 MG (400 MG, 3 IN 1 D)		Tendon Repair Urinary Incontinence Urinary Retention Urinary Tract Infection	Consumer Health Professional	Neurontin (Gabapentin)	PS		
					Atenolol (Atenolol)	C		
					Ranitidine Hydrochloride (Ranitidine Hydrochloride)	C		
					Diazepam (Diazepam)	C		
					Centrum Silver (Ascorbic Acid, Calcium, Minerals Nos, Retinol, Tocopheryl Acetate, Tocopherol (Tocopherol)	C		
					Irbesartan (Irbesartan)	C		
					Hydrochlorothiazide (Hydrochlorothiazide)	C		
					Clopidogrel Sulfate (Clopidogrel Sulfate)	C		

Vicodin (Hydrocodone
Bitartrate,
Paracetamol) C
Paracetamol
(Paracetamol) C

Date:01/19/05ISR Number: 4558631-4Report Type:Expedited (15-DaCompany Report #2005007436
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Emotional Disorder	Consumer	Neurontin			
Other		Mental Disorder Overdose		(Gabapentin)	PS		

Date:01/19/05ISR Number: 4558651-XReport Type:Expedited (15-DaCompany Report #2004119995
Age: Gender:Female I/FU:F

Outcome	PT
Disability	Abdominal Operation
Other	Fall Limb Operation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Post Procedural Complication Tremor	Report Source	Product	Role	Manufacturer	Route
3000 MG (600 MG, 5 IN 1 D), ORAL			Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
ORAL				Deltasone (Prednisone)	SS		ORAL
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Rosuvastatin (Rosuvastatin)	C		
				Folic Acid (Folic Acid)	C		
				Trazodone (Trazodone)	C		
				Warfarin (Warfarin)	C		
				Infliximab (Infliximab)	C		
				Ranitidine (Ranitidine)	C		
				Amitriptyline (Amitriptyline)	C		

Date:01/19/05ISR Number: 4558690-9Report Type:Expedited (15-DaCompany Report #2005006446

Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abasia Emotional Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 2 IN 1 D), ORAL		Pain Treatment Noncompliance		Atorvastatin (Atorvastatin)	C		

Captopril
(Captopril) C
Insulin (Insulin) C

Date:01/20/05ISR Number: 4559978-8Report Type:Direct
Age:61 YR Gender:Female I/FU:I

Company Report #CTU 237323

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Stomach Discomfort		Gabapentin 700 Mg /D	PS		
300 MG Q AM,							
400 MG Q HS							

Date:01/20/05ISR Number: 4559993-4Report Type:Direct
Age:89 YR Gender:Female I/FU:I

Company Report #CTU 237419

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Nausea		Neurontin			
		Pharmaceutical Product		(Gabapentin)	PS		
300 MGM 6AM		Complaint					
; 200 MGM		Rash					
NOON ; 100							
MGM 8PM							
				Nortriptyline	C		
				Lorazepam	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochlorothiazide C
 Prevacid C
 Maxaquin C
 Norvasc C
 Duragesic Patch C
 Advair Diskus C

Date:01/20/05ISR Number: 4560097-5Report Type:Direct
 Age:41 YR Gender:Female I/FU:I

Company Report #CTU 237484

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Neurontin	PS		
300 MG (3) AT		Pharmaceutical Product					
QHS		Complaint		Wellbutrin Sr	C		

Date:01/20/05ISR Number: 4560231-7Report Type:Direct
 Age:71 YR Gender:Female I/FU:I

Company Report #CTU 237470

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Faeces Atrial Fibrillation		Neurontin 100 & 300mg. Warner-Lambert	PS	Warner-Lambert	ORAL
ORAL				Pancrease Caps Xanax Extra Strength Tylenol Cap	C C C		

Date:01/21/05ISR Number: 4557879-2Report Type:Periodic
 Age:52 YR Gender:Male I/FU:F

Company Report #US-BRISTOL-MYERS SQUIBB COMPANY-12413118

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hepatitis	Health Professional	Serzone Tabs	PS	Bristol-Myers Squibb Company	ORAL
100mg							

tablets; 1 to

4 tabs at hs.

Stopped

Dec-2000,

Lipitor

SS

On 10mg/day

as of

06-Feb-2000.

Stopped

Dec-2000,

Gabapentin

SS

ORAL

100mg, 300mg, 8

00mg caps. 1-3

hs-tid

prn. Stop

12/00, restart

Gabapentin

SS

ORAL

100mg, 300mg, 8

00mg caps. 1-3

hs-tid

prn. Stop

12/00, restart

Gabapentin

SS

ORAL

100mg, 300mg, 8

00mg caps. 1-3

hs-tid

prn. Stop

12/00, restart

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/21/05ISR Number: 4558787-3Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0340937A
 Age:51 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - UNKNOWN	Abdominal Pain		Zophren	PS	Glaxosmithkline	
Initial or Prolonged 8 WK	Antinuclear Antibody		Neurontin	SS		ORAL
INTRAVENOUS 110MG	Positive Cyclic		Taxotere	SS		
INTRAVENOUS 70MG	Aphthous Stomatitis Cyclic		Caelyx	SS		
250MG Three times per day	Catheter Site Infection		Becilan	C		ORAL
	Diarrhoea					
	Erythema					
	Nikolsky'S Sign					
	Oedema Peripheral					
	Pain					
	Pain In Extremity					
	Pyrexia					
	Rash Erythematous					
	Skin Exfoliation					
	Staphylococcal Infection					
	Stevens-Johnson Syndrome					

Date:01/21/05ISR Number: 4559184-7Report Type:Direct Company Report #CTU 237566
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900MG TID	Fall Movement Disorder		Neurontin 300mg Parke-Davis	PS	Parke-Davis	ORAL
ORAL	Myoclonus					
	Treatment Noncompliance		Citalopram			
			Hydrobromide	C		
			Gabapentin	C		
			Ibuprofen	C		
			Phenobarbital	C		
			Phenytoin	C		

Date:01/21/05ISR Number: 4561120-4Report Type:Expedited (15-DaCompany Report #2005AP00140
Age:75 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 25 MG DAILY	Cardiac Disorder	Health	Atenolol	PS		ORAL
Intervention to PO	Diabetes Mellitus	Professional				
Prevent Permanent 80 MG DAILY	Myocardial Infarction		Lipitor	SS		
Impairment/Damage 300 MG TID PO	Nervous System Disorder		Neurontin	SS		ORAL
	Tremor		Glipizide	SS		ORAL
10 MG BID PO			Prevacid	SS		ORAL
30 MG BID PO			Altace	SS		ORAL
2.5 MG DAILY						
PO			Nitrotab	C		
			Plavix	C		

Date:01/21/05ISR Number: 4561130-7Report Type:Expedited (15-DaCompany Report #US013912
Age:52 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Atelectasis Balance Disorder Convulsion Disorientation 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
		Hypotension Influenza Like Illness Lung Infiltration	Health	Provigil	PS		ORAL
100 MG QD		Myalgia	Professional				
ORAL		Pneumonia		Provigil	SS		ORAL
200 MG QD		Pyuria					
ORAL		Septic Shock		Provigil	SS		ORAL
50 MG ONCE		Urinary Tract Infection					
ORAL		Vomiting		Klonopin	SS		ORAL
2.25 MG QD							
ORAL				Neurontin	SS		ORAL
1200 MG QD							
ORAL				Aciphex	SS		ORAL
60 MG QD ORAL				Hydrochlorothiazide	SS		ORAL
100 MG QD							
ORAL				Doryx	SS		ORAL
100 MG QD							
ORAL				Lexapro	SS		ORAL
10 MG QD ORAL							

Date:01/21/05ISR Number: 4561380-XReport Type:Expedited (15-DaCompany Report #2005008379

Age: Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Aneurysm	Health	Neurontin			
Other		Gastroenteritis Viral	Professional	(Gabapentin)	PS		
100 MG (100							

Pain

Treatment Noncompliance

Oxycodone	
Hydrochloride	
(Oxycodone	
Hydrochloride)	C
Lorazepam	
(Lorazepam)	C
Escitalopram	
(Escitalopram)	C
Atenolol (Atenolol)	C
Rabeprazole Sodium	
(Rabeprazole Sodium)	C
Thyroid (Thyroid)	C
Lidocaine	
Hydrochloride	
(Lidocaine	
Hydrochloride)	C

Date:01/21/05ISR Number: 4561422-1Report Type:Expedited (15-DaCompany Report #2004086847

Age: Gender:Female I/FU:F

Outcome

PT

Other

Alopecia
 Confusional State
 Depression
 Diverticulitis
 Dysarthria
 Dysuria
 Feeling Abnormal
 Hypertension
 Metrorrhagia
 Oedema Peripheral
 Pain In Extremity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Prescribed Overdose Suicide Attempt Uterine Leiomyoma	Report Source	Product	Role	Manufacturer	Route
100 MG (100 MG, 1 IN 1 D), ORAL		Vulvovaginal Disorder	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Zolmitriptan (Zolmitriptan)	C		
				Sumatriptan Succinate (Sumatriptan Succinate)	C		
				Axotal (Old Form) (Butalbital, Caffeine, Paracetamol)	C		
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		
				Alprazolam (Alprazolam)	C		
				Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
				Promethazine (Promethazine)	C		
				Topiramate (Topiramate)	C		
				Lomotil (Atropine Sulfate, Diphenoxylate Hydrochloride)	C		
				Verapamil Hydrochloride (Verapamil Hydrochloride)	C		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Neurontin			
		Drug Ineffective		(Gabapentin)	PS		ORAL
1800 MG (600		Lung Disorder					
MG, 3 IN 1		Pain					
D), ORAL		Tremor		Xanax Tablet			
				(Alprazolam)	SS		
0.5 MG				Morphine (Morphine)	SS		
				Gabapentin			
				(Gabapentin)	SS		
2400 MG (600							
MG, 4 IN 1 D)				Montelukast Sodium			
				(Montelukast Sodium)	C		

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Freedom Of Information (FOI) Report

Theophylline (Theophylline)	C
Udramil (Trandolapril, Verapamil Hydrochloride)	C
Insulin Novolin 70/30 (Insulin Human Semisynthetic, Insulin Isophane Human Semisynthetic)	C
Duloxetine Hydrochloride (Duloxetine Hydrochloride)	C
Orphenadrine Citrate (Orphenadrine Citrate)	C
Furosemide (Furosemide)	C
Magnesium Hydroxide (Magnesium Hydroxide)	C
Vicodin (Hydrocodone Bitartrate, Paracetamol)	C
Isosorbide Mononitrate (Isosorbide Mononitrate)	C
Lisinopril (Lisinopril)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C

Date:01/21/05ISR Number: 4561429-4Report Type:Expedited (15-DaCompany Report #2005006045
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged SEE IMAGE	Anticonvulsant Drug Level Increased	Consumer	Dilantin Kapseals (Phenytoin Sodium)	PS		ORAL
Other	Aortic Stenosis		Neurontin			

600 MG (300	Condition Aggravated	(Gabapentin)	SS	ORAL
MG, 2 IN 1	Convulsion			
D), ORAL	Dementia			
	Fatigue	Phenytoin		
	Hunger	(Phenytoin)	SS	
	Incontinence	Risperidone		
1 MG (0.5 MG,	Loss Of Consciousness	(Risperidone)	SS	ORAL
2 IN 1 D),	Medication Error			
ORAL	Overdose			
	Pharmaceutical Product	Furosemide		
	Complaint	(Furosemide)	C	
	Prostatic Disorder			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/21/05ISR Number: 4562212-6Report Type:Expedited (15-DaCompany Report #2005009641

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Acne	Consumer	Neurontin			
Initial or Prolonged	Bone Pain		(Gabapentin)	PS		
Other	Drug Ineffective		Anovlar			
	Drug Tolerance		(Ethinylestradiol,			
	Fall		Norethisterone			
ORAL	Inadequate Analgesia		Acetate)	SS		ORAL
	Multiple Sclerosis					
	Scar					
	Scratch					
	Spinal Disorder					
	Thrombosis					
	Vaginal Neoplasm					
	Weight Decreased					

Date:01/21/05ISR Number: 4562254-0Report Type:Expedited (15-DaCompany Report #2004071171

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Drug Eruption	Health	Neurontin			
Initial or Prolonged		Professional	(Gabapentin)	PS		
(800 MG)			Dilantin Suspension			
			(Phenytoin Sodium)			
			(Phenytoin Sodium)	SS		

Date:01/21/05ISR Number: 4562301-6Report Type:Expedited (15-DaCompany Report #2005006149

Age:39 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Abnormal Behaviour	Consumer	Neurontin			
	Drug Abuser		(Gabapentin)	PS		ORAL
ORAL	Dysarthria					
	Inadequate Analgesia					
	Mental Disorder					

Mobility Decreased
 Mutism
 Oral Intake Reduced
 Pneumonia
 Staring

Date:01/21/05ISR Number: 4562881-0Report Type:Expedited (15-DaCompany Report #2005011644
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anaemia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other		Drug Interaction					
3.6 GRAM,		Fatigue	Professional				
ORAL : 3.2		Hyperhidrosis	Company				
GRAM		Liver Disorder	Representative				
(INTERVAL:		Pneumonia					
DAILY) ORAL		Renal Disorder		Esomeprazole	C		
		Stevens-Johnson Syndrome		Celecoxib	C		
				Moxifloxacin	C		
				Hydrochloride	C		
				Ferrous Sulfate	C		
				Supradyn (Ferrous			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Carbonate, Magnesium
 Phosphate Monobasic,
 Manganese Sulfate,
 Molybdenum C
 Herbal Preparation C
 Morphine Sulfate C

Date:01/21/05ISR Number: 4562958-XReport Type:Expedited (15-DaCompany Report #2005009946
 Age:89 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Macular Degeneration	Foreign Health Professional	Neurontin (Gabapentin)	PS		

Date:01/21/05ISR Number: 4562959-1Report Type:Expedited (15-DaCompany Report #2004119747
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1500 MG (500	Arthralgia Drug Ineffective	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other	MG, 3 IN 1	Insomnia	Professional				
D), ORAL		Nausea					
		Pain					
		Pain In Jaw					

Date:01/21/05ISR Number: 4563238-9Report Type:Expedited (15-DaCompany Report #2004065345
 Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	900 MG (3 IN	Depressed Level Of Consciousness	Foreign Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1 D), ORAL		Faecal Incontinence	Professional				
		Somnolence	Company	All Other			

Representative

Therapeutic Products

(All Other
 Therapeutic
 Products) C
 Perindopril
 (Perindopril) C
 Insulin (Insulin) C
 Panadeine Co
 (Codeine Phosphate,
 Paracetamol) C

Date:01/21/05ISR Number: 4563719-8Report Type:Expedited (15-DaCompany Report #2005004219

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Body Height Decreased Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG, ORAL		Drug Effect Decreased Nerve Injury Paraesthesia Pharmaceutical Product Complaint Pruritus		Gabapentin (Gabapentin) Levothyroxine Sodium (Levothyroxine Sodium) Losartan Potassium (Losartan Potassium)	SS C C		

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Freedom Of Information (FOI) Report

Famotidine	
(Famotidine)	C
Verapamil	
(Verapamil)	C
Trazodone	
(Trazodone)	C
Tizanidine	
(Tizanidine)	C
Cevimeline	
Hydrochloride	
(Cevimeline	
Hydrochloride)	C
All Other	
Therapeutic Products	
(All Other	
Therapeutic	
Products)	C
Erythromycin	
(Erythromycin)	C
Estrogens Conjugated	
(Estrogens	
Conjugated)	C
All Other	
Therapeutic Products	
(All Other	
Therapeutic	
Products)	C
Multivitamins With	
Minerals	
(Multivitamins With	
Minerals)	C
B-Komplex "Leciva"	
(Calcium	
Pantotrenate,	
Nicotinamide,	
Pyridoxine	C
Fish Oil (Fish Oil)	C
Magnesium	
(Magnesium)	C
Glucosamine	
(Glucosamine)	C
Calcium (Calcium)	C
Acetylsalicylic Acid	
(Acetylsalicylic	
Acid)	C
Paracetamol	
(Paracetamol)	C

Tears Naturale
(Dextran 70,
Hyppromellose) C
Carmellose Sodium
(Carmellose Sodium) C

Date:01/21/05ISR Number: 4563969-0Report Type:Expedited (15-DaCompany Report #2005011247
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability		Depression Dystonia	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
1200 MG (300 MG, 4 IN 1							
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Freedom Of Information (FOI) Report

D), ORAL

Losartan (Losartan)	C
Atorvastatin (Atorvastatin)	C
Acetylsalicylic Acid (Acetylsalicylic Acid)	C
Lercanidipine Hydrochloride (Lercanidipine Hydrochloride)	C
Paracetamol (Paracetamol)	C

Date:01/21/05ISR Number: 4563978-1Report Type:Expedited (15-DaCompany Report #2005011385
Age:22 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1600 MG (1600 Disability MG), ORAL	Condition Aggravated Convulsion Pharmaceutical Product Complaint	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL

Date:01/24/05ISR Number: 4559313-5Report Type:Expedited (15-DaCompany Report #FR-JNJFOC-20040800636
Age:51 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - INTRAVENOUS Initial or Prolonged OROPHARINGEAL	Acute Generalised Exanthematous Pustulosis Antinuclear Antibody Positive Aphthous Stomatitis Nausea	Health Professional	Caelyx Becilan Neurontin Neurontin Neurontin Taxotere	PS SS SS SS SS		

Skin Infection
Stevens-Johnson Syndrome
Vomiting

Radiation Therapy SS
Zophren SS

Date:01/24/05ISR Number: 4561149-6Report Type:Expedited (15-DaCompany Report #2005012558
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Disability Other 300 MG (100 MG, 3 IN 1 D), ORAL		Balance Disorder Somnolence	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
				Nifedipine (Nifedipine)	C		
				Omeprazole (Omeprazole)	C		
				Tolterodine L-Tartrate (Tolterodine L-Tartrate)	C		
				Atorvastatin (Atorvastatin)	C		
				Combivent (Ipratropium Bromide, Salbutamol Sulfate)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Tears Naturale
 (Dextran 70,
 Hypromellose) C
 Lactulose
 (Lactulose) C
 Symbicort Turbuhaler
 Draco (Budesonide
 , Formoterol
 Fumarate) C Draco
 Panadeine Co
 (Codeine Phosphate,
 Paracetamol) C

Date:01/24/05ISR Number: 4561343-4Report Type:Expedited (15-DaCompany Report #2004111293
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Movement Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (300		Palpitations					
MG, 1 IN 1		Somnolence					
D), ORAL		Speech Disorder					
ORAL				Zoloft (Sertraline)	SS		ORAL

Date:01/24/05ISR Number: 4561659-1Report Type:Expedited (15-DaCompany Report #2005014440
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Inadequate Analgesia	Consumer	Neurontin (Gabapentin)	PS		
		Neuropathy		Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	SS		
		Pain In Extremity		Oxycocet (Oxycodone Hydrochloride, Paracetamol)	SS		

Date:01/24/05ISR Number: 4562178-9Report Type:Expedited (15-DaCompany Report #2005011627
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Inadequate Analgesia Overdose	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG, ORAL		Treatment Noncompliance Weight Increased		Aii Other Non-Therapeutic Products Paracetamol (Paracetamol)	SS C		

Date:01/24/05ISR Number: 4562181-9Report Type:Expedited (15-DaCompany Report #2004064324
Age: Gender:Male I/FU:F

Outcome	PT
Death	Acrophobia
Other	Anger Anxiety Asthenia

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Dose	Duration	Completed Suicide Decreased Interest Delusion	Report Source	Product	Role	Manufacturer	Route
1200 MG (600 MG, 2 IN 1 D)		Disturbance In Attention Drug Ineffective Feeling Abnormal	Consumer	Neurontin (Gabapentin)	PS		
		Hallucination, Auditory Irritability Memory Impairment Mental Impairment Mood Swings Obsessive-Compulsive Disorder Ochlophobia Panic Attack Paranoia Restlessness Self-Medication Sleep Disorder Treatment Noncompliance		Paroxetine Hydrochloride (Paroxetine Hydrochloride) Risperidone (Risperidone) Hydroxyzine Embonate (Hydroxyzine Embonate) Venlafaxine Hydrochloride (Venlafaxine Hydrochloride) Quetiapine Fumarate (Quetiapine Fumarate) Famotidine (Famotidine) Narine Repetabs (Loratadine, Pseudoephedrine Sulfate) Lansoprazole (Lansoprazole) Clarithromycin (Clariythromycin)	SS C C C C		

Date:01/24/05ISR Number: 4562227-8Report Type:Expedited (15-DaCompany Report #2004088763

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		No Adverse Drug Effect	Consumer Health Professional	Neurontin (Gabapentin) All Other Therapeutic Products	PS		

(All Other
Therapeutic
Products)

SS

Date:01/24/05ISR Number: 4562243-6Report Type:Expedited (15-DaCompany Report #2004077254
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams Condition Aggravated	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
600 MG, ORAL		Hypoaesthesia Lyme Disease Multiple Sclerosis Pain	Professional	Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Montelukast Sodium (Montelukast Sodium) Beclomethasone Dipropionate	C C		

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(Beclomethasone
Dipropionate) C
Fluticasone
Propionate
(Fluticasone
Propionate) C
Orciprenaline
Sulfate
(Orciprenaline
Sulfate) C
Mometasone Furoate
(Mometasone Furoate) C
Salbutamol
(Salbutamol) C

Date:01/24/05ISR Number: 4562245-XReport Type:Expedited (15-DaCompany Report #2004096710
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin			
Disability		Depression		(Gabapentin)	PS		ORAL
Other		Discomfort					
ORAL		Enuresis		Phenytoin Sodium			
		Pollakiuria		(Phenytoin Sodium)	C		
		Pruritus		Phenobarbital			
		Urinary Incontinence		(Phenobarbital)	C		
				Valproate Semisodium			
				(Valproate Semisodium)	C		
				Escitalopram			
				(Escitalopram)	C		

Date:01/24/05ISR Number: 4562355-7Report Type:Expedited (15-DaCompany Report #2004089776
Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin			
Disability		Decreased Activity	Health	(Gabapentin)	PS		ORAL
Other		Diabetes Mellitus					
(300 MG, 1		Diabetic Neuropathy	Professional				
D), ORAL		Drug Ineffective		Gabapentin			

Euphoric Mood
Intermittent Claudication
Medication Error
Pain In Extremity

(Gabapentin) SS
Oxycodone
Hydrochloride
(Oxycodone
Hydrochloride) C

Date:01/24/05ISR Number: 4562670-7Report Type:Expedited (15-DaCompany Report #2005010105
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNKNOWN	900 MG	Agranulocytosis Neutropenic Sepsis (300	Foreign Health	Gabapentin (Gabapentin)	PS		
MG, 3 IN 1		Upper Respiratory Tract Infection	Professional	Lansoprazole (Lansoprazole)	C		
				Morphine Sulfate (Morphine Sulfate)	C		
				Senna (Senna)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/24/05ISR Number: 4562671-9Report Type:Expedited (15-DaCompany Report #2005011390
 Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Sedation Somnolence	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
400 MG (100 MG, 4 IN 1 D), ORAL				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL
240 MG (120 MG, 2 IN 1 DAY), ORAL				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
				Clavulin (Amoxicillin Trihydrate, Clavulanate Potassium)	C		
				Atorvastatin (Atorvastatin)	C		
				Esomeprazole (Esomeprazole)	C		
				Amitriptyline (Amitriptyline)	C		
				Diclofenac (Diclofenac)	C		
				Dexamethasone (Dexamethasone)	C		
				Atenolol (Atenolol)	C		

Date:01/24/05ISR Number: 4562680-XReport Type:Expedited (15-DaCompany Report #2005011611
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Condition Aggravated Hepatitis	Foreign Health	Neurontin (Gabapentin)			
1200 MG (400 MG, 3 IN 1 D)			Professional Company Representative		PS		

Date:01/25/05ISR Number: 4560171-3Report Type:Expedited (15-DaCompany Report #200412840FR
Age:51 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Abdominal Pain Antinuclear Antibody Positive Blood Potassium Decreased Blood Sodium Decreased Clostridial Infection Diarrhoea Enterobacter Infection Erythema Multiforme Haemoglobin Decreased Inflammation

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Muscle Spasms Red Blood Cell Count Decreased					
		Staphylococcal Infection Stevens-Johnson Syndrome		Taxotere	PS	Aventis Pharmaceuticals Inc.	
INTRAVENOUS							
		Systemic Lupus Erythematosis Toxic Skin Eruption		Zophren Neurontin Caelyx	SS SS SS		ORAL
INTRAVENOUS							
		White Blood Cell Count Decreased		Becilan	C		ORAL
Dose unit:							
units							

Date:01/25/05ISR Number: 4561792-4Report Type:Expedited (15-DaCompany Report #US012745
Age:29 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Aphasia	Health	Gabitril	PS		ORAL
4 MG BID ORAL							
		Dizziness	Professional	Celexa	SS		ORAL
Initial or Prolonged 40 UNITS QHS							
ORAL		Drug Interaction	Company				
		Dystonia	Representative	Neurontin	SS		ORAL
300 MG ORAL							
		Hallucination		Xanax	SS		ORAL
0.5 UNIT PRN							
ORAL		Hypoglycaemia					
		Oedema Peripheral		Klonopin	SS		ORAL
0.5 UNIT BID							
ORAL		Vision Blurred					
				Lamictal	SS		ORAL
150 UNIT QD							
ORAL							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abortion Spontaneous Drug Exposure During Pregnancy	Foreign Health Professional	Inderal (Propranolol Hydrochloride, Tablet)	PS		ORAL
40 MG		Pregnancy	Other	Clonazepam (Clonazepam)	SS		ORAL
250 MG				Gabapentin (Gabapentin)	SS		ORAL
800 MG				Sertraline (Sertraline)	SS		
100 MG 1X PER 1 DAY				Tolterodine (Tolterodine)	SS		ORAL
2 MG 1X PER 1 DAY				Zopiclone (Zopiclone)	SS		ORAL
1 MG 2X PER 1 DAY							

Outcome
 Life-Threatening
 Hospitalization -
 Initial or Prolonged
 Disability
 Other
 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
600MG MORNING		Disturbance In Attention		Neurontin 600 Mg	PS		
600MG NIGHT		Eye Rolling					
30 MG		Mental Disorder		Paxil 30 Mg	SS		
MORNINGS		Overdose					
		Pain					
		Renal Disorder					
		Suicidal Ideation					
		Suicide Attempt					
		Tinnitus					

Date:01/25/05ISR Number: 4570047-3Report Type:Direct Company Report #CTU 237823
Age:55 YR Gender:Male I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300 MG PO BID	2 MON	Aura Condition Aggravated		Gabapentin 300 Mg (Neurontic Generic)	PS		ORAL
	50MG PO Q HS	2 MON	Pharmaceutical Product Complaint		Primidone 50mg (Mysoline Generic)	SS		ORAL
			Simple Partial Seizures		Tegretol Depakote	C C		

Date:01/26/05ISR Number: 4564171-9Report Type:Expedited (15-DaCompany Report #2005013649
Age: Gender: I/FU:I

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	ORAL		Dysphagia Neoplasm Malignant	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
	ORAL			Professional	Tranexamic Acid (Tranexamic Acid)	SS		ORAL

Date:01/26/05ISR Number: 4564178-1Report Type:Expedited (15-DaCompany Report #2004CG02461

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Required	Cerebrovascular Accident	Foreign	Zomig	PS		
Intervention to	Facial Pain	Health	Neurontin	SS		
Prevent Permanent	Facial Palsy	Professional	Zoloft	SS		ORAL
1 DF QD PO						
Impairment/Damage	Migraine	Other	Melodia	SS		
			Ibuprofen	C		
			Lexomil	C		

Date:01/26/05ISR Number: 4564586-9Report Type:Expedited (15-DaCompany Report #2004091185

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Headache	Foreign	Neurontin			
Initial or Prolonged	Phlebothrombosis	Health	(Gabapentin)	PS		ORAL
600 MG (200						
MG, 3 IN 1	Retinal Vascular	Professional				
D), ORAL	Thrombosis	Company				
	Vitreous Haemorrhage	Representative	Madopar (Benserazide			
			Hydrochloride,			
			Levodopa)	C		
			Cyanocobalamin			
			(Cyanocobalamin)	C		

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Freedom Of Information (FOI) Report

Date:01/26/05ISR Number: 4564691-7Report Type:Expedited (15-DaCompany Report #2005013661
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other (1800 MG), ORAL		Dysaesthesia Hypoaesthesia Neuropathy Peripheral Paraesthesia	Foreign Health Professional	Gabapentin (Gabapentin) Salbutamol (Salbutamol) Bendroflumethiazide (Bendroflumethiazide) Valsartan (Valsartan) Tocopherol (Tocopherol)	PS C C C C		ORAL

Date:01/26/05ISR Number: 4565156-9Report Type:Expedited (15-DaCompany Report #2005013644
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening 900 MG (300 MG, 3 IN 1 D), ORAL		Hypoglycaemia Loss Of Consciousness Tremor	Foreign Health Professional	Gabapentin (Gabapentin) Omeprazole (Omeprazole)	PS SS		ORAL
20 MG (20 MG, 1 IN 1 D) ,ORAL				Polyspectran Os (Bacitracin, Hydrocortisone Insulin Human Injection, Isophane Insulin	C C C C		

Alfuzosin C
 Blood And Related C
 Products

Date:01/26/05ISR Number: 4565677-9Report Type:Expedited (15-DaCompany Report #2005013770
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG (300 Other MG 1 IN 1 D) ORAL	Abasia Blood Glucose Increased Colonic Obstruction Diabetes Mellitus Diabetes Mellitus Inadequate Control Diabetic Coma Drug Screen Positive Gastric Disorder Hepatitis A Hepatitis C Loss Of Consciousness	Consumer	Neurontin (Gabapentin) Quetiapine Fumarate(Quetiapine Fumarate) Trazodone (Trazodone) Vicodin (Hydrocodone Bitartrate, Paracetamol) Alprazolam(Alprazola m	PS C C C C C		ORAL

FDA - Adverse Event Reporting System (AERS)

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Date:01/26/05ISR Number: 4566510-1Report Type:Direct
Age:75 YR Gender:Female I/FU:I

Company Report #CTU 238025

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Dyspnoea		Neurontin	PS		

Date:01/26/05ISR Number: 4566546-0Report Type:Direct
Age:21 YR Gender:Female I/FU:I

Company Report #CTU 238188

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 400MG 1 TID		Condition Aggravated Convulsion		Gabapentin 400mg	PS		

Date:01/26/05ISR Number: 4566554-XReport Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 238196

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ONE PO QD [LIFETIME]		Hypersensitivity Pharmaceutical Product Complaint Rash		Neurontin 400 One Po Qd	PS		ORAL

Date:01/26/05ISR Number: 4566645-3Report Type:Expedited (15-DaCompany Report #2005014940
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (300 MG, 3 IN 1 D), ORAL		Abnormal Behaviour Cholelithiasis Dental Caries Drug Ineffective Flatulence	Consumer	Neurontin (Gabapentin) Vicodin (Hydrocodone	PS		ORAL

(7.5), ORAL	Headache	Bitartrate,		
	Nausea	Paracetamol)	SS	ORAL
	Pain	Lithium Carbonate		
	Skin Depigmentation	(Lithium Carbonate)	C	
	Treatment Noncompliance	Risperidone		
	Weight Decreased	(Risperidone)	C	
		Sertraline		
		Hydrochloride		
		(Sertraline		
		Hydrochloride)	C	

Date:01/26/05ISR Number: 4566688-XReport Type:Expedited (15-DaCompany Report #2005014990
Age: Gender:Male I/FU:I

Outcome	PT
Other	Arthritis
	Bronchial Infection
	Bronchitis Chronic
	Condition Aggravated
	Convulsion
	Drug Ineffective
	Hepatic Enzyme Increased
	Hepatitis A
	Hepatitis B
	Influenza
	Insomnia
	Irritability

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG (100 MG, 1 IN 1 D), ORAL		Lung Disorder Neuropathy Oedema Peripheral Pain In Extremity Pneumonia	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (300 MG, IN 1 D), ORAL				Dilantin Kapseals (Phenytoin Sodium)	SS		ORAL
				Water (Water)	SS		
				Carbamazepine (Carbamazepine)	SS		
				Amitriptyline (Amitriptyline)	C		
				Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride)	C		

Date:01/26/05ISR Number: 4566690-8Report Type:Expedited (15-DaCompany Report #2004117371
Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ORAL		Fibromyalgia Herpes Zoster	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Hot Flush Hyperhidrosis Insomnia Medication Error Nausea	Professional	Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
				Oxycodone (Oxycodone)	C		
				Amitriptyline (Amitriptyline)	C		

Date:01/26/05ISR Number: 4566693-3Report Type:Expedited (15-DaCompany Report #2005012866
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:01/26/05ISR Number: 4566694-5Report Type:Expedited (15-DaCompany Report #2004081555
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anxiety Drug Dependence	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Memory Impairment Somnolence	Professional				

Freedom Of Information (FOI) Report

Date:01/26/05ISR Number: 4566695-7Report Type:Expedited (15-DaCompany Report #2004009355

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG (600 Other MG, 3 IN 1 D), ORAL	Abdominal Injury Abdominal Pain Accident Back Disorder Back Pain Blood Cholesterol Increased Body Temperature	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
20 MG (20 MG, 1 IN 1 D),ORAL	Decreased Cardiac Disorder Chest Pain Condition Aggravated		Lipitor (Atorvastatin)	SS		ORAL
1000 MG (1 IN 2 D), ORAL	Contusion Coordination Abnormal Disorientation Drug Ineffective Drug Intolerance Erectile Dysfunction Facial Pain Feeling Abnormal Feeling Cold Gastrointestinal Disorder Gastrooesophageal Reflux Disease Gingival Pain Glossodynia Hostility Hypersomnia Hyporeflexia Impaired Healing Injury Irritability Limb Injury Medication Error		Lithium (Lithium) Naproxen (Naproxen) Amoxicillin (Amoxicillin) Rofecoxib (Rofecoxib) All Other Therapeutic Products (All Other Therapeutic Products) Levothyroxine Sodium (Levothyroxine Sodium) Vitamins (Vitamins) Diltiazem Hydrochloride (Diltiazem Hydrochloride)	SS SS SS SS SS C C C C		ORAL

Movement Disorder
Myalgia
Neck Pain
Oesophageal Spasm
Overdose
Pain In Jaw
Post Procedural Pain
Sedation
Sleep Disorder
Somnolence
Tenderness
Tooth Abscess
Tremor
Vertigo

Date:01/26/05ISR Number: 4569088-1Report Type:Direct
Age:55 YR Gender:Male I/FU:I

Company Report #CTU 238153

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Ineffective Pharmaceutical Product Complaint		Gabapentin	PS		

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Date:01/26/05ISR Number: 4570119-3Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 238016

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 600 MG TID	Coma		Neurontin	PS		
Initial or Prolonged	Flushing Foaming At Mouth Overdose					

Date:01/27/05ISR Number: 4562853-6Report Type:Expedited (15-DaCompany Report #PHBS2005CH00950
Age:70 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 100 mg, PRN	Abdominal Pain Lower Anaemia		Voltaren	PS	Novartis Sector: Pharma	
SUBCUTANEOUS	Haematoma		Fraxiparine	SS		
3 mg/day	Haemoglobin Decreased		Marcoumar	SS		
40 mg/day	International Normalised		Prednisone	SS		
0.125 mg/day	Ratio Decreased Prothrombin Time Prolonged		Neurontin Pantozol Digoxin	SS C C		
0.5 tab/day	Shock Surgery		Amiloride W/Hydrochlorothiazid e	C		
100 mg/day			Tramal	C		
600 IU/day			Miacalcic	C		

Date:01/27/05ISR Number: 4565242-3Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 238452

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

1	CAPSULE PO		Abdominal Pain Upper		Gabapentin 400mg	PS	Greenstone	ORAL
			Drug Ineffective					
	THREE TIMES A		Neck Pain					
DAY	2	WK	Pharmaceutical Product		Ranitidine	C		
			Complaint		Premarin	C		
					Synthroid	C		

Date:01/27/05ISR Number: 4567122-6Report Type:Expedited (15-DaCompany Report #S04-USA-05512-01
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Aphasia	Health	Celexa			
Other		Dizziness	Professional	(Citalopram			
		Drug Interaction		Hydrobromide)	PS		ORAL
40 MG QHS PO		Dystonia		Gabitril			
		Hallucination, Auditory		(Tiagabine			
		Hypoglycaemia		Hydrochloride)	SS		
4 MG BID		Oedema Peripheral		Neurontin			
		Vision Blurred		(Gabapentin)	SS		
1800 MG QD				Lamictal	SS		
100 MG QAM				Lamictal	SS		ORAL
50 MG QHS				Klonopin	SS		
0.5 MG BID				Xanax (Alprazolam)	C		
0.5 MG PRN							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/27/05ISR Number: 4567295-5Report Type:Expedited (15-DaCompany Report #2005014959

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Anxiety	Consumer	Neurontin			
Initial or Prolonged	Balance Disorder		(Gabapentin)	PS		
	Bipolar Disorder		Geodon			
	Blood Potassium Increased		(Ziprasidone)	SS		ORAL
120 MG ORAL	Blood Pressure Increased					
	Bronchitis		Gabapentin			
	Condition Aggravated		(Gabapentin)	SS		
	Crying		Bupirone			
	Depression		Hydrochloride			
	Difficulty In Walking		(Buspirone			
	Dizziness		Hydrochloride)	SS		
	Fatigue		Olanzapine			
	Muscle Spasms		(Olanzapine)	SS		
			All Other			
			Non-Therapeutic			
			Products	SS		
			Diphenhydramine			
			Hydrochloride	C		
			Clomipramine	C		
			Pantoprazole	C		
			Metoprolol			
			(Metoprolol)	C		
			Amlodipine Besilate			
			(Amlodipine			
			Besilate)	C		
			Valsartan			
			(Valsartan)	C		
			Pravastatin Sodium			
			(Pravastatin Sodium)	C		
			Triamcinolone			
			(Triamcinolone)	C		
			Ciclopirox Olamine			
			(Ciclopirox Olamine)	C		

Date:01/27/05ISR Number: 4567323-7Report Type:Expedited (15-DaCompany Report #2004099089
Age:74 YR Gender:Female I/FU:I

Outcome	PT
Other	Burning Sensation
	Chapped Lips
	Cheilitis
	Glossitis
	Glossodynia
	Hyperkeratosis
	Lip Exfoliation
	Oral Pain
	Pharyngolaryngeal Pain

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Stomatitis Throat Irritation Tongue Exfoliation	Report Source	Product	Role	Manufacturer	Route
500 MG (100 MG, 5 IN 1 D) ORAL			Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
				Metoprolol Succinate	C		
				Estrogens Conjugated	C		
				Levothyroxine Sodium	C		
				Hydrochlorothiazide	C		
				Warfarin Sodium	C		
				Diazepam	C		
				Olmесartan Medoxomil	C		

Date:01/27/05ISR Number: 4567432-2Report Type:Expedited (15-DaCompany Report #PTWYE359621JAN05
Age:35 YR Gender:Female I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
150 MG 1X PER 1 DAY		Dizziness Drug Withdrawal Syndrome Feeling Cold Feeling Hot Hyperhidrosis	Other	Efexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release, 0)	PS		ORAL
600 MG 1X PER 1 DAY		Malaise		Gabapentin (Gabapentin)	SS		ORAL
80 MG 1X PER 1 DAY				Propranolol Hydrochloride (Propranolol Hydrochloride)	SS		ORAL

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Appetite Disorder	Foreign	Neurontin			
(3 IN 1 D),		Asthma	Health	(Gabapentin)	PS		ORAL
ORAL		Blood Glucose Increased	Professional				
		Blood Pressure Increased		Prednisone(Prednison			
		Complex Regional Pain		e)	C		
		Syndrome		All Otehr			
		Condition Aggravated		Non-Therapeutic			
		Drug Ineffective		Products (All Other			
		Drug Withdrawal Syndrome		Non-Therapeutic			
		Fatigue		Products)	C		
		Fibromyalgia		All Otehr			
		Hernia Repair		Therapeutic Products			
		Incision Site		(All Other			
		Complication		Therapeutic	C		
		Muscle Atrophy		Amitriptyline			
		Nausea		Hydrochloride			
		Weight Increased		(Amitriptyline			
				Hydrochloride0	C		
				Cyclobenzaprine			
				Hydrochloride			
				(Cyclobenzaprine			

Freedom Of Information (FOI) Report

Hydrochloride) C
 Clonazepam
 (Clonazepam) C

Date:01/28/05ISR Number: 4565795-5Report Type:Expedited (15-DaCompany Report #2005006201
 Age:36 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Chills	Foreign	Aldactone			
Life-Threatening		Coagulation Factor V	Health	(Spironolactone)	PS		
Hospitalization -		Level Decreased	Professional	Neurontin			
Initial or Prolonged		Hepatic Encephalopathy		(Gabapentin)	SS		ORAL
1200 MG (400		Hepatic Failure					
MG, INTERVAL:		Platelet Count Decreased					
EVERY DAY)		Prothrombin Time					
ORAL		Prolonged		Zidovudine			
		Pyrexia		W/Lamivudine			
				(Lamivudine,			
				Zidovudine)	SS		ORAL
300 MG (150							
MG, INTERVAL:							
EVERY DAY),							
ORAL							
				Peginterferon			
				Alfa-2b			
				(Peginterferon			
				Alfa-2b)	SS		
SUBCUTANEOUS	SUBCUTANEOUS						
				Ribavirin			
				(Ribavirin)	SS		ORAL
1 GRAM (1 IN							
1 D), ORAL							
(2 IN 1 D),				Kaletra (Lopinavir,			
				Ritonavir)	SS		ORAL
ORAL							

Buprenorphine
 Hydrochloride
 (Buprenorphine
 Hydrochloride) C
 Minocycline
 (Minocycline) C
 Clarithromycin
 (Clarithromycin) C
 Pentamidine
 Isethionate
 (Pentamidine
 Isethionate) C

Date:01/28/05ISR Number: 4565803-1Report Type:Expedited (15-DaCompany Report #2004097842
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Body Height Below Normal Drug Exposure During Pregnancy Foetal Growth Retardation Late Developer Premature Baby Pyloric Stenosis Weight Gain Poor	Foreign Health Professional	Neurontin (Gabapentin) Phenobarbital (Phenobarbital) Clobazam (Clobazam)	PS SS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4565806-7Report Type:Expedited (15-DaCompany Report #2005017130

Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Cardiac Failure	Foreign	Neurontin			
Initial or Prolonged	Chronic Obstructive	Health	(Gabapentin)	PS		
	Pulmonary Disease	Professional				
	Pneumonia	Company				
	Respiratory Failure	Representative				

Date:01/28/05ISR Number: 4565922-XReport Type:Expedited (15-DaCompany Report #2005013692

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Abdominal Pain Lower	Foreign	Neurontin			
Initial or Prolonged	Anaemia	Health	(Gabapentin)	PS		ORAL
ORAL						
Other	Fall	Professional	Prednisone			
	Muscle Haemorrhage		(Prednisone)	SS		ORAL
ORAL						
	Platelet Function Test		Heparin-Fraction			
	Abnormal		Calcium Salt			
	Shock		(Heparin-Fraction			
			Calcium Salt)	SS		
SUBCUTANEOUS	SUBCUTANEOUS					
			Diclofenac Sodium			
			(Diclofenac Sodium)	SS		ORAL
ORAL						
			Phenprocoumon			
			(Phenprocoumon)	SS		ORAL
(3 MG) ORAL						
			Pantoprazole Sodium			
			(Pantoprazole			
			Sodium)	C		
			Digoxin (Digoxin)	C		
			Moduretic "Msd"	C		
			Tramadol			
			Hydrochloride			
			(Tramadol			
			Hydrochloride)	C		
			Calcitonin Salmon			
			(Calcitonin, Salmon)	C		

Date:01/28/05ISR Number: 4565975-9Report Type:Expedited (15-DaCompany Report #2005016474
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Condition Aggravated Epilepsy	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
2400 MG (2400 MG ORAL)			Professional				
			Other	Valproate Sodium (Valproate Sodium)	C		

Date:01/28/05ISR Number: 4568194-5Report Type:Expedited (15-DaCompany Report #PTWYE359621JAN05
Age:35 YR Gender:Female I/FU:F

Outcome	PT
Other	Dizziness Drug Withdrawal Syndrome Feeling Abnormal Feeling Cold Feeling Hot Hyperhidrosis

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Malaise
Treatment Noncompliance

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
150 MG 1 X PER 1 DAY, ORAL		Foreign Health Professional Other	Efexor Xr (Venlafaxine Hydrochloride, Capsule, Extended Release)	PS		ORAL
600 MG 1 X PER 1 DAY, ORAL			Gabapentin (Gabapentin)	SS		ORAL
80 MG 1 X PER 1 DAY, ORAL			Propranolol Hydrochloride (Propranolol Hydrochloride)	SS		ORAL

Date:01/28/05ISR Number: 4575334-0Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 238581

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Ineffective		Gabapentin	PS		

Date:01/28/05ISR Number: 4575375-3Report Type:Direct
Age:44 YR Gender:Male I/FU:I

Company Report #CTU 238574

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MGM 4X		Drug Ineffective		Neurontin	PS		

Rash Pruritic

DAILY

Date:01/28/05ISR Number: 4575383-2Report Type:Direct
Age:88 YR Gender:Female I/FU:I

Company Report #CTU 238569

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation		Neurontin	PS		
100 MG 1-BID		Neuropathy Pharmaceutical Product Complaint					

Date:01/28/05ISR Number: 4575500-4Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 238596

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG BID		Drug Hypersensitivity		Gabapentin 100mg	PS		
		Nausea Pharmaceutical Product Complaint Stomach Discomfort					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/28/05ISR Number: 4575522-3Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 238620

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 0.5 MG AT		Abnormal Dreams Bruxism		Lorazepam Unknown	0.5 Mg PS		ORAL
BEDTIME ORAL 300 MG TWICE PER DAY ORAL				Neurontin Pfizer	300 Mg SS	Pfizer	ORAL
				Acetaminophen Triamt/Hctz	C C		

Date:01/28/05ISR Number: 4575525-9Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 238589

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 100 MG TID PO 1 MON		Drug Ineffective Inadequate Analgesia Insomnia Pain Pharmaceutical Product Complaint		Gabapentin	PS		ORAL

Date:01/28/05ISR Number: 4575598-3Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 238646

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - SEE IMAGE Initial or Prolonged		Abdominal Discomfort Bowel Sounds Abnormal		Neurontin Nexium	PS C		

Other	Convulsion	Protonix	C
	Faeces Discoloured	Prevacid	C
	Haematemesis	Aciphex	C
	Hair Colour Changes	Ultram	C
	Medication Error	Mobic	C
	Melaena	Celebrex	C
	Muscle Twitching	Herbal Products	C
	Refusal Of Treatment By Patient	Omega 3 Products	C
	Stool Analysis Abnormal		
	Vomiting		

Date:01/28/05ISR Number: 4575698-8Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 238583

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Gabapentin			
		Drug Tolerance Decreased		(Neurontin)	PS		
300 MG TWO							
		Paraesthesia					
TID							
		Pharmaceutical Product Complaint					

Hospitalization -	Confusional State	Durogesic	PS
TRANSDERMAL			
Initial or Prolonged	Fall	Neurontin	SS
OROPHARINGEAL			
	Malaise	Actiskenan	SS
OROPHARINGEAL			
	Spinal Compression	Kardegic	C
UNKNOWN			
	Fracture	Burinex	C
UNKNOWN			
		Cordarone	C
UNKNOWN			

Date:01/31/05ISR Number: 4565640-8Report Type:Expedited (15-DaCompany Report #DE-JNJFOC-20050106982
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Drug Abuser		Tramal	PS		
OROPHARINGEAL						
Initial or Prolonged	Psychotic Disorder		Tramal	SS		
OROPHARINGEAL						
			Neurontin	SS		
OROPHARINGEAL						
			Neurontin	SS		
OROPHARINGEAL						

Date:01/31/05ISR Number: 4568222-7Report Type:Expedited (15-DaCompany Report #2005015729
Age:35 YR Gender:Female I/FU:I

Outcome	PT
Other	Drug Withdrawal Syndrome
	Feeling Cold
	Feeling Hot

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Hyperhidrosis Syncope	Report Source	Product	Role	Manufacturer	Route
600 MG (600 MG, 1 IN 1 D), ORAL			Foreign Health Professional	Gabapentin (Tablets) (Gabapentin)	PS		ORAL
150 MG (150 MG, 1 IN 1 D), ORAL				Venlafaxine (Venlafaxine)	SS		ORAL
80 MG (DAILY), ORAL				Propranolol (Propranolol)	SS		ORAL

Date:01/31/05ISR Number: 4568233-1Report Type:Expedited (15-DaCompany Report #2005016492
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNKNOWN	900 MG	Agranulocytosis Chills (300 Decreased Immune Responsiveness	Foreign Health Professional	Gabapentin (Gabapentin)	PS		
MG, 3 IN 1 D), UNKNOWN		Malaise Neutropenic Sepsis Upper Respiratory Tract Infection		Lansoprazole Morphine Sulfate	C C		

Date:01/31/05ISR Number: 4568386-5Report Type:Expedited (15-DaCompany Report #2005016760
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cardiac Arrest Sinus Tachycardia	Foreign Health Professional	Gabapentin (Gabapentin) Levetiracetam (Levetiracetam) Carbamazepine (Carbamazepine)	PS SS SS		

Date:01/31/05ISR Number: 4568389-0Report Type:Expedited (15-DaCompany Report #2005011611

Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Cytolytic Hepatitis	Foreign Health Professional	Neurontin (Gabapentin)	PS		
1200 MG (400 MG, 3 IN 1 D)		Hepatitis	Company Representative	Carbamazepine (Carbamazepine)	SS		ORAL
500 MG (2 IN 1 D), ORAL							

Date:01/31/05ISR Number: 4568474-3Report Type:Expedited (15-DaCompany Report #2004119079

Age: Gender:Male 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

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, 3 IN 1 D)		Renal Failure Acute	Health Professional	Neurontin (Gabapentin)	PS		
				Lisinopril (Lisinopril)	C		
				Atenolol (Atenolol)	C		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		
				Atorvastatin (Atorvastatin)	C		
				Hydrochlorothiazide (Hydrochlorothiazide)	C		

Date:01/31/05ISR Number: 4569772-XReport Type:Expedited (15-DaCompany Report #2004009355
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abdominal Injury
Initial or Prolonged	Abdominal Pain
Other	Accident
	Back Disorder
	Back Pain
	Blood Cholesterol Increased
	Chest Pain
	Contusion
	Coordination Abnormal
	Disorientation
	Drug Ineffective
	Drug Intolerance
	Erectile Dysfunction
	Facial Pain
	Feeling Cold
	Gastrooesophageal Reflux Disease

Gingival Disorder
Glossodynia
Hostility
Hypersomnia
Hyporeflexia
Impaired Healing
Injury
Irritability
Limb Injury
Medication Error
Mobility Decreased
Muscle Spasms
Myalgia
Neck Pain
Oesophageal Spasm
Pain In Extremity
Pain In Jaw
Sedation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Sleep Disorder Somnolence Tooth Abscess	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL		Treatment Noncompliance Tremor Vertigo	Professional				
				Lipitor (Atorvastatin)	SS		ORAL
20 MG (20 MG, 1 IN 1 D), ORAL							
				Lithium (Lithium) (Lithium)	SS		
				Naproxen (Naproxen) (Naproxen)	SS		
				Amoxicillin (Amoxicillin)	SS		ORAL
1000 MG (1 IN 2 D), ORAL							
				Rofecoxib (Rofecoxib)	SS		
				Levothyroxine (Levothyroxine Sodium)	C		
				Vitamins (Vitamins)	C		
				Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C		

Date:01/31/05ISR Number: 4570702-5Report Type:Expedited (15-DaCompany Report #2005016291
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Activities Of Daily	Consumer	Neurontin			

Living Impaired	(Gabapentin)	PS
Foot Fracture	Diazepam (Diazepam)	
Intervertebral Disc Degeneration	(Diazepam)	SS
Musculoskeletal Stiffness	Ibuprofen	
Pain	(Ibuprofen)	SS
Rheumatoid Arthritis	Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS
Sciatica	Panadeine Co (Codeine Phosphate, Paracetamol)	SS

Date:01/31/05ISR Number: 4575429-1Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 238665

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required		Gastrointestinal Pain		Gabapentin 300mg			
Intervention to 2700MG DAY		Gastrooesophageal Reflux		Teva	PS	Teva	ORAL
Prevent Permanent ORAL		Disease					
Impairment/Damage		Pharmaceutical Product Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:01/31/05ISR Number: 4601976-XReport Type:Periodic
Age:53 YR Gender:Female I/FU:I

Company Report #140604USA

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Pain Upper	Consumer	Gabapentin	PS		ORAL
400 MILLIGRAM		Anorexia					
BID ORAL		Haematochezia		Xanax Xr	C		
		Headache		Minopril	C		
		Nausea					

Date:01/31/05ISR Number: 4601979-5Report Type:Periodic
Age:47 YR Gender:Male I/FU:I

Company Report #140691USA

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Abnormal Behaviour	Consumer	Gabapentin	PS		ORAL
300 MILLIGRAM		Agitation					
Initial or Prolonged		Akinesia		Oxycodone	C		
TID ORAL		Aphasia		Lexapro	C		
		Dysarthria		Zanaflex	C		
		Dystonia		Oxycontin	C		
		Fall					
		Fatigue					

Date:02/01/05ISR Number: 4567161-5Report Type:Expedited (15-DaCompany Report #US012745
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Aphasia		Lamictal	PS	Glaxosmithkline	ORAL
Hospitalization -		Diplopia		Gabitril	SS		ORAL
4MG Twice per		Dizziness					
Initial or Prolonged	15 DAY						
day		Drug Interaction		Celexa	SS		ORAL
Other		Dystonia					
40UNIT At							
night							

300MG	Hallucination	Neurontin	SS	ORAL
Variable dose	Hypoglycaemia			
.5MG As	Oedema Peripheral	Xanax	SS	ORAL
required	Vision Blurred			
.5MG Twice	Visual Acuity Reduced	Klonopin	SS	ORAL
per day		Alcohol	C	
		Cocaine	C	

Date:02/01/05ISR Number: 4567451-6Report Type:Expedited (15-DaCompany Report #US-MERCK-0501USA04023
Age:55 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Balance Disorder
Initial or Prolonged	Chills
	Convulsion
	Diarrhoea
	Disorientation
	Dizziness
	Electroencephalogram
	Abnormal
	Fatigue
	Feeling Abnormal
	Hypotension
	Influenza Like Illness
	Lung Infiltration
	Myalgia

Other	Depression	Consumer	Neurontin	
1800 MG	Migraine		(Gabapentin)	PS
(DAILY)	Neuralgia			
	Nuclear Magnetic		Lansoprazole	
	Resonance Imaging Brain		(Lansoprazole)	C
	Abnormal			

Date:02/01/05ISR Number: 4569380-0Report Type:Expedited (15-DaCompany Report #2004076197
Age:60 YR Gender:Male I/FU:F

Outcome	PT
Other	Abdominal Pain
	Abdominal Rigidity
	Abnormal Dreams
	Amnesia
	Anxiety
	Arthritis
	Asthenia
	Back Pain
	Chest Discomfort
	Depressed Mood
	Depression

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness Drug Ineffective Electroencephalogram	Report Source				
300 MG (100 MG, 3 IN 1 D)		Abnormal Electromyogram Abnormal Fatigue Headache	Consumer	Lipitor (Atorvastatin) Neurontin (Gabapentin)	PS SS		
		Loss Of Consciousness Medication Error Migraine Musculoskeletal Stiffness Neck Pain Nerve Conduction Studies Abnormal Road Traffic Accident Sleep Disorder Tinnitus Transient Ischaemic Attack					

Date:02/01/05ISR Number: 4569506-9Report Type:Expedited (15-DaCompany Report #2004085994
Age:74 YR Gender:Male I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
200 MG (600 MG, 3 IN 1 D), ORAL		Blood Alkaline Phosphatase Increased Bronchiectasis Gamma-Glutamyltransferase Increased Pyrexia Respiratory Tract Infection	Foreign Health Professional Company Representative	Neurontin (Tablets) (Gabapentin) Carbamazepine (Carbamazepine) Clonazepam (Clonazepam) Tramadol Hydrochloride (Tramadol Hydrochloride) Lidocaine	PS SS SS SS		ORAL
INTRAVENOUS	INTRAVENOUS						

INTRA VENOUS INTRA VENOUS (Lidocaine) SS

Date:02/01/05ISR Number: 4569671-3Report Type:Expedited (15-DaCompany Report #2004097842
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged PLACENTAL		Drug Exposure During Pregnancy	Foreign Health	Neurontin (Gabapentin)	PS		
Other PLACENTAL		Growth Retardation Late Developer	Professional	Phenobarbital(Phenob arbital)	SS		
PLACENTAL		Premature Baby		Clobazam (Clobazam)	SS		
PLACENTAL		Pyloric Stenosis Small For Dates Baby					

Date:02/01/05ISR Number: 4575837-9Report Type:Direct Company Report #CTU 238764
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Personality Change		Neurontin 600mg	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/05ISR Number: 4570255-1Report Type:Expedited (15-DaCompany Report #2005017734
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State	Foreign	Gabapentin			
		Pruritus	Health	(Gabapentin)	PS		ORAL
300 MG (100			Professional				
MG, 3 IN 1							
D), ORAL							

Date:02/02/05ISR Number: 4570547-6Report Type:Expedited (15-DaCompany Report #RENA-10959
Age:57 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Hyperkeratosis	Foreign	Renagel	PS		ORAL
0.8 G TID PO							
Initial or Prolonged		Mobility Decreased	Health	Beloc Zok	SS		
		Morphoea	Professional	Neurontin	SS		
300 MG QD			Company	Phos-Ex	C		
			Representative	Einsalpha	C		
			Other	Vigantoleten	C		
				Restex	C		
				Rohypnol	C		
				Marcumar	C		

Date:02/02/05ISR Number: 4570550-6Report Type:Direct Company Report #CTU 239015
Age:28 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Abnormal Behaviour		Nuerontin 300mg	PS		ORAL
4 TIMES DAY							
		Confusional State					
1200MG ORAL							
		Crying		Zoloft 100mg	SS		ORAL
1 THEN 2 DAY							
		Depression					
200MG ORAL							

Drug Ineffective
 Feeling Abnormal
 Mood Swings
 Overdose
 Personality Change
 Suicidal Ideation

Date:02/02/05ISR Number: 4570629-9Report Type:Direct
 Age:48 YR Gender:Female I/FU:I

Company Report #CTU 238992

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Pain		Gabapentin Cap Mg Greenstone	300 PS	Greenstone	ORAL
TWO, 8 HOUR							
600	ORAL						

Date:02/02/05ISR Number: 4571137-1Report Type:Expedited (15-DaCompany Report #2004071270
 Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anorexia Fall	Foreign Health	Neurontin (Gabapentin)	PS		
900 MG (300		Insomnia	Professional				
MG, 3 IN 1		Nausea	Company				
D),		Vertigo Vomiting	Representative	Panadeine Co (Codeine Phosphate, Paracetamol) Morphine Sulfate	C		

Freedom Of Information (FOI) Report

(Morphine Sulfate) C
 Clonazepam
 (Clonazepam) C

Date:02/02/05ISR Number: 4571145-0Report Type:Expedited (15-DaCompany Report #2005019189
 Age:18 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 3600 MG, ORAL		Alanine Aminotransferase Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1200 MG (400 MG, 3 IN 1 D), ORAL		Aspartate Aminotransferase Increased	Professional Company Representative	Carbamazepine (Carbamazepine)	SS		ORAL
		Drug Hypersensitivity Drug Rash With Eosinophilia And Systemic Symptoms Face Oedema Hepatomegaly Leukocytosis Lymphadenopathy Lymphocyte Morphology Abnormal Lymphocyte Transformation Test Positive Oedema Peripheral Pyrexia Rash					

Date:02/02/05ISR Number: 4571232-7Report Type:Expedited (15-DaCompany Report #US012745
 Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 4 MG BID ORAL		Aphasia	Health	Gabitril	PS		ORAL
40 UNITS QHS		Diplopia	Professional	Celexa	SS		ORAL

ORAL	Dizziness	Company			
300 MG ORAL	Drug Interaction	Representative	Neurontin	SS	ORAL
0.5 UNIT PRN	Dystonia		Xanax	SS	ORAL
ORAL	Hallucination				
0.5 UNIT BID	Hypoglycaemia		Klonopin	SS	ORAL
ORAL	Oedema Peripheral				
150 MG QD	Vision Blurred		Lamictal	SS	ORAL
ORAL					

Date:02/02/05ISR Number: 4571300-XReport Type:Expedited (15-DaCompany Report #2004100399
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Depression	Consumer	Neurontin			
Initial or Prolonged	Mental Disorder		(Gabapentin)	PS		
Other	Pain		Risperidone			
ORAL	Pancreatitis		(Risperidone)	SS		ORAL
	Sexual Dysfunction		All Other			
	Weight Increased		Therapeutic Products			
			(All Other			
			Therapeutic			
			Products)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/05ISR Number: 4571817-8Report Type:Expedited (15-DaCompany Report #2005014990

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Bronchial Infection Bronchitis Chronic	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG (100 MG, 1 IN 1 D), ORAL		Convulsion Drug Ineffective					
300 MG (300 MG, 1 IN 1 D), ORAL		Gastrointestinal Disorder Haemorrhoids Hepatitis A Hepatitis B		Dilantin Kapseals (Phenytoin Sodium)	SS		ORAL
		Hypoaesthesia Influenza Insomnia Irritability Neuropathy Peripheral Oedema Peripheral Pain In Extremity Paraesthesia Peripheral Coldness Pneumonia Sleep Disorder		Water (Water) Carbamazepine (Carbamazepine) Amitriptyline (Amitriptyline) Diphenhydramine Hydrochloride (Diphenhydramine Hydrochloride)	SS SS C C		

Date:02/02/05ISR Number: 4578199-6Report Type:Direct

Age: Gender:Male I/FU:I

Company Report #CTU 238871

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other PO		Anger Anxiety		Neurontin Generic	PS		ORAL

Date:02/02/05ISR Number: 4578214-XReport Type:Direct

Age:65 YR Gender:Female I/FU:I

Company Report #CTU 238868

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache		Neurontin 300 Mg Tab	PS		
300 MG AT		Nausea					
BEDTIME							

Date:02/02/05ISR Number: 4578225-4Report Type:Direct Company Report #CTU 238865
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated		Gabapentin 600 Mg	PS		ORAL
2 PO TID		Convulsion Drug Ineffective Pharmaceutical Product Complaint					

Date:02/02/05ISR Number: 4578321-1Report Type:Direct Company Report #CTU 238860
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
100MG PRN		Drug Effect Decreased		Gabapentin 100mg	PS		
		Nausea Pharmaceutical Product Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/02/05ISR Number: 4578359-4Report Type:Direct
 Age:88 YR Gender:Female I/FU:I

Company Report #CTU 238891

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 100 MG PO Initial or Prolonged DAILY	6	MON	Depressed Level Of Consciousness	Gabapentin	PS		ORAL
DOSE UNKNOWN	6	MON	Dialysis	Carbamazepine	SS		
			Nephritis Interstitial	Lisinopril	C		
			Renal Failure Acute	Metoprolol	C		
				Plavix	C		
				Isosorbide	C		
				Lasix	C		

Date:02/03/05ISR Number: 4577968-6Report Type:Direct
 Age:66 YR Gender:Female I/FU:I

Company Report #CTU 239169

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Disability 3 X DAILY 300 MG			Asthenia Chest Pain Headache	Gabapentin 300 Mg Greenstone Mf	PS	Greenstone	
			Pharmaceutical Product Complaint Renal Pain				

Date:02/04/05ISR Number: 4573003-4Report Type:Expedited (15-DaCompany Report #2005021397
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged			Foreign Health Professional Company Representative	Neurontin (Gabapentin) Nsaid'S (Nsaid'S) Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	PS SS		
			Coma Drug Toxicity Overdose				

Fentanyl	
(Fentanyl)	SS
Oxazepam	
(Oxazepam)	SS
Paracetamol	
(Paracetamol)	SS
Sotalol	
Hydrochloride	
(Sotalol	
Hydrochloride)	C

Date:02/04/05ISR Number: 4573004-6Report Type:Expedited (15-DaCompany Report #2004116429
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Congenital Anomaly		Congenital Anomaly	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
600 MG (300		Drug Exposure During					
MG, 2 IN 1		Pregnancy	Professional				
D), ORAL		Hypermetropia					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/04/05ISR Number: 4573049-6Report Type:Expedited (15-DaCompany Report #8008648
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Cardiac Arrest	Foreign Health Professional Other	Keppra Carbamazepine Gabapentin	PS SS SS		

Date:02/04/05ISR Number: 4574015-7Report Type:Expedited (15-DaCompany Report #2004100270
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Delusion Illusion Impaired Driving Ability	Consumer	Neurontin (Gabapentin)	PS		

Date:02/04/05ISR Number: 4574021-2Report Type:Expedited (15-DaCompany Report #2004099657
 Age:90 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG, ORAL Disability ORAL		Abasia Blindness Contusion	Consumer	Neurontin (Gabapentin) Celebrex (Celecoxib)	PS SS		ORAL ORAL
Other		Delusion Dizziness Dysstasia Fall Feeling Abnormal Illusion Joint Injury Thinking Abnormal Vertigo		Quinine (Quinine) Centrum Silver (Ascorbic Acid, Calcium, Minerals Nos, Retinol, Tocopheryl Acetate, Calcium (Calcium) Ascorbic Acid (Ascorbic Acid)	C C C C		

Date:02/04/05ISR Number: 4574030-3Report Type:Expedited (15-DaCompany Report #2005019377
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypercalcaemia Renal Failure Acute	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG, ORAL			Company Representative	Vicodin (Hydrocodone Bitartrate, Paracetamol) Olanzapine/Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	SS SS		
25 MG							

Date:02/04/05ISR Number: 4574036-4Report Type:Expedited (15-DaCompany Report #2004074283
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged Other		Balance Disorder Deafness Pain Tinnitus	Consumer Health Professional	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/04/05ISR Number: 4574174-6Report Type:Expedited (15-DaCompany Report #2004019184

Age:31 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Bruxism Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (1 IN 1 D), ORAL		Diarrhoea					
50 MG (50 MG, 1 IN 1 D), ORAL		Drug Ineffective Feeling Abnormal Gastrointestinal Disorder		Zoloft (Sertraline)	SS		ORAL
		Herpes Virus Infection Migraine Pain In Jaw Sleep Disorder Tooth Fracture Vision Blurred Weight Decreased		Ibuprofen (Ibuprofen)	C		

Date:02/04/05ISR Number: 4574231-4Report Type:Expedited (15-DaCompany Report #2005020587

Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Alopecia Disease Recurrence	Health Professional	Neurontin (Gabapentin)	PS		ORAL
(300 MG), ORAL		Intervertebral Disc Protrusion Pain In Extremity Surgical Procedure Repeated		Naproxen (Naproxen) Vicodin (Hydrocodone Bitartrate, Paracetamol) Propacet (Dextropropoxyphene Napsilate, Paracetamol) Carisoprodol (Carisoprodol) Fexofenadine	C C C		

Hydrochloride
 (Fexofenadine
 Hydrochloride) C
 Rofecoxib
 (Rofecoxib) C

Date:02/04/05ISR Number: 4574342-3Report Type:Expedited (15-DaCompany Report #2005022076
 Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 10 MG (10 MG, Disability 1 IN 1 D), Other ORAL		Cellulitis Difficulty In Walking Fall Foot Fracture Hypoaesthesia Intervertebral Disc Protrusion Myalgia Neuropathy Wheelchair User	Health Professional	Lipitor (Atorvastatin) Neurontin (Gabapentin) Alendronate Sodium (Alendronate Sodium) Acetylsalicylic Acid (Acetylsalicylic Acid)	PS SS SS C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/04/05ISR Number: 4574455-6Report Type:Expedited (15-DaCompany Report #2004059295

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Consumer	Neurontin (Tablets)			
Other		Anxiety	Health	(Gabapentin)	PS		ORAL
1800 MG (1		Asthma	Professional				
D), ORAL		Balance Disorder		Zyrtec-D 12 Hour			
		Chronic Obstructive		(Pseudoephedrine,			
		Pulmonary Disease		Cetirizine)	SS		
2 IN 1 D		Condition Aggravated		Mirapex			
		Disturbance In Attention		(Pramipexole)			
		Dizziness		(Pramipexole)	SS		
2 IN 1 D		Multiple Allergies		Clonazepam			
		Sinus Disorder		(Clonazepam)	SS		
2 IN 1 D		Tooth Extraction		Mirtazapine			
				(Mirtazapine)	SS		
				Amitriptyline			
				(Amitriptyline)	SS		
				Montelukast Sodium			
				(Montelukast Sodium)	C		
				Combivent			
				(Ipratropium			
				Bromide, Salbutamol			
				Sulfate)	C		
				Mirtazapine			
				(Mirtazapine)	C		
				Esomeprazole			
				(Esomeprazole)	C		
				Combivent			
				(Ipratropium			
				Bromide, Salbutamol			
				Sulfate)	C		
				Salbutamol			
				(Salbutamol)	C		
				Calamine/Camphor/Dip			
				henhydramine			
				(Calamine, Camphor,			
				Diphenhydramine)	C		
				Bupropion			
				Hydrochloride			

(Bupropion
Hydrochloride) C

Date:02/04/05ISR Number: 4574872-4Report Type:Expedited (15-DaCompany Report #2005008280
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Lung Disorder	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL		Pain Pharmaceutical Product	Professional				
0.5 MG		Complaint Tremor		Xanax Tablet (Alprazolam)	SS		
2400 MG (600 MG, 4 IN 1 D)				Morphine (Morphine) Gabapentin (Gabapentin)	SS SS		
				Montelukast Sodium (Montelukast Sodium) Theophylline	C		

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(Theophylline)	C
Udramil	
(Trandolapril,	
Verapamil	
Hydrochloride)	C
Insulin Novolin	
70/30 (Insulin Human	
Semisynthetic,	
Insulin Isophane	
Human Semisynthetic)	C
Duloxetine	
Hydrochloride	
(Duloxetine	
Hydrochloride)	C
Orphenadrine Citrate	
(Orphenadrine	
Citrate)	C
Furosemide	
(Furosemide)	C
Magnesium Hydroxide	
(Magnesium	
Hydroxide)	C
Vicodin (Hydrocodone	
Bitartrate,	
Paracetamol)	C
Isosorbide	
Mononitrate	
(Isosorbide	
Mononitrate)	C
Lisinopril	
(Lisinopril)	C
Acetylsalicylic Acid	
(Acetylsalicylic	
Acid)	C

Date:02/04/05ISR Number: 4575818-5Report Type:Expedited (15-DaCompany Report #2005013644
 Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Coma	Foreign	Neurontin			
		Hypoglycaemia	Health	(Gabapentin)	PS		ORAL
900 MG (300		Loss Of Consciousness	Professional				
MG, 3 IN 1							

Tremor

D), ORAL

20 MG (20 MG,

1 IN 1 D),

ORAL

SUBCUTANEOUS

SUBCUTANEOUS

SUBCUTANEOUS

SUBCUTANEOUS

Omeprazole
(Omeprazole)

SS

ORAL

Insulin Human
Injection, Isophane
(Insulin Human
Injection, Isophane) SS

Insulin (Insulin) SS

Polyspectran Os
(Bacitracin,
Hydrocortisone
Acetate, Neomycin
Sulfate, Polymyxin B C
Alfuzosin
(Alfuzosin) C
Blood And Related
Products (Blood And

Freedom Of Information (FOI) Report

Related Products) C
 Prednisolone
 (Prednisolone) C

Date:02/04/05ISR Number: 4575885-9Report Type:Expedited (15-DaCompany Report #2005006212
 Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Amnesia Blood Thyroid Stimulating	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL		Hormone Increased Fatigue	Professional				
		High Density Lipoprotein Increased Hypercholesterolaemia Hypersomnia Hypertriglyceridaemia Low Density Lipoprotein Abnormal Weight Increased		Estradiol (Estradiol) Dydrogesterone (Dydrogesterone) Clonazepam (Clonazepam) Tramadol Hydrochloride (Tramadol Hydrochloride)	C C C C		

Date:02/04/05ISR Number: 4576069-0Report Type:Expedited (15-DaCompany Report #2004104710
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Grand Mal Convulsion Incontinence	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
100 MG (100 MG , 1 IN 1 D) , ORAL			Professional				
ORAL				Valproate Sodium (Valproate Sodium) Carbamazepine	SS		ORAL

800 MG(400

(Carbamazepine)

SS

ORAL

MG, 2 IN 1

D), ORAL

Date:02/04/05ISR Number: 4576156-7Report Type:Expedited (15-DaCompany Report #2004045858

Age:81 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (TID Disability INTERVAL: EVERY DAY)	Cervical Myelopathy Dizziness Drug Tolerance Decreased Feeling Drunk Medication Error Muscular Weakness Osteoarthritis Pain Paraesthesia Paralysis Somnolence Spinal Cord Compression Vertigo	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Tramadol/Acetaminophen (Paracetamol, Tramadol) Diosin (Dosin) Rofecoxib (Rofecoxib) Atorvastatin (Atorvastatin) Fluoxetine (Fluoxetine)	PS C C C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/04/05ISR Number: 4576188-9Report Type:Expedited (15-DaCompany Report #2005021394

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Faecal Incontinence	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL		Spinal Disorder	Professional				

Date:02/04/05ISR Number: 4576190-7Report Type:Expedited (15-DaCompany Report #2004056277

Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Abdominal Pain	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
100 MG (100		Abdominal Rigidity					
MG, 1 IN 1		Acute Generalised	Professional				
D), ORAL		Exanthematous Pustulosis					
		Antinuclear Antibody Positive		Pyridoxine Hydrochloride (Pyridoxine Hydrochloride)	SS		
750 MG (250		Aphthous Stomatitis					
MG, 3 IN 1		Blood Albumin Decreased					
D),		Blood Chloride Decreased					
		Blood Creatinine		Ondansetron Hydrochloride (Ondansetron Hydrochloride)	SS		
		Decreased					
		Blood Creatinine Increased					
		Blood Glucose Increased					
		Blood Potassium Decreased		Docetaxel (Docetaxel)	SS		
INTRAVENOUS	110 MG	Blood Urea Decreased					
MG, 1 IN 3		Catheter Related					
WK),		Infection					
INTRAVENOUS		Clostridial Infection					
		Dermatitis Bullous		Doxorubicin			

		Diarrhoea	Hydrochloride	
		Enterobacter Infection		
		General Physical Health	(Doxorubicin	
		Deterioration	Hydrochloride)	SS
INTRAVENOUS	70 MG (70 MG,	Nausea		
1 IN 3 WK),				
		Oedema Peripheral		
INTRAVENOUS		Pain In Extremity		
		Pco2 Decreased		
		Protein Total Decreased		
		Staphylococcal Infection		
		Stevens-Johnson Syndrome		
		Vomiting		

Date:02/07/05ISR Number: 4571666-0Report Type:Expedited (15-DaCompany Report #FRWYE676031MAR04
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disturbance In Attention		Enbrel	PS		
SUBCUTANEOUS							
		Fatigue		Enbrel	SS		
SUBCUTANEOUS	unknown						
		Insomnia		Skenan	SS		ORAL
		Keratitis Herpetic		Neurontin	SS		ORAL
unknown							
		Malaise		Morphine	SS		
		Nausea					
		Somnolence					
		Vomiting					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/07/05ISR Number: 4576407-9Report Type:Expedited (15-DaCompany Report #8006735

Age:7 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	500 MG 2/D PO	Abnormal Behaviour	Foreign	Keppra	PS		ORAL
Other	300 MG 2/D PO	Aggression	Health	Neurontin	SS		ORAL
		Condition Aggravated	Professional	Rivotril	C		
		Electroencephalogram	Company	Lamictal "Burroughs			
		Abnormal	Representative	Wellcome"	C		
		Encephalopathy	Other	.	C		
		Epilepsy		.	C		
		Regressive Behaviour		.	C		
				.	C		
				.	C		
				.	C		

Date:02/07/05ISR Number: 4576755-2Report Type:Expedited (15-DaCompany Report #2005019183

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300 MG (300	Abscess	Consumer	Neurontin	PS		ORAL
	MG, 1 IN 1	Immunosuppression		(Gabapentin)			
		Localised Infection					
				Prednisone Tablet	SS		
				(Prednisone)			

Date:02/07/05ISR Number: 4576793-XReport Type:Expedited (15-DaCompany Report #2005002328

Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	300 MG (100	Balance Disorder	Consumer	Neurontin	PS		
Other		Constipation	Health	(Gabapentin)			

Depression

Professional

Hallucination
Incoherent
Weight Decreased

Alprazolam	
(Alprazolam)	C
Analgesics	
(Analgesics)	C
All Other	
Therapeutic	
Prudcts (All Other	
Therapeutic	
Prudcts)	C
Lovastatin	
(Lovastatin)	C
Metoprolol Tartrate	
(Metoprolol	
Tartrate)	C
Levothyroxine Sodium	
(Levothyroxine	
Sodium)	C
Losartan Potassium	
(Losartan	
Postassium)	C
Torasemide	
(Torasemide)	C
''	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/07/05ISR Number: 4578509-XReport Type:Direct
Age:28 YR Gender:Male I/FU:I

Company Report #CTU 239386

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	300MG 4	Anhedonia Confusional State		Neurontin 300mg Pfizer	PS	Pfizer	ORAL
TIMES A DAY		Crying					
ORAL		Depression					
		Intentional Misuse Mood Swings		Zoloft 100mg Pfizer	SS	Pfizer	ORAL
100MG 2		Personality Change					
TIMES A DAY		Relationship Breakdown					
ORAL		Social Avoidant Behaviour Suicidal Ideation Thinking Abnormal					

Date:02/07/05ISR Number: 4578777-4Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 239388 E

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	30 MG DAILY	Pneumonia Aspiration		Cymbalta Lilly	PS	Lilly	ORAL
Life-Threatening	PO	Respiratory Failure					
Hospitalization -	100 MG TID	Vomiting		Neurontin	SS		ORAL
Initial or Prolonged	PO						

Date:02/07/05ISR Number: 4578839-1Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 239452

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective		Gabapentin 200 Mg Po			

1200 MG PO Pain Tid PS
Pharmaceutical Product
TID Complaint Cymbalta C

Date:02/07/05ISR Number: 4578845-7Report Type:Direct Company Report #CTU 239448
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Neurontin	PS		ORAL
300 MG PO QID		Headache Pharmaceutical Product Complaint					

Date:02/07/05ISR Number: 4687551-XReport Type:Direct Company Report #CTU 239524
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Nausea		Gabapentin 300mg	PS		
Other		Stomach Discomfort					
3 TABS 3 X		Therapeutic Response Unexpected With Drug Substitution					
DAILY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/08/05ISR Number: 4575021-9Report Type:Expedited (15-DaCompany Report #US-AMGEN-US109092

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bone Marrow Depression		Epogen	PS		
SUBCUTANEOUS		Malnutrition		Neurontin	SS		
		Therapeutic Response		Coumadin	C		
		Decreased		Colchicine	C		
		Thrombocytopenia		Amiodarone	C		
				Diltiazem	C		
				Hectoral	C		
				Coreg	C		

Date:02/08/05ISR Number: 4576937-XReport Type:Expedited (15-DaCompany Report #2005008280

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin			
1800 MG (600		Lung Disorder	Health	(Gabapentin)	PS		ORAL
MG, 3 IN 1		Pain	Professional				
MG, 3 IN 1		Pharmaceutical Product					
D), ORAL		Complaint		Xanax Tablet			
(0.5 MG)		Tremor		(Alprazolam)	SS		
				Morphine (Morphine)	SS		
				Gabapentin			
				(Gabapentin)	SS		
2400 MG (600							
MG, 4 IN 1 D)				Montelukast Sodium			
				(Montelukast Sodium)	C		
				Theophylline			
				(Theophylline)	C		
				Udramil			
				(Trandolapril,			
				Verapamil			
				Hydrochloride)	C		
				Insulin Novolin			

70/30 (Insulin Human Semisynthetic, Insulin Isophane Human Semisynthetic)	C
Duloxetine Hydrochloride (Duloxetine Hydrochloride)	C
Orphenadrine Citrate (Orphenadrine Citrate)	C
Furosemide (Furosemide)	C
Magnesium Hydroxide (Magnesium Hydroxide)	C
Vicodin (Hydrocodone Bitartrate, Paracetamol)	C
Iisosorbide Mononitrate (Isosorbide Mononitrate)	C
Lisinopril (Lisinopril)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Acetylsalicylic Acid
(Acetylsalicylic
Acid) C

Date:02/08/05ISR Number: 4576939-3Report Type:Expedited (15-DaCompany Report #2005008379

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Aneurysm	Health	Neurontin			
Other		Disease Progression	Professional	(Gabapentin)	PS		
100 MG (100							
MG, 1 IN 1D)		Fatigue					
		Feeling Abnormal		Oxycodone			
		Gastroenteritis Viral		Hydrochloride			
		Movement Disorder		(Oxycodone			
		Neuropathy		Hydrochloride)	C		
		Visual Acuity Reduced		Lorazepam			
				(Lorazepam)	C		
				Escitalopram			
				(Escitalopram)	C		
				Atenolol (Atenolol)	C		
				Rabeprazole Sodium			
				(Rabeprazole Sodium)	C		
				Thyroid (Thyroid)	C		
				Lidocaine			
				Hydrochloride			
				(Lidocaine			
				Hydrochloride)	C		

Date:02/08/05ISR Number: 4576943-5Report Type:Expedited (15-DaCompany Report #2004112247

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ankle Fracture	Consumer	Neurontin			
		Balance Disorder	Company	(Gabapentin)	PS		
900 MG (1 D)							
		Fall	Representative	Oxcarbazepine			
				(Oxcarbazepine)	SS		

Age:61 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Acute Respiratory
Initial or Prolonged	Distress Syndrome
Other	Blood Albumin Decreased
	Blood Alkaline
	Phosphatase Increased
	Blood Pressure Increased
	Crepitations
	Drug Interaction
	Ejection Fraction
	Decreased
	Electrocardiogram Qt
	Prolonged
	Haemoglobin Decreased
	Heart Rate Increased
	Mitral Valve Incompetence
	Pulmonary Oedema
	Respiratory Rate

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Increased Torsade De Pointes Ventricular Tachycardia White Blood Cell Count	Report Source	Product	Role	Manufacturer	Route
50 MG (1 IN 1 D), ORAL		Increased	Literature	Sertraline (Sertraline)	PS		ORAL
900 MG (3 IN 1 D)				Gabapentin (Gabapentin)	SS		
INTRAVENOUS	INTRAVNEOUS			Fentanyl (Fentanyl)	SS		
ORAL				Midazolam (Midazolam)	SS		
				Methadone (Methadone)	SS		ORAL
				Enalapril (Enalapril)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Digoxin (Digoxin)	C		

Date:02/08/05ISR Number: 4576953-8Report Type:Expedited (15-DaCompany Report #2004009355

Age: Gender:Male I/FU:F

Outcome PT
Hospitalization - Abdominal Pain
Initial or Prolonged Accident
Other Back Disorder
Blood Cholesterol
Increased
Chest Pain
Condition Aggravated
Contusion
Disorientation
Drug Effect Decreased
Erectile Dysfunction
Facial Pain
Feeling Cold

Gastrooesophageal Reflux
Disease
Gingival Pain
Glossodynia
Hostility
Hypersomnia
Hyporeflexia
Impaired Healing
Injury
Movement Disorder
Muscle Spasms
Myalgia
Oesophageal Spasm
Overdose
Pain In Extremity
Pain In Jaw
Sedation
Skeletal Injury
Sleep Disorder
Somnolence
Tenderness
Tooth Abscess

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Freedom Of Information (FOI) Report

Dose	Duration	Tremor Vertigo	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, 3 IN 1 D), ORAL			Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
20 MG (20 MG, 1 IN 1 D), ORAL				Lipitor (atorvastatin)	SS		ORAL
1000 MG (1 IN 2 D), ORAL				Lithium (Lithium) Naproxen (Naproxen) Amoxicillin (Amoxicillin)	SS SS SS		ORAL
				Rofecoxib (Rofecoxib) All Other Therapeutic Products(All Other Therapeutic Products0 Levothyroxine Sodium (Levothyroxine Sodium) Vitamins (Vitamins0 Diltiazem Hydrochloride (Diltiazem Hydrochloride)	SS C C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional Company Representative	Neurontin (Gabapentin) Vicodin (Hydrocodone Bitartrate, Paracetamol)	PS C		

Date:02/08/05ISR Number: 4577122-8Report Type:Expedited (15-DaCompany Report #2004071171
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (600 Other MG, 1 IN 1 D)		Drug Hypersensitivity Drug Interaction Pyrexia	Health Professional Company Representative	Neurontin (Gabapentin) Dilantin Suspension (Phenytoin Sodium) (Phenytoin Sodium) Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	PS SS C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/08/05ISR Number: 4577166-6Report Type:Expedited (15-DaCompany Report #2004065605

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Health Professional Company Representative	Neurontin (Gabapentin) Vicodin (Hydrocodone Bitartrate, Paracetamol)	PS SS		

Date:02/08/05ISR Number: 4577177-0Report Type:Expedited (15-DaCompany Report #2005023350

Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Dependence Pharmaceutical Product Complaint	Health Professional	Neurontin (Gabapentin) Dextropropoxyphene (Dextropropoxyphene)	PS C		

Date:02/08/05ISR Number: 4577474-9Report Type:Expedited (15-DaCompany Report #2005023348

Age:21 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Other	1500 MG (500 MG, 3 IN 1	Cough Retching	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL

D), ORAL

Valproate Sodium (Valproate Sodium)	C
Clobazam (Clobazam)	C
Anovlar (Ethinylestradiol, Norethisterone Acetate)	C
Zuclopenthixol (Zuclopenthixol)	C

Date:02/08/05ISR Number: 4579232-8Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 239761

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
600 MG PO TID		Drug Ineffective		Neurontin Generic	PS		ORAL
		Pain					
		Pharmaceutical Product					
		Complaint					

Date:02/09/05ISR Number: 4577755-9Report Type:Expedited (15-DaCompany Report #2004AL000570
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Cardiac Arrest	Literature	Kadian (Morphine			
		Cardio-Respiratory Arrest	Health	Sulfate Sustained			
		Completed Suicide	Professional	Release) Capsules,			
		Overdose		100 Mg (Alpharma)	PS	Alpharma	ORAL
PO		Pupil Fixed		Trazodone			
				Hydrochloride			
				Tablets, 100 Mg			
				(Purepac)	SS	Purepac	ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

PO			Benzodiazepine Derivatives	SS		ORAL
PO			Diazepam Tablets Ups, 10 Mg (Purepac)	SS	Purepac	ORAL
PO			Risperidone	SS		ORAL
PO			Levodopa	SS		ORAL
PO			Cyclobenzaprine	SS		ORAL
PO			Trihexyphenidyl	SS		ORAL
PO			Propoxyphene	SS		ORAL
PO			Venlafaxine	SS		ORAL
PO			Gabapentin	SS		ORAL

Date:02/09/05ISR Number: 4578460-5Report Type:Expedited (15-DaCompany Report #2005022624
 Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other	150 MG (150 MG, QD INTERVAL; EVERY DAY),	Back Pain Economic Problem Pain Rheumatoid Arthritis Sciatic Nerve Injury	Consumer	Norpace Capsule (Disopyramide Phosphate)	PS		ORAL
	ORAL			Neurontin (Gabapentin)	SS		ORAL
	1200 MG (400 MG, TID INTERVAL:						

EVERY DAY),

ORAL

Furosemide	
(Furosemide)	C
Metoprolol Succinate	
(Metoprolol	
Succinate)	C
Levothyroxine Sodium	
(Levothyroxine	
Sodium)	C
Potassium Chloride	
(Potassium Chloride)	C
Isosorbide	
Mononitrate	
(Isosorbide	
Mononitrate)	C
Esomeprazole	
(Esomeprazole)	C
Carisoprodol	
(Carisoprodol)	C
Glipizide	
(Glipizide)	C

Date:02/09/05ISR Number: 4578521-0Report Type:Expedited (15-DaCompany Report #2004033914

Age:28 YR Gender:Female I/FU:F

Outcome	PT
Life-Threatening	Anaemia
Hospitalization -	Epilepsy
Initial or Prolonged	Gastric Disorder
Other	Generalised Oedema
	Hepatic Enzyme 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
		Liver Disorder Pleural Effusion Pneumonia	Report Source				
2100 MG (D), ORAL		Pulmonary Oedema Tachycardia Tachypnoea Weight Decreased	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
1 IN 1 D, ORAL				Dalacin Ovulos (Clindamycin Hydrochloride) (Clindamycin Palmitate Udramil (Trandolapril, Verapamil Hydrochloride)	SS SS		ORAL
10 MG (10 MG, 1 IN 1 D), ORAL				Torasemide (Torasemide)	SS		ORAL
				Irbesartan (Irbesartan) Moxonidine (Moxonidine) Insulin (Insulin) Initard (Insulin, Insulin Injection, Isophane)	C C C C		

Date:02/09/05ISR Number: 4578778-6Report Type:Direct
Age:66 YR Gender:Male I/FU:I

Company Report #CTU 239966

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG 2		Drug Ineffective		Gabapentin	PS		

CAPSULES
Pharmaceutical Product
Complaint
Date:02/09/05ISR Number: 4578788-9Report Type:Direct Company Report #CTU 239970
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Tolerance Decreased Dyspepsia Gastrointestinal Disorder Pharmaceutical Product Complaint		Neurontin	PS		

Date:02/09/05ISR Number: 4578797-XReport Type:Direct Company Report #CTU 239987
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
200 MG AM		Pharmaceutical Product		Gabapentin	PS		
NOON 4P		Complaint					
1200MG AT HS							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/09/05ISR Number: 4578804-4Report Type:Direct
Age:54 YR Gender:Female I/FU:I

Company Report #CTU 239952

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective Pharmaceutical Product Complaint		Neurontin 600 Mg 1 Po Tid Generic By Purepac Pharm	PS	Purepac Pharm	ORAL
600 MG 1 PO							
TID							

Date:02/09/05ISR Number: 4578805-6Report Type:Direct
Age:22 YR Gender:Male I/FU:I

Company Report #CTU 239953

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated		Gabapentin 600 Mg	PS		
Other		Convulsion					
3 Q AM, 2 Q P							
3 Q HS		Pharmaceutical Product Complaint					

Date:02/09/05ISR Number: 4579262-6Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 239979

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dysgeusia Tongue Coated		Neurontin 300 Mg	PS		

Date:02/09/05ISR Number: 4579266-3Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 239936

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Headache		Neurontin	PS		
Other		Malaise					
300 MG TWO							
TID							

Vomiting

Date:02/09/05ISR Number: 4579311-5Report Type:Expedited (15-DaCompany Report #2004100399

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Hospitalization -	Anxiety	Consumer	Neurontin			
Initial or Prolonged	Depression		(Gabapentin)	PS		
Other	Memory Impairment		Risperidone			
ORAL	Mental Disorder		(Respiridone0	C		ORAL
	Nerve Injury		All Other			
	Pain		Therapeutic Products			
	Pancreatitis		(All Other			
	Sexual Dysfunction		Therapeutic			
	Weight Increased		Products0	C		

Date:02/09/05ISR Number: 4579319-XReport Type:Expedited (15-DaCompany Report #2004048907

Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abdominal Distension
Initial or Prolonged	Anxiety
Other	Arterial Injury
	Atrial Fibrillation
	Biliary Tract Disorder
	Blood Calcium Decreased

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Hydrochloride (Fluoxetine Hydrochloride)	C
Raloxifene Hydrochloride (Raloxifene Hydrochloride)	C
Lithium Carbonate (Lithium Carbonate)	C
Tramadol Hydrochloride (Tramadol Hydrochloride)	C
Atenolol (Atenolol)	C
Trazodone (Trazodone)	C
Propacet	

Freedom Of Information (FOI) Report

(Dextropropoxyphene Napsilate, Paracetamol)	C
Estratest Hs (Estrogens Esterified, Methyltestosterone)	C
Estradiol (Estradiol)	C
Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C
Nefazodone Hydrochloride (Nefazodone Hydrochloride)	C
Zolmitriptan (Zolmitriptan)	C
Sodium Fluoride (Sodium Fluoride)	C
Amoxicillin (Amoxicillin)	C
Meperidine W/Promethazine (Pethidine, Promethazine)	C
Azithromycin (Azithromycin)	C
Aquatab C (Dextromethorphan, Guaifenesin, Phenylpropanolamine)	C
Prednisone (Prednisone)	C
Pravastatin Sodium (Pravastatin Sodium)	C
Clindamycin (Clindamycin)	C
Chlorhexidine (Chlorhexidine)	C
Vicodin (Hydrocodone Bitartrate, Paracetamol)	C
Docusate Sodium (Docusate Sodium)	C
Risperidone (Risperidone)	C

Mirtazapine (Mirtazapine)	C
Bupropion Hydrochloride (Bupropion Hydrochloride)	C
Propranolol Hydrochloride (Propranolol Hydrochloride)	C
Sumatriptan Succinate (Sumatriptan Succinate)	C

Freedom Of Information (FOI) Report

Methocarbamol	
(Methocarbamol)	C
Naproxen (Naproxen)	C
Narine Repetabs	
(Loratadine,	
Pseudoephedrine	
Sulfate)	C
Temazepam	
(Temazepam)	C
Quetiapine Fumarate	
(Quetiapine	
Fumarate)	C
Robitussin A-C/Old	
Form/ (Codeine	
Phosphate,	
Guaifenesin,	
Pheniramine Maleate)	C
Flurazepam	
Hydrochloride	
(Flurazepam	
Hydrochloride)	C
Mometasone Furoate	
(Mometasone Furoate)	C
Adapalene	
(Adapalene)	C
Ronatic	
(Chlorphenamine	
Tannate, Mepyramine	
Tannate,	
Phenylephrine	C
Modafinil	
(Modafinil)	C
Tizanidine	
Hydrochloride	
(Tizanidine	
Hydrochloride)	C
Estrogens Conjugated	
(Estrogens	
Conjugated)	C
Estradiol	
(Estradiol)	C
Tiagabine	
Hydrochloride	
(Tiagabine	
Hydrochloride)	C
Sildenafil Citrate	
(Sildenafil Citrate)	C

Butorphanol Tartrate
(Butorphanol
Tartrate) C
Trimethobenzamide
Hydrochloride
(Trimethobenzamide
Hydrochloride) C
Lithium Carbonate
(Lithium Carbonate) C
Valproate Semisodium
(Valproate
Semisodium) C
Beclometasone
Dipropionate

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Beclometasone
Dipropionate) C
Clavulin
(Amoxicillin
Trihydrate,
Clavulanate
Potassium) C

Date:02/10/05ISR Number: 4577387-2Report Type:Expedited (15-DaCompany Report #DE-GLAXOSMITHKLINE-D0045025A
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Bladder Disorder		Lamictal	PS	Glaxosmithkline	ORAL
50MG Twice							
per day	5	MON					
		Blood Aldosterone					
800MG Three		Increased		Neurontin	SS		ORAL
times per day		Drug Interaction					
		Face Oedema					
		Residual Urine Volume					

Date:02/10/05ISR Number: 4579598-9Report Type:Direct Company Report #CTU 240048
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
800 MG TWO		Rash		Gabapentin 800 Mg	PS		
TID		Urticaria					

Date:02/10/05ISR Number: 4579670-3Report Type:Expedited (15-DaCompany Report #2004AL000000791
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Pacemaker Insertion Completed Suicide	Literature Health Professional	Verapamil Hydrochloride Tablets, 120 Mg			

PO	Hyperglycaemia	(Purepac)	PS	ORAL
	Hyperkalaemia	Metformin		
	Hypotension	Hydrochloride		
	Intentional Misuse	Tablets, 1000 Mg		
	Somnolence	(Purepac)	SS	ORAL
PO		Metoprolol	SS	ORAL
PO		Gabapentin	SS	
		Levothyroxine	SS	ORAL
PO				

Date:02/10/05ISR Number: 4579844-1Report Type:Direct Company Report #CTU 240101
 Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	3 X A DAY	Hepatic Cyst		Mobic 7.5 Mg	PS		ORAL
ORAL				Gabapentin 100 Mg	SS		ORAL
3X A DAY							
ORAL							

Date:02/10/05ISR Number: 4579871-4Report Type:Expedited (15-DaCompany Report #2004AL000701
 Age:46 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
		Areflexia Blood Pressure Diastolic Decreased Blood Pressure Systolic	Literature Health Professional	Trazodone Hydrochloride Tablets, 100 Mg (Purepac)	PS	Purepac	ORAL
PO		Increased Brain Oedema		Alprazolam Tablets Usp, 2 Mg (Purepac)	SS	Purepac	ORAL
PO		Coma Completed Suicide		Oxycodone (Long-Acting)	SS		ORAL
PO		General Physical Health		Olanzapine	SS		ORAL
PO		Deterioration		Cyclobenzaprine	SS		ORAL
PO		Heart Rate Irregular		Gabapentin	SS		ORAL
PO		Hypotension		Celecoxib	SS		ORAL
PO		Hypoxia Intentional Misuse Pupil Fixed Tachycardia					

Date:02/10/05ISR Number: 4579888-XReport Type:Expedited (15-DaCompany Report #2005023287

Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (600 Other MG, 3 IN 1 D), ORAL		Confusional State Creatinine Renal Clearance Decreased Fall	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
20 MG (5 MG, IN 1 D), ORAL		Lumbar Vertebra Injury Malaise		Morphine Sulfate (Morphine Sulfate)	SS		ORAL

(25 MCG,
EVERY 2
WEEKS)

Fentanyl (Fentanyl) SS
Acetylsalicylate
Lysine
(Acetylsalicylate
Lysine) C
Bumetanide
(Bumetanide) C
Amiodarone
(Amiodarone) C

Date:02/10/05ISR Number: 4579905-7Report Type:Expedited (15-DaCompany Report #K200500170
Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cardiac Disorder	Health	Altace Capsules			
Other		Diabetes Mellitus	Professional	(Ramipril) Capsule,			
		Myocardial Infarction	Other	2.5mg	PS		ORAL
2.5 MG, QD,		Nervous System Disorder					
ORAL		Tremor		Atenolol (Atenolol)			
				Tablet, 25mg	SS		ORAL
25 MG, QD,							
ORAL				Lipitor /Net			
				(Atorvastatin			
				Calcium) 80mg	SS		
80 MG, QD				Neurontin			
				(Gabapentin)			

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Freedom Of Information (FOI) Report

300 MG, TID, ORAL	Capsule, 300mg	SS	ORAL
10 MG BID, ORAL	Glipizide (Glipizide) Tablet, 10mg	SS	ORAL
30 MG, BID, ORAL	Prevacid (Lansoprazole), Capsule, 30mg	SS	ORAL
	Nitrotab (Glyceryl Trinitrate)	C	
	Plavix (Clopidogrel Sulfate)	C	

Date:02/10/05ISR Number: 4580034-7Report Type:Expedited (15-DaCompany Report #2004107080
Age:36 YR Gender:Male I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (300 MG, 1 IN 1 D), ORAL		Dizziness Dyspepsia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
400 MG (200 MG, 2 IN 1 D), ORAL		Hypersensitivity Pain	Professional	Celebrex (Celecoxib)	SS		ORAL
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged (10 MG, 1 IN Other 1 D)	Anxiety Blood Creatine Phosphokinase Increased Blood Testosterone Decreased Chest Pain Demyelination Depression Drug Toxicity	Consumer Health Professional	Lipitor (Atorvastatin)	PS		
0.5 MG	Feeling Abnormal Hypertension Hypoaesthesia Hypothyroidism Inadequate Analgesia Lethargy Muscle Tightness Muscle Twitching Muscular Weakness Myalgia Nervousness Pain In Extremity Sleep Apnoea Syndrome Thinking Abnormal		Neurontin (Gabapentin) Bextra (Valdecoxib) Levothyroxine Sodium (Levothyroxine Sodium) Perindopril Erbumine (Perindopril Erbumine) Hydrochlorothiazide (Hydrochlorothiazide) Topiramate (Topiramate) Naproxen Sodium (Naproxen Sodium) Acetylsalicylic Acid (Acetylsalicylic Acid) Levothyroxine Sodium (Levothyroxine Sodium)	SS SS SS SS SS C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/11/05ISR Number: 4582308-2Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 240225

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin (Generic)			
Other		Drug Hypersensitivity		#1	PS		
		Pharmaceutical Product					
300 MG TWO HS		Complaint		Neurontin (Generic)			
				#2	SS		
300 MG TWO HS							

Date:02/11/05ISR Number: 4582320-3Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 240223

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin (Generic)			
		Pharmaceutical Product			PS		
		Complaint					
		Urticaria					

Date:02/11/05ISR Number: 4583504-0Report Type:Expedited (15-DaCompany Report #2004114437
 Age:81 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin			
Other		Condition Aggravated	Consumer	(Gabapentin)	PS		
		Macular Degeneration					
1200 MG (1 IN		Neuropathic Pain					
1 D)		Pain		All Other			
		Treatment Noncompliance		Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	C		
				Dyazide			
				(Hydrochlorothiazide			
				, Triamterene)	C		

Outcome	PT
Hospitalization -	Activities Of Daily
Initial or Prolonged	Living Impaired
Disability	Aplastic Anaemia
Other	Arthralgia
	Asthenia
	Balance Disorder
	Burns Second Degree
	Coma
	Coordination Abnormal
	Crying
	Disturbance In Attention
	Dizziness
	Dry Mouth
	Fall
	Headache
	Hemiparesis
	Impaired Driving Ability
	Movement Disorder
	Muscle Spasms
	Nerve Injury
	Nervousness
	Pain

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Pain In Extremity Palpitations Panic Attack					
1200 MG		Paralysis Red Blood Cell Count Increased Speech Disorder	Consumer Health Professional	Neurontin (Gabapentin) Dilantin (Phenytoin Sodium)	PS SS		ORAL
(TID), ORAL		Walking Aid User Wheelchair User		Acetylsalicylic Acid (Acetylsalicylic Acid) Vitamins With Minerals (Vitamins With Minerals) Megestrol (Megestrol)	SS SS C		

Date:02/11/05ISR Number: 4583583-0Report Type:Expedited (15-DaCompany Report #2004051426
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dizziness Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG, ORAL		Dystonia Hallucination Hypoglycaemia		Xanax Tablet (Alprazolam) (Alprazolam)	SS		ORAL
ORAL		Oedema Peripheral Speech Disorder		Tiagabine Hydrochloride (Tiagabine Hydrochloride)	SS		ORAL
8 MG (4 MG, 2 IN 1 D), ORAL				Lamotrigine (Lamotrigine)	SS		ORAL
1 D, ORAL				Citalopram Hydrobromide (Citalopram			

1 IN 1 D,	Hydrobromide)	SS	ORAL
ORAL			
2 IN 1 D,	Clonazepam (Clonazepam)	SS	ORAL
ORAL			

Date:02/11/05ISR Number: 4583753-1Report Type:Expedited (15-DaCompany Report #2012272
Age:42 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Accidental Death	Consumer	Oxycontin Tablets			
Hospitalization -	Accidental Overdose	Health	(Oxycodone			
Initial or Prolonged	Asthenia	Professional	Hydrochloride)	PS		
SEE IMAGE						
Other	Back Pain	Other	Diazepam (Diazepam)	SS		
Required	Drug Abuser		Oxazepam (Oxazepam)	SS		
Intervention to	Erectile Dysfunction		Temazepam			
Prevent Permanent	Haematemesis		(Temazepam)	SS		
Impairment/Damage	Headache		Lorazepam			
	Iron Deficiency Anaemia		(Lorazepam)	SS		
	Mallory-Weiss Syndrome		Cannabnoids			
	Tachycardia		(Cannabis)	SS		
	Tooth Disorder		Diphenhydramine			
	Tooth Extraction		Hydrochloride			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Diphenhydramine Hydrochloride)	SS
Gabapentin (Gabapentin)	SS
Claritin (Loratadine)	C
Zoloft (Sertraline Hydrochloride)	C
Altace (Ramipril)	C
Allopurinol (Allopurinol)	C
Depakote (Valproate Semisodium)	C
Soma (Carisoprodol)	C
Relafen (Nabumetone)	C

Date:02/11/05ISR Number: 4583843-3Report Type:Expedited (15-DaCompany Report #2005022478
Age:73 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Glucose Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Drug Ineffective					
		Dyspnoea		Amitriptyline Hydrochloride			
		Inadequate Analgesia		(Amitriptyline Hydrochloride)	SS		
		Medical Device Complication		Glipizide	C		
		Post Procedural Complication		Amiodarone	C		
		Post Procedural Pain		Acetylsalicylic Acid	C		
		Rib Fracture		Citalopram Hydrobromide	C		
		Triple Vessel Bypass Graft		Simvastatin	C		
				Zolpidem Tartrate	C		
				Furosemide	C		
				Potassium	C		

Date:02/11/05ISR Number: 4583845-7Report Type:Expedited (15-DaCompany Report #2005022019
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Cystitis	Consumer	Neurontin			

Initial or Prolonged	Depression	(Gabapentin)	PS
600 MG (200			
Other	Diarrhoea		
MG, 3 IN 1 D)			
	Disturbance In Attention	Baclofen (Baclofen)	
	Drug Interaction	(Baclofen)	SS
130 MG			
	Hallucination	Paracetamol	C
	Hypotension	Ibuprofen	C
	Insomnia	Vicodin (Hydrocodone	
	Neuralgia	Bitartrate,	
	Pain In Extremity	Paracetamol)	C
	Psychotic Disorder		
	Sensory Disturbance		
	Social Avoidant Behaviour		
	Syncope		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/05ISR Number: 4579864-7Report Type:Expedited (15-DaCompany Report #200511219GDDC

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Rifampicin	PS	Aventis	
		Drug Interaction				Pharmaceuticals Inc.	

dose: UNKNOWN

DOSAGE

dose: UNKNOWN

DOSAGE

Date:02/14/05ISR Number: 4579993-8Report Type:Expedited (15-DaCompany Report #US-ROCHE-394642

Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Aphasia		Klonopin	PS	Roche	ORAL
Hospitalization -		Dizziness		Lamictal	I		ORAL
Initial or Prolonged		Drug Interaction		Neurontin	I		ORAL
		Dystonia		Xanax	I		ORAL
		Hallucination		Gabitril	I		ORAL
15 DAY							
		Hypoglycaemia		Celexa	I		ORAL
EVERY NIGHT.		Oedema Peripheral					

Date:02/14/05ISR Number: 4580323-6Report Type:Periodic

Company Report #US-GLAXOSMITHKLINE-A0432317A

Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
25MG Twice		Drug Exposure During		Lamictal	PS	Glaxosmithkline	ORAL
per day		Pregnancy					
150MG Unknown		Pruritus		Depakote Er	SS		ORAL
150MG Per day				Effexor	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/05ISR Number: 4584813-1Report Type:Expedited (15-DaCompany Report #2004067531

Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation	Consumer	Neurontin			
Hospitalization -		Anxiety	Health	(Gabapentin)	PS		ORAL
600 MG (300		Arthralgia	Professional				
Initial or Prolonged		Asthenia					
MG, 2 IN 1		Atherosclerosis		Benadryl			
Disability		Back Pain		(Diphenhydramine)	SS		ORAL
D), ORAL		Brain Oedema		Oxycodone			
Other		Burning Sensation		Hydrochloride			
ORAL		Cardio-Respiratory Arrest		(Oxycodone			
		Cardiomegaly		Hydrochloride)	SS		ORAL
1-3 TABLETS		Coma					
Q12H, ORAL		Coronary Artery		Hydrocodone			
		Atherosclerosis		(Hydrocodone)	SS		
1-2 EVERY 4-6		Drug Ineffective					
HOURS (10		Drug Interaction		Zolpidem Tartrate			
		Drug Level Increased		(Zolpidme Tartrate)	SS		ORAL
10 MG (10 MG,		Drug Screen Positive					
1 IN 1 D),		Erectile Dysfunction					
ORAL		Excoriation		Citalopram			
		Fall		Hydrobromide			
		General Physical Health		(Citalopram			
		Deterioration		Hydrobromide)	SS		ORAL
80 MG (80		Haematemesis					
MG,1 IN 1 D),		Hepatic Steatosis					
ORAL		Hypertensive Heart		Amitriptyline			
		Disease		Hydrochloride(
		Hypoaesthesia		Amitriptyline			
		Insomnia		Hydrochloride)	C		

Laceration	Fentanyl	C
Lumbar Radiculopathy	Cyclobenzaprine	
Major Depression	Hydrochloride	
Mood Swings	(Cyclobenzaprine	
Muscle Spasms	Hydrochloride)	C
Muscle Spasticity	Bupropion	
Muscle Tightness	Hydrochloride	
Musculoskeletal Pain	(Bupropion	
Osteoarthritis	Hydrochloride)	C
Pain	Quinine Sulfate	
Paraesthesia	(Quinine Sulfate)	C
Polysubstance Abuse	Sertraline	
Post Procedural Pain	Hydrochloride	
Pulmonary Oedema	(Sertraline	
Renal Cyst	Hydrochloride)	C
Respiratory Depression	Trazodone	
Snoring	(Trazodone)	C
Suicide Attempt	Hydroxyzine Embonate	
Tearfulness	(Hydroxyzine	
	Embonate)	C
	Carisoprodol	
	(Carisoprodol)	C
	Paroxetine	
	Hydrochloride	
	(Paroxetine	
	Hydrochloride)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/14/05ISR Number: 4584821-0Report Type:Expedited (15-DaCompany Report #2005023528
 Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	5 MG (5 MG, 1 IN 1 D), ORAL	Back Pain Blood Glucose Increased	Consumer	Glucotrol (Glipizide)	PS		ORAL
	100 MG (100 MG, 1 IN 1 D), ORAL	Cough Muscle Spasms Pain		Neurontin (Gabapentin)	SS		ORAL
		Treatment Noncompliance					
		Viral Upper Respiratory Tract Infection					

Date:02/14/05ISR Number: 4585337-8Report Type:Expedited (15-DaCompany Report #2005024395
 Age:25 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	1800 MG (600 MCG, 3 IN 1 D), ORAL	Disorientation Drug Interaction	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
		Hip Surgery					
		Nerve Injury					
		Suicidal Ideation		Glucosamine W/Chondroitin Sulfates (Ascorbic Acid, Chondroitin Sulfate, Glucosamine Ibuprofen Paracetamol Propacet (Dextropropoxyphene Napsylate, Paracetamol)	SS C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2100 MG Other 50 MG (50 MG, 11IN 1 D)		Contusion Crying Hemiparesis Neuropathy Paraesthesia Speech Disorder Tremor Visual Disturbance	Consumer	Neurontin (Gabapentin) Topiramate (Topiramate) Zolpidem Tartrate (Zolpidem Tartrate) Tramadol Hydrochloride (Tramadol Hydrochloride) Propacet (Dextropropoxyphene Napsilate, Paracetamol)	PS SS C C C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury Asphyxiation	Consumer	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Zoloft (Sertraline) SS

Date:02/14/05ISR Number: 4586718-9Report Type:Expedited (15-DaCompany Report #2005025147

Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1200 MG (1 Other D), ORAL	Anti-Hbc Antibody Positive	Foreign	Gabapentin (Tablets) (Gabapentin)	PS		ORAL
		Cholestasis					
		Prothrombin Time Prolonged Somnolence		Amitriptyline Hydrochloride (Amitriptyline Hydrochloride) Fentanyl (Fentanyl) Clonazepam (Clonazepam)	SS C C		

Date:02/14/05ISR Number: 4586757-8Report Type:Expedited (15-DaCompany Report #2005022778

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Completed Suicide Suicidal Ideation	Consumer	Zoloft (Sertraline) Neurontin (Gabapentin) Levothyroxine Sodium (Levothyroxine Sodium)	PS SS C		

Date:02/15/05ISR Number: 4584806-4Report Type:Expedited (15-DaCompany Report #2005AL000500

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other PO		Dyspnoea Erythema Rash	Health Professional	Gabapentin Tablets, 600 Mg (Purepac)	PS	Purepac	ORAL

Date:02/15/05ISR Number: 4587270-4Report Type:Expedited (15-DaCompany Report #2005024040

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury Asphyxiation	Consumer	Zoloft (Sertraline) Neurontin (Gabapentin)	PS SS		

Date:02/15/05ISR Number: 4587275-3Report Type:Expedited (15-DaCompany Report #2004030610

Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Myalgia Myopathy	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Oedema Peripheral Pain In Extremity	Health Professional Other	Diclofenac Sodium (Diclofenac Sodium) Venostasin Forte (Horse Chestnut Extract, Thiamine	C		

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Freedom Of Information (FOI) Report

Hydrochloride) C
 All Other
 Therapeutic Products C

Date:02/15/05ISR Number: 4587386-2Report Type:Expedited (15-DaCompany Report #2005027210
 Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Foreign	Neurontin			
		Hallucination	Health	(Gabapentin)	PS		ORAL
300 MG (300							
MG, 1 IN 1 D)		Sleep Disorder	Professional				
ORAL							
				Oxycodone			
				(Oxycodone)	SS		ORAL
80 MG (40 MG,							
2 IN 1 D)							
ORAL							

Date:02/15/05ISR Number: 4587499-5Report Type:Expedited (15-DaCompany Report #2005013649
 Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysphagia	Foreign	Gabapentin			
		Neoplasm Malignant	Health	(Gabapentin)	PS		ORAL
ORAL							
			Professional	Tranexamic Acid			
				(Tranexamic Acid)	SS		ORAL
ORAL							

Date:02/15/05ISR Number: 4587695-7Report Type:Expedited (15-DaCompany Report #2005026792
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin			

Exostosis (Gabapentin) PS ORAL
 100 MG (100
 MG, 1 IN 1
 D), ORAL

Lansoprazole
 (Lansoprazole) C
 Celecoxib
 (Celecoxib) C

Date:02/15/05ISR Number: 4588007-5Report Type:Expedited (15-DaCompany Report #2005014940
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abnormal Behaviour Cholelithiasis	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL		Dental Caries Drug Ineffective	Professional				
(6 IN 1 D), ORAL		Flatulence Headache Insomnia Skin Discolouration		Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		ORAL
		Weight Decreased		Lithium Carbonate (Lithium Carbonate) Risperidone (Risperidone) Sertraline Hydrochloride (Sertraline)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Hydrochloride) C
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C

Date:02/16/05ISR Number: 4583767-1Report Type:Expedited (15-DaCompany Report #200418611US
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Chest Pain Dizziness		Lantus	PS	Aventis Pharmaceuticals Inc.	
SUBCUTANEOUS	Dose unit:						

units

dose: 20/125

Glucotrol Xl	SS	ORAL
Metformin	SS	ORAL
Lotensin Hct	SS	ORAL
Plavix	SS	ORAL
Crestor	SS	ORAL
Celebrex	SS	ORAL
Ecotrin	SS	ORAL
Isosorbide	SS	ORAL
Neurontin	SS	ORAL
Ssri	SS	ORAL
Protonix	C	ORAL

Date:02/16/05ISR Number: 4586871-7Report Type:Direct Company Report #CTU 240572
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300MG PO BID		Diarrhoea Nausea		Generic Neurontin Gabapentin	PS		ORAL
		Vomiting		..	SS		

Date:02/16/05ISR Number: 4586882-1Report Type:Direct Company Report #CTU 240563
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Neuropathic Pain		Gabapentin	PS		
		Pain					
		Pharmaceutical Product					
		Complaint					

Date:02/16/05ISR Number: 4588441-3Report Type:Direct Company Report #CTU 240528
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Gabapentin 300 Mg			
		Pharmaceutical Product		Tab	PS		
12 BID		Complaint		Gabapentin 300 Mg			
				Caps	SS		
12 BID							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/16/05ISR Number: 4588782-XReport Type:Expedited (15-DaCompany Report #2005026789
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2400 MG (1 D), ORAL		Abasia Back Pain Balance Disorder	Consumer	Gabapentin (Gabapentin)	PS		ORAL
ORAL		Choking Cognitive Disorder Drug Ineffective Dystonia		Nefazodone Hydroocxhloride (Nefazodone Hydroocxhloride)	SS		ORAL
		Feeling Abnormal Headache Loss Of Consciousness Nuclear Magnetic Resonance Imaging Brain Abnormal Pain Pharyngolaryngeal Pain Road Traffic Accident Somnolence Speech Disorder Tremor Vision Blurred Walking Aid User		Escitalopram (Escitalopram) Clonazepam (Clonazepam)	SS C		

Date:02/16/05ISR Number: 4588894-0Report Type:Expedited (15-DaCompany Report #2005025068
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Balance Disorder Drug Ineffective	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Fall Pain		Carbamazepine (Carbamazepine)	SS		ORAL
ORAL		Walking Aid User		Lamotrigine (Lamotrigine)	SS		ORAL
				Hydrochlorothiazide			

(Hydrochlorothiazide) C
Atorvastatin (Atorvastatin) C
Amlodipine Besilate (Amlodipine Besilate) C
Atenolol (Atenolol) C
Lisinopril (Lisinopril) C

Date:02/16/05ISR Number: 4588900-3Report Type:Expedited (15-DaCompany Report #2005020356
Age:61 YR Gender:Female I/FU:F

Outcome PT
Hospitalization - Blood Albumin Decreased
Initial or Prolonged Blood Alkaline
Other Phosphatase Increased
Blood Pressure Increased
Drug Interaction
Ejection Fraction
Decreased
Electrocardiogram Qt

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Prolonged Haemoglobin Decreased Heart Rate Increased Hypotension	Report Source				
50 MG (1 IN 1 D), ORAL		Pulmonary Oedema Respiratory Distress Respiratory Rate	Literature	Sertraline (Sertraline)	PS		ORAL
900 MG (3 IN 1 D)		Increased Torsade De Pointes Ventricular Dysfunction		Gabapentin (Gabapentin)	SS		
ORAL		Ventricular Tachycardia White Blood Cell Count Increased		Methadone Hydrochloride (Methadone) (Methadone Hydrochloride)	SS		ORAL
INTRAVENOUS	INTRAVENOUS			Fentanyl (Fentanyl)	SS		
				Midazolam (Midazolam)	SS		
				Enalapril (Enalapril)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Digoxin (Digoxin)	C		

Date:02/16/05ISR Number: 4588914-3Report Type:Expedited (15-DaCompany Report #2004056018

Age:41 YR Gender:Female I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (1 D)		Abnormal Behaviour Amnesia	Consumer Health	Neurontin (Gabapentin)	PS		
INTRATHECAL	9 MG (1 D),	Dysstasia Feeling Hot	Professional	Morphine (Morphine)	SS		
INTRATHECAL		Granuloma Heart Rate Increased		Baclofen (Baclofen) All Other	SS		

Hot Flush
 Hyperhidrosis
 Insomnia
 Loss Of Consciousness
 Medical Device
 Complication
 Mental Disorder
 Nerve Injury
 Oedema
 Pain
 Speech Disorder
 Unevaluable Event
 Weight Increased

Non-Therapeutic
 Products (All Other
 Non-Therapeutic
 Product) SS
 Bupivacaine
 (Bupivacaine) C

Date:02/16/05ISR Number: 4589231-8Report Type:Direct
 Age:35 YR Gender:Female I/FU:I

Company Report #CTU 240556

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Gastric Disorder		Neurontin	PS		ORAL
400 MG, 3							
TABLETS TID,							
PO							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/16/05ISR Number: 4589284-7Report Type:Expedited (15-DaCompany Report #USA-2005-0018816

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Tolerance Increased	Consumer Company Representative	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		
	80 MG, TID			Lorcet (Paracetamol, Hydrocodone Bitartrate)	SS		
	400 MG, TID			Neurontin (Gabapentin)	SS		
	7.5 MG, BID			Soma (Carisoprodol)	SS		
	300 MG, HS			Tranxene (Clorazepate Dipotassium)	SS		
				Trazodone(Trazadone)	SS		

Date:02/16/05ISR Number: 4589410-XReport Type:Expedited (15-DaCompany Report #2005026776

Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Arm Amputation Hypermetropia	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
	2400 MG (800 MG, 3 IN 1 D), ORAL	Macular Degeneration	Professional				
		Macular Oedema	Company				
		Refusal Of Treatment By Patient	Representative	Fentanyl (Fentanyl)	C		
		Retinopathy		Topiramate (Topiramate)	C		
		Road Traffic Accident		Sertraline Hydrochloride (Sertraline Hydrochloride)	C		
				Clorazepate Dipotassium			

(Clorazepate
 Dipotassium) C
 Diazepam (Diazepam) C
 Clonazepam
 (Clonazepam) C

Date:02/16/05ISR Number: 4589512-8Report Type:Expedited (15-DaCompany Report #2005025718
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Amnesia Convulsion Rash Generalised Swelling Face Treatment Noncompliance	Consumer	Phenytoin Suspension (Phenytoin Sodium) Neurontin Phn Capsules (Gabapentin)	PS SS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL				Phenobarbital (Phenobarbital)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/16/05ISR Number: 4589543-8Report Type:Expedited (15-DaCompany Report #2005026800

Age:84 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 800 MG (1 D), ORAL		Condition Aggravated Oliguria Thrombocytopenia	Foreign Health Professional Other	Neurontin (Gabapentin) Danaparoid Sodium (Danaparoid Sodium)	PS SS		ORAL
	2250 UNITS (1 D), SUBCUTANEOUS						
	3000 MG (1 D), ORAL			Pristinamycin (Pristinamycin)	SS		ORAL
	200 MG (200 MG, 1 IN 1 D), ORAL			Amiodarone Hydrochloride (Amiodarone Hydrochloride)	SS		ORAL
				Acetylsalicylate Lysine (Acetylsalicylate Lysine) Heparin Calcium (Heparin Calcium)	C C		

Date:02/16/05ISR Number: 4589583-9Report Type:Expedited (15-DaCompany Report #2005027238

Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Confusional State	Foreign	Neurontin			

Initial or Prolonged ORAL	General Physical Health	Health	(Gabapentin)	PS	ORAL
	Deterioration Parkinson'S Disease	Professional	Analgesics (Analgesics)	C	

Date:02/16/05ISR Number: 4600120-2Report Type:Direct Company Report #CTU 240556
 Age:35 YR Gender:Female I/FU:I

Outcome Dose 400 MG THREE TID PO	Duration	PT Pharmaceutical Product Complaint	Report Source	Product Neurontin	Role PS	Manufacturer	Route ORAL
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Date:02/17/05ISR Number: 4586488-4Report Type:Direct Company Report #CTU 240663
 Age: Gender:Male I/FU:I

Outcome Dose Other 800 MG 2 PO Q AM 1 PO NOON 2 PO Q HS	Duration	PT Rash Therapeutic Product Ineffective	Report Source	Product Gabapentin	Role PS	Manufacturer	Route ORAL
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Date:02/17/05ISR Number: 4586506-3Report Type:Direct Company Report #CTU 240660
 Age: Gender:Female I/FU:I

Outcome Other	PT Pharmaceutical Product Complaint Vaginal Candidiasis
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vulvovaginal Discomfort

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG CAP			Gabapentin	PS		

Date:02/17/05ISR Number: 4586685-8Report Type:Direct
 Age: Gender:Male I/FU:I Company Report #CTU 240645

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin 1200 Mg Po	PS		ORAL
1200 MG PO		Pain		Tid			
TID		Pharmaceutical Product					
		Complaint					
		Therapy Non-Responder		Cymbalta	C		

Date:02/17/05ISR Number: 4586728-1Report Type:Direct
 Age:67 YR Gender:Male I/FU:I Company Report #CTU 240633

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Gabapentin 300 Mg Po	PS		ORAL
300 MG PO Q		Abdominal Discomfort		Q Hs			
HS		Nausea					

Date:02/17/05ISR Number: 4586732-3Report Type:Direct
 Age:41 YR Gender:Female I/FU:I Company Report #CTU 240636

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Gabapentin	PS		
Other		Aggression					
		Anxiety					
		Discomfort					

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Dilantin	PS		
100 MG 3 TID		Drug Tolerance Decreased					
1 WKS		Pharmaceutical Product		Neurontin	SS		
400 MG 3 TID		Complaint					

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Pressure Diastolic Decreased	Foreign Health Professional	Neurontin (Gabapentin)	PS		
30 MG		Oedema Mouth					
		Oropharyngeal Swelling Swelling Face	Company Representative Other	Panadeine Co (Codeine Phosphate, Paracetamol)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/17/05ISR Number: 4590182-3Report Type:Expedited (15-DaCompany Report #2004048907

Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2000 MG (400 Other MG, 5 IN 1 D)	Abdominal Distension Abdominal Injury Abnormal Behaviour Agitation Anger Anxiety Arterial Injury Atrial Fibrillation Biliary Tract Disorder Blood Calcium Decreased Body Dysmorphic Disorder Chest Pain Cough Depression Diaphragmatic Injury Feeling Guilty Gun Shot Wound Haematocrit Decreased Haemoglobin Decreased Haemothorax Hepatic Trauma Injury Insomnia Intercostal Neuralgia Nephrogenic Diabetes Insipidus Pain Pericardial Effusion Pleural Effusion Pleuritic Pain Pneumatoxis Cystoides Intestinalis Pneumonia Pneumothorax Psychiatric Symptom Pulmonary Embolism Respiratory Failure Scar Self Esteem Decreased Suicidal Ideation	Consumer	Neurontin (Gabapentin) Cefalexin (Cefalexin) Warfarin Sodium (Warfarin Sodium) Phenylpropanolamine (Phenylpropanolamine) Cyproheptadine Hydrochloride (Cyproheptadine Hydrochloride) Yohimbine (Yohimbine) Cetirizine Hydrochloride (Cetirizine Hydrochloride) Adapalene (Adapalene) Meloxicam (Meloxicam) Aripiprazole (Aripiprazole) Ketorolac Tromethamine (Ketorolac Tromethamine) Isocom (Dichloralphenazone, Isometheptene Mucate, Paracetamol) Clonazepam (Clonazepam) Olanzapine (Olanzapine) Zolpidem Tartrate	PS C C C C C C C C C C C C C C		

Suicide Attempt
Traumatic Shock
Venous Injury

(Zolpidem Tartrate)	C
Fluoxetine	
Hydrochloride	
(Fluoxetine	
Hydrochloride)	C
Raloxifene	
Hydrochloride	
(Raloxifene	
Hydrochloride)	C
Lithium Carbonate	
(Lithium	
Carbonate)	C
Tramadol	
Hydrochloride	
(Tramadol	
Hydrochloride)	C
Atenolol	
(Atenolol)	C

Freedom Of Information (FOI) Report

Trazodone	
(Trazodone)	C
Propacet	
(Dextropropoxyphene	
Napsilate,	
Paracetamol)	C
Estratest Hs	
(Estrogens	
Esterified,	
Methyltestosterone)	C
Estradiol	
(Estradiol)	C
Venlafaxine	
Hydrochloride	
(Venlafaxine	
Hydrochloride)	C
Nefazodone	
Hydrochloride	
(Nefazodone	
Hydrochloride)	C
Zolmitriptan	
(Zolmitriptan)	C
Sodium Fluoride	
(Sodium Fluoride)	C
Amoxicillin	
(Amoxicillin)	C
Meperidine	
W/Promethazine	
(Pethidine,	
Promethazine)	C
Azithromycin	
(Azithromycin)	C
Aquatab C	
(Dextromethorphan,	
Guaifenesin,	
Phenylpropanolamine)	C
Prednisone	
(Prednisone)	C
Pravastatin Sodium	
(Pravastatin Sodium)	C
Clindamycin	
(Clindamycin)	C
Chlorhexidine	
(Chlorhexidine)	C
Vicodin (Hydrocodone	
Bitartrate,	
Paracetamol)	C

Docusate Sodium	C
(Docusate Sodium)	
Risperidone	
(Risperidone)	C
Mirtazapine	
(Mirtazapine)	C
Bupropion	
Hydrochloride	
(Bupropion	
Hydrochloride)	C
Propranolol	
Hydrochloride	
(Propranolol	
Hydrochloride)	C

Freedom Of Information (FOI) Report

Sumatriptan Succinate (Sumatriptan Succinate)	C
Methocarbamol (Methocarbamol)	C
Naproxen (Naproxen)	C
Narine Repetabs (Loratadine, Pseudoephedrine Sulfate)	C
Temazepam (Temazepam)	C
Quetiapine Fumarate (Quetiapine Fumarate)	C
Robitussin A-C/Old Form/ (Codeine Phosphate, Guaifenesin, Pheniramine Maleate)	C
Flurazepam Hydrochloride (Flurazepam Hydrochloride)	C
Mometasone Furoate (Mometasone Furoate)	C
Adapalene (Adapalene)	C
Ronatic (Chlorphenamine Tannate, Mepyramine Tannate, Phenylephrine)	C
Modafinil (Modafinil)	C
Tizanidine Hydrochloride (Tizanidine Hydrochloride)	C
Estrogens Conjugated (Estrogens Conjugated)	C
Estradiol (Estradiol)	C
Tiagabine Hydrochloride	

(Tiagabine
Hydrochloride) C
Sildenafil Citrate
(Sildenafil Citrate) C
Butorphanol Tartrate
(Butorphanol
Tartrate) C
Trimethobenzamide
Hydrochloride
(Trimethobenzamide
Hydrochloride) C
Lithium Carbonate
(Lithium Carbonate) C
Valproate Semisodium

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Valproate Semisodium) C
 Beclometasone Dipropionate (Beclometasone Dipropionate) C
 Clavulin (Amoxicillin Trihydrate, Clavulanate Potassium) C

Date:02/17/05ISR Number: 4590490-6Report Type:Expedited (15-DaCompany Report #2004009909
 Age:59 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG (200 Other MG, 3 IN 1 D), ORAL	Blood Creatinine Increased Condition Aggravated Encephalopathy Myoclonus Oedema Peripheral	Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL 60 MG (2 N 1 D), ORAL			Oxycocet (Paracetamol, Oxycodone Hydrochloride)	SS		ORAL
			Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL
62.5 MG (2 IN 1 D), ORAL			Bosentan (Bosentan)	SS		ORAL
			Warfarin (Warfarin) Furosemide (Furosemide) Prednisone (Prednisone) Epoprostenol Sodium	C C C		

(Epoprostenol Sodium)	C
Pantoprazole (Pantoprazole)	C
Metformin Hydrochloride (Metformin Hydrochloride)	C
Bactrim (Sulfamethoxazole, Trimethoprim)	C
Oxygen (Oxygen)	C
Lisinopril (Lisinopril)	C
Metolazone (Metolazone)	C

Date:02/17/05ISR Number: 4590496-7Report Type:Expedited (15-DaCompany Report #2005022778

Age: Gender: I/FU:F

Outcome	PT
Death	Completed Suicide
Other	Injury Asphyxiation

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pharmaceutical Product Complaint Suicidal Ideation	Report Source	Product	Role	Manufacturer	Route
			Consumer	Zoloft (Sertraline)	PS		
				Neurontin (Gabapentin)	SS		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		

Date:02/17/05ISR Number: 4590511-0Report Type:Expedited (15-DaCompany Report #2003008068

Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability		Activities Of Daily Living Impaired	Consumer	Neurontin (Gabapentin)	PS		
Other 1200 MG (TID), ORAL		Anaemia Aplastic Anaemia Arthralgia Asthenia Balance Disorder Burns Second Degree Coma Condition Aggravated Convulsion Coordination Abnormal Disturbance In Attention Dizziness Drug Intolerance Dry Mouth Fall Grip Strength Decreased Headache Hemiparesis Ill-Defined Disorder Impaired Driving Ability Loss Of Consciousness Mental Impairment Muscle Spasms Nervousness Pain	Professional	Dilantin (Phenytoin Sodium)	SS		ORAL
				Acetylsalicylic Acid (Acetylsalicylic Acid)	SS		
				Vitamins With Minerals (Vitamins With Minerals)	SS		
				Megestrol (Megestrol)	C		

Panic Attack
Paralysis
Red Blood Cell Count
Increased
Speech Disorder

Date:02/18/05ISR Number: 4586473-2Report Type:Expedited (15-DaCompany Report #US-ROCHE-376030
Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening REPORTED		Aphasia		Klonopin	PS	Roche	ORAL
START DATE AS PRIOR TO 16 JANUARY 2001.		Dizziness					
		Drug Interaction					
		Dystonia					
DOSING AMOUNT REPORTED AS:		Hallucination		Lamictal	I		ORAL
		Hypoglycaemia		Neurontin	I		ORAL
		Vision Blurred					

"2/1/1/2".

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

15 DAY

	Xanax	I	ORAL
	Gabitril	I	ORAL
	Celexa	I	ORAL

Date:02/18/05ISR Number: 4586890-0Report Type:Expedited (15-DaCompany Report #US-ROCHE-376030
 Age:29 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening REPORTED	Dizziness		Klonopin	PS	Roche	ORAL
START DATE AS PRIOR TO 16 JANUARY 2001.	Drug Interaction Dystonia Hypoglycaemia					
DOSING AMOUNT REPORTED AS: "2/1/1/2".	Oedema Peripheral Vision Blurred		Lamictal Neurontin	I I		ORAL ORAL

15 DAY

	Xanax	I	ORAL
	Gabitril	I	ORAL
	Celexa	I	ORAL

Date:02/18/05ISR Number: 4586937-1Report Type:Expedited (15-DaCompany Report #US-ROCHE-394642
 Age:29 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening REPORTED	Aphasia		Klonopin	PS	Roche	ORAL
Hospitalization - START DATE AS Initial or Prolonged PRIOR TO 16 JANUARY 2001.	Dizziness Drug Interaction Dystonia					

Hallucination	Lamictal	I	ORAL
Hypoglycaemia	Neurontin	I	ORAL
Oedema Peripheral	Xanax	I	ORAL
Vision Blurred	Gabitril	I	ORAL

15 DAY

Celexa	I	ORAL
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EVERY NIGHT.

Date:02/18/05ISR Number: 4587761-6Report Type:Expedited (15-DaCompany Report #US-AMGEN-US109092
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anisocytosis		Epogen	PS		
SUBCUTANEOUS		Blood Albumin Decreased		Neurontin	SS		
		Bone Marrow Depression		Coumadin	C		
		Elliptocytosis		Colchicine	C		
		Iron Deficiency		Amiodarone	C		
		Malnutrition		Diltiazem	C		
		Therapeutic Response		Hectoral	C		
		Decreased		Coreg	C		
		Thrombocytopenia					

Date:02/18/05ISR Number: 4588616-3Report Type:Expedited (15-DaCompany Report #GXKR2005GB00583
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Foreign	Rifampicin (Ngx)			
		Drug Interaction	Health	(Rifampicin)	PS		
			Professional	Gabapentin			
			Other	(Ngx)(Gabapentin)	SS		ORAL

ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/18/05ISR Number: 4590005-2Report Type:Direct
Age:78 YR Gender:Female I/FU:I

Company Report #CTU 240807

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased Insomnia		Gabapentin 100 Mg Teva	PS	Teva	ORAL
100 MG		Pharmaceutical Product					
NIGHTLY ORAL		Complaint					

Date:02/18/05ISR Number: 4590271-3Report Type:Direct
Age:66 YR Gender:Male I/FU:I

Company Report #CTU 240828

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product Complaint		Neurontin (Gabapentin)	PS		
600 MG TID		Therapeutic Product Ineffective		Hydrocodone Enalapril Glucophage Avodart	C C C C		

Date:02/18/05ISR Number: 4590284-1Report Type:Direct
Age: Gender: I/FU:I

Company Report #CTU 240799

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Acidosis		Ceftriaxone 2 Gm Iv			
Other		Body Temperature		Daily/Roche	PS	Roche	
INTRAVENOUS	2 GM DAY IV	Increased Dehydration Haemolytic Anaemia Heart Rate Increased Intervertebral Discitis Multi-Organ Failure Renal Failure Respiratory Failure Sepsis		Enoxaparin Gabapentin Ondansetron Apap Hydroxyzine Clonidine Percocet-10	SS SS SS SS SS SS SS		

Date:02/18/05ISR Number: 4590338-XReport Type:Direct
Age:50 YR Gender:Male I/FU:I

Company Report #CTU 240796

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Neurontin (Generic)	PS		
900 MG TID							

Date:02/18/05ISR Number: 4591303-9Report Type:Expedited (15-DaCompany Report #2005028496
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Asthenia	Foreign	Neurontin (Tablets)			
(600 MG),ORAL		Diplopia	Consumer	(Gabapentin)	PS		ORAL
		Dizziness		Carbamazepine			
		Feeling Abnormal		(Carbamazepine)	C		
		Hearing Impaired		Baclofen (Baclofen)	C		
		Nervousness		Calcium With Vitamin			
		Somnolence		D (Calcium			
		Visual Disturbance		Phosphate, Calcium			
		Vomiting		Sodium Lactate,			
				Ergocalciferol)	C		
				Thiamine			
				Hydrochloride			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Thiamine
Hydrochloride) C
Phenobarbital Sodium
(Phenobarbital
Sodium) C
Phenytoin
(Phenytoin) C

Date:02/18/05ISR Number: 4591670-6Report Type:Expedited (15-DaCompany Report #2005028641
Age:47 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 1800 MG (1 Initial or Prolonged D), ORAL	Coma Intentional Misuse	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL			Hydrocodone (Hydrocodone)	SS		ORAL
ORAL			Carisprodol (Carisoprodol)	C		ORAL

Date:02/18/05ISR Number: 4591802-XReport Type:Expedited (15-DaCompany Report #S04-USA-05512-01
Age:29 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Other 40 MG QHS PO	Aphasia Diplopia	Health Professional	Celexa (Citalopram Hydrobromide)	PS		ORAL
4 MG BID	Dizziness Drug Interaction		Gabitril (Tiagabine Hydrochloride)	SS		
1800 MG QD	Dystonia Hallucination, Auditory		Neurontin (Gabapentin)	SS		
0.5 MG PRN	Hypoglycaemia		Xanax (Alprazolam)	SS		
0.5 MG BID	Oedema Peripheral		Klonopin (Clonazepam)	SS		

100 MG QAM

Lamictal

SS

50 MG QHS

Lamictal

SS

Date:02/18/05ISR Number: 4594002-2Report Type:Expedited (15-DaCompany Report #2005028590

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysphagia	Foreign	Neurontin			
		Parkinson'S Disease	Health	(Gabapentin)	PS		ORAL
ORAL			Professional	Sinemet (Carbidopa, Levodopa)	C		
				Carbamazepine (Carbamazepine)	C		

Date:02/22/05ISR Number: 4591495-1Report Type:Expedited (15-DaCompany Report #2005027778

Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Interaction	Consumer	Neurontin			
		Grand Mal Convulsion		(Gabapentin)	PS		
				Carbamazepine (Carbamazepine)	SS		
				Doxycycline (Doxycycline)	SS		

200 MG

(100MG,2 IN 1

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Freedom Of Information (FOI) Report

D)

Combivent
 (Ipratropium
 Bromide, Salbutamol
 Sulfate) C
 Salbutamol
 (Salbutamol) C
 Multivitamins
 (Ascorbic Acid,
 Ergocalciferol,
 Folic Acid,
 Nicotinamide, C
 Guaifenesin
 (Guaifenesin) C
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C
 Glipizide
 (Glipizide) C

Date:02/22/05ISR Number: 4591553-1Report Type:Expedited (15-DaCompany Report #2004112247
 Age:74 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other UNKNOWN	900 MG	Ankle Fracture Arthritis (3 IN Asthenia	Consumer Company Representative	Neurontin (Gabapentin)	PS		
1 D), UNKNOWN		Balance Disorder Confusional State		Oxcarbazepine (Oxcarbazepine)	SS		
UNKNOWN	UNKNOWN	Depression Dizziness Drug Ineffective Dyskinesia Fall Feeling Abnormal Feeling Drunk Head Injury Headache Hypotonia		Alendronate Sodium (Alendronate Sodium) Thyroid (Thyroid) Naproxen Sodium (Naproxen Sodium) Calcium Carbonate (Calcium Carbonate) Acetylsalicylic Acid (Acetylsalicylic Acid)	C C C C C C		

Influenza	Centrum Silver	
Insomnia	(Ascorbic Acid,	
Memory Impairment	Calcium, Minerals	
Mental Impairment	Nos, Retinol,	
Muscle Twitching	Tocopheryl Acetate,	C
Neuropathy Peripheral	Fish Oil (Fish Oil)	C
Subdural Haemorrhage	Atorvastatin	
Vision Blurred	(Atorvastatin)	C
	Carbamazepine	
	(Carbamazepine)	C
	Phenytoin Sodium	
	(Phenytoin Sodium)	C
	Carbamazepine	
	(Carbamazepine)	C
	Phenytoin	
	(Phenytoin)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/22/05ISR Number: 4591931-0Report Type:Expedited (15-DaCompany Report #2004111569
Age:77 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 400 MG (100 Other MG, 4 IN 1 D), ORAL	Gait Disturbance Joint Dislocation Postoperative Memory Impairment Refusal Of Treatment By Patient Sleep Attacks Somnolence Stupor Surgical Procedure Repeated	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Terazosin Hydrochloride (Terazosin Hydrochloride)	C		
			Potassium (Potassium)	C		
			Levothyroxine Sodium (Levothyroxine Sodium)	C		

Date:02/22/05ISR Number: 4591977-2Report Type:Expedited (15-DaCompany Report #2005028609
Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 800 MG (400 MG, 2 AT BEDTIME), ORAL	Blood Urine Present Haemorrhage	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Phenytoin Sodium (Phenytoin Sodium)	C		
			All Other Therapeutic Products (All Other Therapeutic Products)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Cough	Foreign	Neurontin			
Hospitalization -		Dysphagia	Health	(Gabapentin)	PS		ORAL
1200 MG (400							
Initial or Prolonged		Retching	Professional				
MG, 3 IN 1							
Other			Other				
D), ORAL				Valproate Sodium			
				(Valpoate			
				Sodium)	C		
				Clobazam			
				(Clobazam)	C		
				Anovlar			
				(Ethinylestradiol,			
				Norethisterone			
				Acetate)	C		
				Zuclopenthixol			
				(Zuclopenthixol)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/23/05ISR Number: 4591357-XReport Type:Direct
 Age:75 YR Gender:Male I/FU:I

Company Report #CTU 241084

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
ONE CAPSULE		Disease Recurrence		Gabapentin 300mg	PS	Greenstone	
TWICE DAILY		Drug Ineffective					
		Phantom Pain					
		Pharmaceutical Product					
		Complaint					

Date:02/23/05ISR Number: 4591374-XReport Type:Direct
 Age:52 YR Gender:Female I/FU:I

Company Report #CTU 241240

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Neurontin	PS		ORAL
400 MG TID		Pharmaceutical Product					
PO	5 YR	Complaint					

Date:02/23/05ISR Number: 4591375-1Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 241241

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Neurontin 400mg	PS		
BREAKFAST &		Electroencephalogram					
Initial or Prolonged		Abnormal		Neurontin	SS		
SUPPER							
LUNCH & HS							

Date:02/23/05ISR Number: 4591387-8Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 241244

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

2 CAP 3 Abdominal Rigidity Gabapentin 400 Mg PS ORAL
 Foreign Body Trauma
 TIMES DAILY Muscle Spasms
 PO Pharmaceutical Product
 Complaint

Date:02/23/05ISR Number: 4591398-2Report Type:Direct Company Report #CTU 241246
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Gabapentin	PS		ORAL
800 MG PO BID		Hallucination		Aminophylline	C		
		Sleep Walking		Lexapro	C		

Date:02/23/05ISR Number: 4591564-6Report Type:Direct Company Report #CTU 241192
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Chest Discomfort		Gabapentin			
ONE CAPSULE		Diarrhoea		(Neurontin) 100 Mg	PS		
THREE TIMES A		Nasal Congestion					
DAY							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/23/05ISR Number: 4594772-3Report Type:Expedited (15-DaCompany Report #KII-2005-0015236
Age:46 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged Other	Blood Pressure Diastolic Decreased Bradyphrenia Confusional State	Study Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		ORAL
ORAL			Gabapentin (Gabapentin)	SS		ORAL
ORAL	Depressed Level Of Consciousness					
	Heart Rate Increased Miosis Multiple Drug Overdose Oxygen Saturation Decreased Pupillary Reflex Impaired Respiratory Rate Decreased					

Date:02/23/05ISR Number: 4594891-1Report Type:Expedited (15-DaCompany Report #2005030141
Age:81 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG (600 MG, 3 IN 1 D), ORAL	Anaphylactic Reaction	Foreign Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Carbamazepine (Carbamazepine)	SS		
0.6 MG (0.2 MG, 3 IN 1 D), ORAL			Buprenorphine (Buprenorphine)	SS		ORAL
			Diclofenac (Diclofenac)	C		

Dyazide
 (Hydrochlorothiazide
 , Triamterene) C
 Risedronate Sodium
 (Risedronate Sodium) C
 Calcium (Calcium) C

Date:02/23/05ISR Number: 4595021-2Report Type:Expedited (15-DaCompany Report #2005028624
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (60 Other MG, 2 IN 1 D), ORAL		Hypotonia Pain Sedation Surgery Treatment Noncompliance	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Vicodin (Hydrocodone Bitartrate, Paracetamol) Cyclobenzaprine Hydrochloride Paroxetine Hydrochloride Clonazepam Lansoprazole Levothyroxine Sodium	C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Celecoxib C

Date:02/23/05ISR Number: 4595155-2Report Type:Expedited (15-DaCompany Report #2004121829
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Anxiety Fear Mental Disorder Nightmare Panic Reaction Pharmaceutical Product Complaint Suicidal Ideation	Consumer Health Professional	Gabapentin (Gabapentin) Venlafaxine Hydrochloride (Venlafaxine Hydrochloride) Risperidone (Risperidone) Zolpidem Tartrate (Zolpidem Tartrate) Escitalopram (Escitalopram)	PS C C C		

Date:02/23/05ISR Number: 4595160-6Report Type:Expedited (15-DaCompany Report #USA-2005-0018816
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer Health Professional Company	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		
80 MG, TID			Representative	Lorcet (Paracetamol, Hydrocodone Bitartrate) Neurontin (Gabapentin)	SS SS		
400 MG, TID				Soma (Carisoprodol) Tranxene (Clorazepate Dipotassium)	SS SS		
7.5 MG, BID				Atrazodone			

300 MG, HS

(Trazodone)

SS

Date:02/23/05ISR Number: 4595342-3Report Type:Expedited (15-DaCompany Report #S05-ESP-00586-01

Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	30 MG QD PO	Speech Disorder	Foreign Health	Seropram (Citalopram)	PS		ORAL
	2.5 MG QD PO		Professional Other	Lorazepam (Lorazepam)	SS		ORAL
	600 MG QD PO			Gabapentin	SS		ORAL

Date:02/23/05ISR Number: 4595680-4Report Type:Direct

Company Report #CTU 241229

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening	1 GM BID	Neutrophil Count		Sulfasalazine	PS		
	100 MG TID	Decreased		Gabapentin	SS		
(PAIN)		White Blood Cell Count					
		Decreased					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/23/05ISR Number: 4692996-8Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 241061

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nausea		Neurontin	PS		
		Pharmaceutical Product Complaint					

Date:02/24/05ISR Number: 4591171-5Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 241360

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Gabapentin	PS		
300MG 3X		Headache					
		Pharmaceutical Product Complaint					
DAILY							

Date:02/24/05ISR Number: 4591226-5Report Type:Direct
Age:50 YR Gender:Male I/FU:I

Company Report #CTU 241346

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Neurontin (Generic Form)	PS		
900 MG TID							

Date:02/24/05ISR Number: 4595706-8Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 241257

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Gabapentin 300 Mg	PS		ORAL
1 CAP PO QHS							
		Pharmaceutical Product Complaint					

Date:02/24/05ISR Number: 4595714-7Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 241260

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Drug Ineffective		Gabapentin 600 Mg	PS		
1/2 PILL QID							
Other		Pharmaceutical Product Complaint		Darvocet N 100	C		

Date:02/24/05ISR Number: 4595725-1Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 241263

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
600 MG ONE		Agitation		Gabapentin	PS		
THREE TIMES		Disorientation					
DAILY		Memory Impairment					
		Mood Swings					
		Pharmaceutical Product Complaint					
		Tremor					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/24/05ISR Number: 4595808-6Report Type:Direct
Age:64 YR Gender:Female I/FU:I

Company Report #CTU 241249

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea		Gabapentin	PS		ORAL
2 PO TID		Oropharyngeal Swelling Pharmaceutical Product Complaint					

Date:02/24/05ISR Number: 4595809-8Report Type:Direct
Age: Gender: I/FU:I

Company Report #CTU 241248

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
INTRACTABLE		Drug Toxicity		Gabapentin	PS		
EPILEPSY		Nystagmus Pharmaceutical Product Complaint					

Date:02/25/05ISR Number: 4594942-4Report Type:Expedited (15-DaCompany Report #2005031290
Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abasia	Consumer	Lipitor			
10 MG (10 MG,		Drug Effect Decreased		(Atorvastatin)	PS		ORAL
1 IN 1 D),		Drug Ineffective					
ORAL		Feeling Hot And Cold					
		Hypoaesthesia		Neurontin			
ORAL		Insomnia		(Gabapentin)	SS		ORAL
		Neuropathy		Fluoxetine			
		Pain		Hydrochloride			
		Sedation		(Fluoxetine			
ORAL		Spinal Fracture		Hydrochloride)	SS		ORAL

Vision Blurred

Irbesartan	
(Irbesartan)	C
Digoxin (Digoxin)	C
Warrarin Sodium	
(Warfarin Sodium)	C
Escitalopram	
(Escitalopram)	C

Date:02/25/05ISR Number: 4596128-6Report Type:Expedited (15-DaCompany Report #A001-OCT-3720

Age:51 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Contusion Face Injury Fall Mental Status Changes	Study Health Professional	Aricept (Donepezil Hydrochloride) Marinol (Dronabinol) Kytril (Granisetron)	PS SS SS		
INTRAVENOUS	1 IN 1 D,						
INTRAVENOUS				Oxycontin (Oxycodone Hydrochloride)	SS		
1 IN 12 HR				Neurontin (Gabapentin) Ambien (Zolpidem Tartrate) Aranesp (Darbepoetin Alfa) Morphine Sulfate (Morphine Sulfate)	SS SS C C		

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Freedom Of Information (FOI) Report

Chlorpromazine	
(Chlorpromazine)	C
Oxy Ir (Oxycodone	
Hydrochloride)	C
Prograf (Tacrolimus)	C
Protonix	
(Pantoprazole)	C
Prandin	
(Repaglinide)	C
Zoloft (Sertraline	
Hydrochloride)	C
Glipizide	
(Glipizide)	C
Procardia	
(Nifedipine)	C
Magnesium	
(Magnesium)	C
Levaquin	
(Levofloxacin)	C
Pentamidine	
(Pentamidine)	C
Pericolace	
(Peri-Colace)	C
Dulcolax Suppository	
(Bisacodyl)	C
Baby Aspirin	
(Acetylsalicylic	
Acid)	C
Lasix (Furosemide)	C
Leukine	
(Sargramostim	C
Dexamethasone	
(Dexamethasone)	C
Adriamycin	
(Doxorubicin)	C
Cisplatinum	
(Cisplatin)	C
Senekot (Senna	
Fruit)	C
Inderal (Propranolol	
Hydrochloride)	C
Multivitamins	
(Multivitamins)	C
Insulin (Insulin)	C
Aldactone	
(Spironolactone)	C
Megace (Megestrol	

Date:02/25/05ISR Number: 4596837-9Report Type:Expedited (15-DaCompany Report #KII-2005-0015263
Age:58 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Blood Creatine Increased
Initial or Prolonged	Blood Pressure Decreased
	Blood Urea Increased
	Depressed Level Of
	Consciousness
	Disorientation
	Feeling Abnormal

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Medication Error	Report Source	Product	Role	Manufacturer	Route
		Infection Lethargy					
PRN, ORAL		Myoclonus	Study	Morphine Sulfate	PS		ORAL
PRN, ORAL			Health	Hydromorphone Hcl	SS		ORAL
ORAL			Professional Other	Gabapentin (Gabapentin)	SS		ORAL
ORAL				Hypnotics And Sedatives()	SS		ORAL
				Antibiotics	C		
				Insulin (Insulin)	C		
				Serum Lipid Reducing Agents	C		
				Antiplatelet	C		
				Asa	C		
				Cefoxitin (Cefoxitin)	C		

Date:02/25/05ISR Number: 4597551-6Report Type:Direct
Age:61 YR Gender:Female I/FU:I

Company Report #CTU 241380

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
BID		Pain		Neurontin Generic	PS		
		Paraesthesia Pharmaceutical Product Complaint Somnolence Stomach Discomfort					

Date:02/28/05ISR Number: 4592757-4Report Type:Expedited (15-DaCompany Report #US-SOLVAY-00204002739
Age:18935 DYGender:Male I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Daily Dose:		Anaemia		Marinol	PS		ORAL

Hospitalization -			Confusional State			
unk.						
Initial or Prolonged			Convulsion	Aranesp	SS	
SUBCUTANEOUS	Daily	Dose:	Face Injury			
200 iu.						
Daily Dose:			Fall	Oxycontin	SS	ORAL
40 mg.	31	DAY	Hepatic Neoplasm			
Daily Dose:			Malignant	Aricept	SS	ORAL
unk.			Mental Status Changes			
Daily Dose:			Nausea	Neurontin	SS	ORAL
unk.			Pancytopenia			
INTRAVENOUS	Daily	Dose: 1	Vomiting	Kytril	SS	
mg.	1	DAY				
UNKNOWN	Daily	Dose:		Morphine Sulfate	C	
15 mg.						
Frequency:As						
needed	55	DAY		Zoloft	C	ORAL
Daily Dose:						
100 mg.				Procardia	C	ORAL
Daily Dose:						
120 mg.				Dexamethasone	C	
INTRAVENOUS	Daily	Dose:				
10 mg.						
Frequency:Onc						
e	1	DAY		Leukine	C	
SUBCUTANEOUS	Daily	Dose:				
500 mg.	7	DAY				

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

INTRAVENOUS	Daily Dose:		Lasix	C	
40 mg.					
Frequency:Onc					
e	1 DAY				
Daily Dose:			Asa	C	ORAL
81 mg.					
RECTAL	Daily Dose: 5		Dulcolax	C	
mg.					
Frequency:As					
needed					
UNKNOWN	Daily Dose: 2		Pericolace	C	
DF.					
RESPIRATORY					
(INHALATION)	Daily Dose:				
unk.					
Daily Dose:			Levaquin	C	ORAL
500 mg.	8 DAY				
Daily Dose:			Magnesium	C	ORAL
800 mg.					
SUBCUTANEOUS	Daily dose:		Darbepoetin Alfa	C	
unknown	87 DAY				
Daily Dose:			Megace	C	ORAL
unk.					

Daily Dose:		Aldactone	C	ORAL
unk.				
UNKNOWN	Daily Dose:	Insulin	C	
unk.				
UNKNOWN	Daily Dose:	Multivitamins	C	
unk.				
Daily Dose:		Inderal	C	ORAL
unk.				
UNKNOWN	Daily Dose:	Senecot	C	
unk.				
SUBCUTANEOUS	Daily Dose:	Cisplatinum	C	
unk.				
SUBCUTANEOUS	Daily Dose:	Adriamycin	C	
unk.				
Daily Dose:		Glipizide	C	ORAL
500 mg.				
Daily Dose:		Prandin	C	ORAL
0.1 mg.				
UNKNOWN	Daily Dose:	Protonix	C	
40 mg.				
UNKNOWN	Daily Dose: 8	Prograf	C	
mg.				
UNKNOWN	Daily Dose:	Chlorpromazine	C	
25 mg.				
Frequency:As				
needed				

UNKNOWN Daily Dose: 4

Prograf C

mg.

Oxycodone C

ORAL

Daily Dose:

30 mg.

Date:02/28/05ISR Number: 4592953-6Report Type:Expedited (15-DaCompany Report #ES-GLAXOSMITHKLINE-B0372968A
Age:35 YR Gender:Male I/FU:F

Outcome PT
Life-Threatening Granulocytopenia
Haemoglobin Decreased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Leukopenia White Blood Cell Count Decreased	Report Source	Product	Role	Manufacturer	Route
600MG Twice per day	21 DAY			Zidovudine	PS	Glaxosmithkline	ORAL
300MG Per day				Epivir	SS	Glaxosmithkline	ORAL
1.2G Three times per day				Gabapentin	SS		ORAL
UNKNOWN 80MG Twice per day	600MG Per day			Efavirenz	C		
				Stavudine	C		ORAL

Date:02/28/05ISR Number: 4595408-8Report Type:Direct
Age:50 YR Gender:Male I/FU:I

Company Report #CTU 241546

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Rash		Neurontin (Generic Form)	PS		
900 MG TID							

Date:02/28/05ISR Number: 4595527-6Report Type:Direct
Age:66 YR Gender:Male I/FU:I

Company Report #CTU 241537

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Ineffective Pharmaceutical Product		Generic Neurontin (Gabapentin)	PS		
600 MG TID		Complaint		Hydrocodone	C		
				Enalapril	C		
				Avodant	C		
				Glucophage	C		

Date:02/28/05ISR Number: 4596211-5Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 241501

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Rash		Gabapentin 800mg Two			
800 MG TWO		Urticaria		Tid	PS		
TID							

Date:02/28/05ISR Number: 4596403-5Report Type:Expedited (15-DaCompany Report #2005027210
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Interaction	Foreign	Neurontin			
Other		Hallucination, Visual	Health	(Gabapentin)	PS		ORAL
900 MG (300		Sleep Disorder	Professional				
MG, 3 IN 1							
D); ORAL							
80 MG (40 MG,				Oxycodone			
2 IN 1 D);				(Oxycodone)	SS		ORAL
ORAL							
				Simvastatin	C		
				Atenolol	C		
				Acetylsalicylic Acid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/05ISR Number: 4596620-4Report Type:Expedited (15-DaCompany Report #2005031559
 Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arthropathy	Literature	Gabapentin			
Hospitalization - 600 MG (300 Initial or Prolonged MG, 2 IN 1 Other D), ORAL		Breast Cancer Metastatic	Health	(Gabapentin)	PS		ORAL
		Cerebellar Syndrome	Professional				
		Coordination Abnormal					
		Fall		Glibenclamide			
		Fear		(Glibenclamide)	C		
		Gait Disturbance		Metoprolol Tartrate			
		Hyporeflexia		(Metoprolol Tartrate)	C		
		Walking Aid User		Hyzaar			
				(Hydrochlorothiazide , Losartan Potassium)	C		
				Fentanyl (Fentanyl)	C		
				Omeprazole			
				(Omeprazole)	C		
				Docusate Sodium			
				(Docusate Sodium)	C		
				Bisacodyl			
				(Bisacodyl)	C		
				Epoetin Alfa			
				(Epoetin Alfa)	C		
				Sucralfate			
				(Sucralfate)	C		

Date:02/28/05ISR Number: 4596814-8Report Type:Expedited (15-DaCompany Report #2004104699
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Circulatory Collapse	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
			Professional Company Representative	Gabapentin (Gabapentin)	SS		

Outcome	PT
Hospitalization -	Abasia
Initial or Prolonged	Abnormal Behaviour
Disability	Arthralgia
Other	Bladder Prolapse
	Blindness
	Contusion
	Delusion
	Dizziness
	Dysstasia
	Exostosis
	Fall
	Illusion
	Intervertebral Disc Space
	Narrowing
	Joint Dislocation
	Knee Deformity
	Mean Cell Volume

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Increased Medication Error Middle Insomnia Noctiphobia					
300 MG, ORAL		Osteoarthritis Osteopenia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
200 MG (200 MG, 1 IN 1 D), ORAL		Red Blood Cell Sedimentation Rate Increased	Professional	Celebrex (Celecoxib)	SS		ORAL
		Spinal Osteoarthritis Thinking Abnormal Vertigo		Quinine (Quinine) Centrum Silver (Ascorbic Acid, Calcium, Minerals Nos, Retinol, Tocopheryl Acetate, Calcium (Calcium) Ascorbic Acid (Ascorbic Acid) Ibuprofen (Ibuprofen) Paracetamol (Paracetamol) Indometacin (Indometacin)	C C C C C C		

Date:02/28/05ISR Number: 4596937-3Report Type:Expedited (15-DaCompany Report #2005019377
Age: Gender:Female I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (1 IN1 Other D), ORAL		Dyspnoea Electrophoresis Protein Abnormal	Health Professional Company	Neurontin (Gabapentin)	PS		ORAL
10/650 MG (1 IN 1 D), ORAL		Faecal Occult Blood Positive Fatigue Hypercalcaemia	Representative	Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		ORAL

Hypothyroidism
 Irritable Bowel Syndrome
 Palpitations
 Pyrexia
 Renal Failure Acute

Olanzapine/Fluoxetine
 Hydrochloride
 (Fluoxetine
 Hydrochloride, ,
 Olanzapine) SS

ORAL

6/25 (1 IN 1

Sinusitis

D), ORAL

Date:02/28/05ISR Number: 4596956-7Report Type:Expedited (15-DaCompany Report #2004051426

Age:29 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG, ORAL		Aphasia Diplopia	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Dizziness Drug Interaction Dystonia		Xanax Tablet (Alprazolam) (Alprazolam)	SS		ORAL
ORAL		Hallucination Hypoglycaemia Oedema Peripheral Vision Blurred		Tiagabine Hydrochloride (Tiagabine Hydrochloride)	SS		ORAL
8 MG (4 MG, 2 IN 1 D), ORAL				Lamotrigine (Lamotrigine)	SS		ORAL
1 D, ORAL				Citalopram			

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Freedom Of Information (FOI) Report

1 IN 1 D, ORAL	Hydrobromide (Citalopram Hydrobromide)	SS	ORAL
2 IN 1 D, ORAL	Clonazepam (Clonazepam)	SS	ORAL

Date:02/28/05ISR Number: 4596993-2Report Type:Expedited (15-DaCompany Report #2005006035
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Cardiac Disorder	Consumer	Neurontin			
Other		Condition Aggravated	Health	(Gabapentin)	PS		
600 MG (300 MG, 2 IN 1 D)		Diabetic Neuropathy	Professional				
		Drug Ineffective		Sulfasalazine			
		Emphysema		(Sulfasalazine)	C		
		Oedema Peripheral		Lisinopril			
		Visual Acuity Reduced		(Lisinopril)	C		
				Glipizide			
				(Glipizide)	C		
				Pravastatin Sodium			
				(Pravastatin Sodium)	C		
				Ezetimibe			
				(Ezetimibe)	C		
				Salbutamol			
				(Salbutamol)	C		
				Furosemide			
				(Furosemide)	C		

Date:02/28/05ISR Number: 4596999-3Report Type:Expedited (15-DaCompany Report #2005032149
Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Asthenia	Consumer	Neurontin			

Initial or Prolonged	Fatigue	(Gabapentin)	PS
900 MG (300			
Other	Gastrooesophageal Reflux		
MG, 3 IN 1 D)			
	Disease	Oxycodone	
	Pneumonia	Hydrochloride	
	Respiratory Depression	(Oxycodone	
	Somnolence	Hydrochloride)	C
		Vicodin (Hydrocodone	
		Bitartrate,	
		Paracetamol)	C
		Diltiazem	
		Hydrochloride	
		(Diltiazem	
		Hydrochloride)	C
		Esomeprazole	
		(Esomeprazole)	C
		Rosuvastatin	
		(Rosuvastatin)	C
		Vitamins (Vitamins)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:02/28/05ISR Number: 4597348-7Report Type:Direct
 Age:80 YR Gender:Female I/FU:I

Company Report #CTU 241498

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product		Gabapentin 100 Mg	PS		ORAL
200 MG PO QD		Complaint Sedation					

Date:02/28/05ISR Number: 4597399-2Report Type:Direct
 Age:73 YR Gender:Female I/FU:I

Company Report #CTU 241495

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diabetic Neuropathy		Neurontin Tab	PS		
600 MG DAILY		Drug Ineffective Neuropathy Paraesthesia					

Date:02/28/05ISR Number: 4597400-6Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 241494

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Irritation		Gabapentin 100 Mg	PS		ORAL
9 QD PO							

Date:02/28/05ISR Number: 4597518-8Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 241481

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Leg Amputation Phantom Pain		Gabapentin 300 Mg Tablets	PS		
900 MG Q AM AND Q HS; 600 MG Q NOON							

Date:02/28/05ISR Number: 4597522-XReport Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 241478

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion		Neurontin 600 Mg Two			
600 MG TWO				Am, One Noon, One Pm	PS		
AM, ONE NOON,							
ONE PM							

Date:02/28/05ISR Number: 4597525-5Report Type:Direct
Age:46 YR Gender:Male I/FU:I

Company Report #CTU 241474

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Pharmaceutical Product		Gabapentin 100 Mg			
Intervention to		Complaint		-Purepac	PS	Purepac	ORAL
100 MG PO							
Prevent Permanent		Rash					
Impairment/Damage		Swelling Face					

Date:03/01/05ISR Number: 4593866-6Report Type:Expedited (15-DaCompany Report #US-ROCHE-394642
Age:29 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

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
REPORTED		Aphasia		Klonopin	PS	Roche	ORAL
START DATE AS		Diplopia					
PRIOR TO 16		Dizziness					
JANUARY 2001.		Drug Interaction					
		Dystonia		Lamictal	I		ORAL
		Hallucination		Neurontin	I		ORAL
		Hypoglycaemia		Xanax	I		ORAL
15 DAY		Oedema Peripheral		Gabitril	I		ORAL
EVERY NIGHT.		Vision Blurred		Celexa	I		ORAL

Date:03/01/05ISR Number: 4597143-9Report Type:Expedited (15-DaCompany Report #2005036213
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Dizziness	Foreign	Gabapentin			
Other		Memory Impairment	Health	(Gabapentin)	PS		ORAL
300 MG (300			Professional				
MG , 1 IN 1							
D), ORAL				Diclofenac			
				(Diclofenac)	C		

Date:03/01/05ISR Number: 4597223-8Report Type:Expedited (15-DaCompany Report #2005033656
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Deafness	Foreign	Neurontin			
Initial or Prolonged		Deafness Neurosensory	Consumer	(Gabapentin)	PS		
Other		Ear Discomfort					

Tremor
Vaginal Haemorrhage

Date:03/01/05ISR Number: 4598228-3Report Type:Direct Company Report #CTU 241701
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Dysaesthesia Pharmaceutical Product Complaint		Neurontin	PS		

Date:03/01/05ISR Number: 4598413-0Report Type:Direct Company Report #CTU 241746
Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Gabapentin 100 Mg Po Bid	PS		ORAL
PO QD							

Date:03/01/05ISR Number: 4598810-3Report Type:Expedited (15-DaCompany Report #2005033652
Age: Gender:Male I/FU:I

Outcome	PT
Other	Disorientation Drug Ineffective

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Dyspnoea Hypertension	Report Source	Product	Role	Manufacturer	Route
			Consumer	Neurontin (Gabapentin)	PS		
				Lorazepam (Lorazepam)	C		
				Atenolol (Atenolol)	C		
				Digoxin (Digoxin)	C		
				Losartan Potassium (Losartan Potassium)	C		
				Amlodipine Besilate (Amlodipine Besilate)	C		
				Vitamins (Vitamins)	C		

Date:03/01/05ISR Number: 4598814-0Report Type:Expedited (15-DaCompany Report #2005032613
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Deafness Unilateral Vertigo					
300 MG (100 MG, 3 IN 1 D), ORAL		Viral Labyrinthitis					

Date:03/01/05ISR Number: 4598848-6Report Type:Expedited (15-DaCompany Report #2004120003
Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anxiety Blood Urine Present	Health Professional	Neurontin (Gabapentin)	PS		
200 MG (200 MG, 1 IN 1 D)		Bundle Branch Block Right	Company				
		Conjunctivitis Convulsion Ear Discomfort	Representative	Propacet (Dextroprooxyphene			

Feeling Abnormal	Napsilate,	
Fluid Overload	Paracetamol)	C
Gait Disturbance	Lisinopril	
Hyponatraemia	(Lisinopril)	C
Hypoxia		
Inappropriate		
Antidiuretic Hormone		
Secretion		
Medication Error		
Muscle Twitching		
Musculoskeletal		
Discomfort		
Neuralgia		
Overweight		
Pain		
Paraesthesia		
Pulmonary Oedema		
Spinal Osteoarthritis		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/01/05ISR Number: 4598854-1Report Type:Expedited (15-DaCompany Report #2004021501
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Balance Disorder Condition Aggravated	Consumer Health	Neurontin (Gabapentin)	PS		
300 MG (DAILY)		Depression	Professional				
		Difficulty In Walking Drug Interaction		Escitalopram (Escitalopram)	SS		
20 MG (DAILY)		Fall Feeling Drunk Gait Disturbance Hallucination, Visual Migraine Nightmare		Allegra-D (Fexofenadine, Pseudoephedrine Hydrochloride) Propacet (Dextropropoxyphene Napsilate, Paracetamol) Pantoprazole (Pantoprazole) Docusate (Docusate)	C C C C		

Date:03/02/05ISR Number: 4598320-3Report Type:Expedited (15-DaCompany Report #USA-2004-0016457
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Anaemia Asthenia Blood Glucose Increased Blood Sodium Decreased	Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet	PS		
20 MG, Q12H,		Confusional State Contusion Eye Injury Fall Fatigue		Oxyir Capsules 5mg (Oxycodone Hdyrochloride, Oxycodone Hydrochloride)Ir	SS		
5 MG, Q4H,		Mental Status Changes Nausea Pancytopenia		Aranesp (Darbepoetin) Protonix	SS		

	Vomiting	(Pantoprazole)	SS
40 MG DAILY		Gabapentin	
		(Gabapentin)	SS
		Donepezil	SS
		Dronabinol	
		(Dronabinol)	SS
		Insulin (Inslin)	SS
		Ambien (Zolpidem	
		Tartrate)	SS
		Senokot(Sennosidea+B	
)	SS
		Aldactone	
		(Spironolactone)	SS
		Megacet (Megestrol	
		Acetate)	SS
		Inderal (Propranolol	
		Hydrochloride)	SS
		Magnesium	
		(Magnesium)	SS
400 MG		Acetylsalicylic Acid	
		(Acetylsalicylic	
		Acid)	SS
81 MG DAILY		Prograf (Tacrolimus)	SS
SEE IMAGE		Kytril (Granisetron)	C
INTRAVENOUS	1 MG,		

Freedom Of Information (FOI) Report

INTRAVENOUS

Chlorpromazine (Chlorpromazine)	C
Glipizide (Glipizide)	C
Procardia	C
Pentamidine (Pentamidine)	C
Peri-Colace (Sennodide A+B, Docusate Sodium)	C
Dulcolax	C
Dexamethasone (Dexamethasone)	C
Cis-Platium (Cisplatin)	C
Zoloft (Sertraline Hydrochloride)	C
Leukine (Sargramostim)	C
Adriamycin (Doxorubicin)	C

Date:03/02/05ISR Number: 4599096-6Report Type:Expedited (15-DaCompany Report #US014711
Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1600 UG PRN		Communication Disorder	Foreign	Fentanyl Citrate	PS		ORAL
ORAL		Disorientation	Study				
600 MG DAILY		Drug Interaction	Health	Gabapentin	SS		
2.5 GR DAILY		Drug Toxicity	Professional	Capecitabine	SS		
			Other	Oxycontin	C		
				Paracetamol	C		
				Dexamethasone	C		
				Lansoprazole	C		
				Laxoberal	C		

Date:03/02/05ISR Number: 4599310-7Report Type:Direct Company Report #CTU 241779
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache		Neurontin	PS		
300 MG 2 TID		Malaise Pharmaceutical Product Complaint Vomiting					

Date:03/02/05ISR Number: 4599315-6Report Type:Direct Company Report #CTU 241789
Age:91 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oedema Peripheral Pharmaceutical Product		Neurontin 100 Mg Pfizer	PS	Pfizer	
100 MG 3 BID		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/05ISR Number: 4599327-2Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 241792

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Neurontin (Generic)	PS		
600 MG TID	1	MON					

Date:03/02/05ISR Number: 4599331-4Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 241801

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation Insomnia		Generic Neurontin	PS		

Date:03/02/05ISR Number: 4599343-0Report Type:Direct
 Age:53 YR Gender:Male I/FU:I

Company Report #CTU 241810

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Gabapentin	PS		
600 TID							
Pharmaceutical Product Complaint							

Date:03/02/05ISR Number: 4599347-8Report Type:Direct
 Age: Gender: I/FU:I

Company Report #CTU 241836

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Headache Pain		Gabapentin	PS		
Pharmaceutical Product Complaint							

Date:03/02/05ISR Number: 4599348-XReport Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 241839

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Dreams		Gabapentin 300 Mg			
		Pharmaceutical Product		Tid	PS		ORAL
300 MG TID PO		Complaint					
DAILY							
				Glyburide	C		
				Warfarin	C		
				Ditropan	C		
				Pravachol	C		
				Furosemide	C		
				Pentoxifylline	C		

Date:03/02/05ISR Number: 4599356-9Report Type:Direct Company Report #CTU 241856
Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour		Neurontin	PS		
400 MG 4 X		Agitation					
DAILY		Pharmaceutical Product					
		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/02/05ISR Number: 4599359-4Report Type:Direct
 Age:46 YR Gender:Male I/FU:I

Company Report #CTU 241859

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product		Neurontin	PS		ORAL
300 MG ONE		Complaint					
PO QID		Syncope					

Date:03/02/05ISR Number: 4599360-0Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 241861

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased		Neurontin			
800 MG QID		Headache		(Gabapentin) Generic	PS		
		Nausea					
		Pharmaceutical Product					
		Complaint					

Date:03/02/05ISR Number: 4599361-2Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 241863

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache		Neurontin			
800 MG QID				(Gabapentin) Generic	PS		

Date:03/02/05ISR Number: 4599363-6Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 241869

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
600MG - 2 PO		Pharmaceutical Product		Neurontin (Generic)			
		Complaint		600 Mg	PS		ORAL

4 X DAY

Date:03/02/05ISR Number: 4599388-0Report Type:Direct
 Age:71 YR Gender:Female I/FU:I

Company Report #CTU 241897

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
400 MG 2 QGS		Condition Aggravated		Neurontin 400 Mg	PS		
		Depression					
		Drug Ineffective					
		Neuropathy					
		Pharmaceutical Product					
		Complaint					

Date:03/02/05ISR Number: 4600008-7Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 241773

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
800 MG 2 TABS		Drug Tolerance Decreased		Neurontin	PS		
QID		Pharmaceutical Product					
		Complaint					

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Freedom Of Information (FOI) Report

Date:03/03/05ISR Number: 4597631-5Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0544971A
 Age:45 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Blister		Lamictal	PS	Glaxosmithkline	ORAL
Other		Convulsion		Neurontin	SS		
5 WK		Per					
UNKNOWN	3000MG	Erythema					
day	7 YR	Oral Mucosal Blistering		Penicillin	SS	Glaxosmithkline	
UNKNOWN		Rash					
		Swelling Face					

Date:03/03/05ISR Number: 4598473-7Report Type:Direct Company Report #CTU 241965
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharmaceutical Product		Neurontin Generic	PS		
		Complaint					
		Urticaria					

Date:03/04/05ISR Number: 4600530-3Report Type:Expedited (15-DaCompany Report #2005038390
 Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Cardiovascular Disorder	Health	Celebrex (Celecoxib)	PS		
Initial or Prolonged		Carpal Tunnel Syndrome	Professional	Neurontin			
Other		Cataract		(Gabapentin)	SS		
SEE IMAGE		Knee Operation		Bextra (Valdecoxib)	SS		
		Neuropathy Peripheral		Bengay (Menthol)	SS		
		Nosocomial Infection		Paracetamol			
		Post Procedural		(Paracetamol)	C		
		Complication		Analgesics			
		Spinal Operation		(Analgesics)	C		
		Staphylococcal Infection					
		Streptococcal Infection					

Date:03/04/05ISR Number: 4601181-7Report Type:Expedited (15-DaCompany Report #2005034411
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Withdrawal Syndrome Suicidal Ideation	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
7200 MG (1800 MG, 4 IN 1 D), ORAL				Xanax Tablet (Alpazolam) Carbamazepine (Carbamazepine) Rosuvastatin (Rosuvastatin)	SS C C		

Date:03/07/05ISR Number: 4600993-3Report Type:Direct Company Report #CTU 242144
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
600 MG 1 PO TID		Headache Pharmaceutical Product Complaint Speech Disorder		Gabapentin Furosemide Spironolactone	PS C C		ORAL

Freedom Of Information (FOI) Report

Nitro Stat	C
Nitroglycerin Patch	C
Klor-Con	C
Plavix	C
Ecotrin	C
Toprol Xl	C
Zocor	C
Neurontin	C
Hydrocodone/Apap	C
Glucophage	C
Avandia	C
Amaryl	C
Fluoxetine	C
Ibuprofen	C
Prevacid	C
Ambien	C
Restoril	C
Sur-O-Lax	C
Promethazine	C
Droperridol	C
Albuterol	C

Date:03/07/05ISR Number: 4602383-6Report Type:Expedited (15-DaCompany Report #2004089776
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Condition Aggravated	Consumer	Neurontin			
Other		Diabetes Mellitus		(Gabapentin)	PS		ORAL
2700 MG (900		Diabetic Neuropathy					
MG, 3 IN 1		Drug Effect Decreased					
D), ORAL		Drug Ineffective		Gabapentin			
		Euphoric Mood		(Gabapentin)	SS		
		Food Interaction		Oxycodone			
		Neuropathic Arthropathy		Hydrochloride			
		Neuropathy		(Oxycodone			
				Hydrochloride)	C		

Date:03/07/05ISR Number: 4602406-4Report Type:Expedited (15-DaCompany Report #2004010780
 Age:69 YR Gender:Male I/FU:F

Outcome	PT
Disability	Actinic Keratosis
Other	Aldolase Increased
	Anhedonia
	Arthralgia
	Arthropathy
	Asthenia
	Blood Cholesterol Increased
	Blood Creatine Phosphokinase Increased
	Blood Creatinine Increased
	Blood Glucose Increased
	Blood Potassium Increased
	Blood Sodium Increased
	Blood Urine Present
	Bundle Branch Block Left
	Diverticulum

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
10 MG (1 IN 1 D)		Consumer	Lipitor (Atorvastatin)	PS		
300 MG, ORAL			Celebrex (Celecoxib) Neurontin (Gabapentin)	SS SS		ORAL
(DAILY), ORAL			Hyzaar (Hydrochlorothiazide , Losartan Potassium)	SS		ORAL
			Influenza Vaccine (Influenza Vaccine) All Other Therapeutic Products (All Other Therapeutic Products)	C C		
			Vicodin (Hydrocodone Bitartrate, Paracetamol) Acetylsalicylic Acid (Acetylsalicylic Acid) Amlodipine Besilate (Amlodipine Besilate)	C C C		

Date:03/07/05ISR Number: 4602461-1Report Type:Expedited (15-DaCompany Report #2004120003

Age:84 YR Gender:Female I/FU:F

Outcome PT
Hospitalization - Arthritis
Initial or Prolonged Aspartate
Other Aminotransferase
Increased
Blood Bicarbonate
Increased

Blood Chloride Decreased
Blood Pressure Diastolic
Decreased
Blood Pressure Systolic
Increased
Blood Urine Present
Bundle Branch Block Right
Cardiac Failure
Congestive
Conjunctivitis
Convulsion
Dyskinesia
Ear Discomfort
Exostosis
Fluid Overload
Gait Disturbance
Haematocrit Decreased
Haemoglobin Decreased
Hypoxia
Inappropriate

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Antidiuretic Hormone Secretion Memory Impairment Movement Disorder	Health Professional	Neurontin (Gabapentin)	PS		
200 MG (200 MG, 1 IN 1 D)		Muscle Spasms Muscle Tightness Muscle Twitching	Company Representative	Propacet (Dectropropoxyphene Napsilate, Paracetamol)	C		
		Myelopathy Neuralgia Overweight Paraesthesia Postictal State Pulmonary Oedema Qrs Axis Abnormal Spinal Disorder Spinal Osteoarthritis		Lisinopril (Lisinopril)	C		

Date:03/07/05ISR Number: 4602487-8Report Type:Expedited (15-DaCompany Report #2005024130
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Back Pain Bedridden Breast Pain Chest Pain	Consumer	Neurontin (Gabapentin)	PS		
600 MG		Drug Ineffective Neck Pain Pain Pain In Extremity Paraesthesia Pharmaceutical Product Complaint		Gabapentin (Gabapentin)	SS		

Date:03/07/05ISR Number: 4602494-5Report Type:Expedited (15-DaCompany Report #2005001941
Age:27 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abortion Threatened	Consumer	Neurontin			

Initial or Prolonged	Drug Exposure During	Health	(Gabapentin)	PS
	Pregnancy	Professional	Valproate Semisodium	
	Pregnancy		(Valproate	
	Vomiting		Semisodium)	SS
			Levetiracetam	
			(Levetiracetam)	SS
			Zonisamide	
			(Zonisamide)	SS

Date:03/07/05ISR Number: 4602495-7Report Type:Expedited (15-DaCompany Report #2005018142

Age:57 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Anger
Initial or Prolonged	Bruxism
Other	Disease Recurrence
	Influenza
	Irritability
	Irritable Bowel Syndrome
	Mood Altered
	Muscle Spasms

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
(600 MG)	ORAL	Nausea Pain Pain In Extremity	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
		Pharmaceutical Product Complaint	Professional	Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
		Trigeminal Neuralgia		Lansoprazole (Lansoprazole)	C		
				Librax (Chlordiazepoxide Hydrochloride, Clidinium Bromide)	C		
				Provella-14 (Estrogens Conjugated, Medroxyprogesterone Acetate)	C		
				Zolpidem Tartrate (Zolpidem Tartrate)	C		
				Prochlorperazine Edisylate (Prochlorperazine Edisylate)	C		
				Ondansetron Hydrochloride (Ondansetron Hydrochloride)	C		
				Oxcarbazepine (Oxcarbazepine)	C		
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		

Date:03/07/05ISR Number: 4602656-7Report Type:Expedited (15-DaCompany Report #2005034856

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 25 MG (25 MG,		Back Injury	Consumer	Zoloft (Sertraline)	PS		ORAL

1 IN 1 D),
 Depression
 ORAL
 Dizziness
 Headache
 Mania
 Neurontin
 (Gabapentin) SS ORAL
 900 MG (300
 Nephrolithiasis
 MG, 3 IN 1
 Pain In Extremity
 D), ORAL
 Post Procedural
 Complication
 Road Traffic Accident
 Stress

Date:03/07/05ISR Number: 4602661-0Report Type:Expedited (15-DaCompany Report #2005024066
 Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin (Gabapentin)	PS		
				Zoloft (Sertraline)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/07/05ISR Number: 4603038-4Report Type:Expedited (15-DaCompany Report #2005037014

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Radius Fracture Ulna Fracture	Foreign Health Professional	Gabapentin (Gabapentin)	PS		

Date:03/08/05ISR Number: 4601375-0Report Type:Expedited (15-DaCompany Report #200510782FR

Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - SUBCUTANEOUS Initial or Prolonged		Hepatitis		Lovenox	PS		
				Neurontin Atarax	SS		ORAL
				/Can/ Rivotril	C C		ORAL ORAL
				Cortancyl 5 Mg	C		ORAL

Date:03/08/05ISR Number: 4603353-4Report Type:Expedited (15-DaCompany Report #2005028590

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Dysphagia	Foreign	Gabapentin			
Other		Parkinson'S Disease	Health	(Gabapentin)	PS		ORAL
ORAL			Professional	Sinemet (Carbidopa, Levodopa)	C		
				Carbamazepine (Carbamazepine)	C		

Date:03/08/05ISR Number: 4603740-4Report Type:Direct Company Report #CTU 242310

Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Coordination Abnormal
Initial or Prolonged

Trazodone PS
Gabapentin SS
Clonazepam SS

Date:03/08/05ISR Number: 4603753-2Report Type:Expedited (15-DaCompany Report #2005004260
Age: Gender:Unknown I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Completed Suicide	Health Professional	Neurontin (Gabapentin)	PS		

Date:03/08/05ISR Number: 4603755-6Report Type:Expedited (15-DaCompany Report #2005018142
Age:57 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Anger
Initial or Prolonged	Bruxism
Other	Condition Aggravated
	Disease Recurrence
	Influenza
	Irritable Bowel Syndrome
	Mood Altered

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23000 MG 1 IN

Cardiac Disorder

(Gabapentin)

PS

1 D

Cardiac Murmur

Chest Pain

Circulatory Collapse

Convulsion

Depression

Dizziness

Electrocardiogram

Abnormal

Emotional Disorder

Headache

Heart Rate Irregular

Loss Of Consciousness

Mental Disorder

Myocardial Infarction

Pain

Palpitations

Somatic Delusion

Suicidal Ideation

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Hospitalization - 0.125 MG PO Initial or Prolonged QD	Fall Hallucination Orthostatic Hypotension	Digoxin 0.125mg	PS	ORAL
600 MG PO BID		Gabapentin	SS	ORAL
+ 900 MG QHS		Isosorbide Dinitrate	SS	ORAL
10 MG PO TID				
+ 20 MG QHS		Carvedilol	SS	ORAL
6.25 MG PO BID				

Date:03/08/05ISR Number: 4604277-9Report Type:Direct Company Report #CTU 242241
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG QHS 0.25 TID PRN	Fall		Gabapentin Trazodone Clonazepam	PS SS SS		

Date:03/08/05ISR Number: 4612301-2Report Type:Periodic Company Report #2005UW01188
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Chest Pain Dizziness	Health Professional Other	Crestor Lantus Glucotrol Xl Metformin Lotensin Hct	PS SS SS SS SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Plavix	SS
Celebrex	SS
Ecotrin	SS
Isosorbide	SS
Neurontin	SS
Ssri	SS
Protonix	C

Date:03/09/05ISR Number: 4604063-XReport Type:Direct
 Age:50 YR Gender:Male I/FU:I

Company Report #CTU 242409

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash		Neurontin (Generic Form)	PS		
900 M TID							

Date:03/09/05ISR Number: 4604070-7Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 242417

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Gabapentin	PS		
600 MG TID							
Drug Effect Decreased Pharmaceutical Product Complaint							

Date:03/09/05ISR Number: 4604074-4Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 242397

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia		Gabapentin Tablet	PS		
600 MG BID							
		Blood Urine Present		Gabapentin Capsule	SS		
600 MG BID							
Constipation Vision Blurred Vomiting Projectile							

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2100 MG Other	Aphasia Asthenia	Consumer Health	Neurontin (Gabapentin)	PS		
100 MG (50 MG, 2 IN 1 D), ORAL	Contusion Crying Feeling Abnormal Headache Hemiparesis Loss Of Consciousness Myalgia Neuropathy Neuropathy Peripheral Paraesthesia Speech Disorder Systemic Lupus Erythematosis Tremor	Professional	Topiramate (Topiramate)	SS		ORAL
			Zolpidem Tartrate (Zolpidem Tartrate)	C		
			Tramadol Hydrochloride (Tramadol Hydrochloride)	C		
			Propacet (Detropropoxyphene Napsilate, Paracetamol)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/09/05ISR Number: 4606127-3Report Type:Expedited (15-DaCompany Report #2005025718

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Amnesia Convulsion Disease Recurrence Liver Function Test Abnormal	Consumer Health Professional	Phenytoin Suspension (Phenytoin Sodium) Neurontin Phn Capsules (Gabapentin)	PS SS		ORAL
900 MG (300 MG,3 IN 1 D), ORAL		Rash Generalised Swelling Face Treatment Noncompliance		Phenobarbital (Phenobarbital) Lamictal (Lamotrigine)	SS C		

Date:03/09/05ISR Number: 4606872-XReport Type:Expedited (15-DaCompany Report #2005038428

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion Haemorrhage	Consumer	Neurontin (Gabapentin) Norvasc (Amlodipine)	PS SS		

Date:03/09/05ISR Number: 4606917-7Report Type:Expedited (15-DaCompany Report #2005038774

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Coma Convulsion Encephalopathy Neurotoxicity Respiratory Failure	Literature	Gabapentin (Gabapentin)	PS		

Date:03/09/05ISR Number: 4606920-7Report Type:Expedited (15-DaCompany Report #2004106809
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury Mental Disorder Overdose	Consumer	Neurontin (Gabapentin)	PS		

Date:03/09/05ISR Number: 4606948-7Report Type:Expedited (15-DaCompany Report #2005039819
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (100 Other MG, 4 IN 1 D), ORAL		Anticonvulsant Drug Level Decreased Grand Mal Convulsion Pharmaceutical Product Complaint	Consumer	Neurontin (Gabapentin) Clopidogrel Sulfate (Clopidogrel Sulfate) Phenytoin Sodium (Phenytoin Sodium)	PS C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/09/05ISR Number: 4606950-5Report Type:Expedited (15-DaCompany Report #2005038787

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coma Convulsion Encephalopathy Neurotoxicity Respiratory Failure	Literature	Gabapentin (Gabapentin)	PS		

Date:03/09/05ISR Number: 4606954-2Report Type:Expedited (15-DaCompany Report #2005037725

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aura Body Temperature	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Increased Convulsion Drug Ineffective Pharmaceutical Product Complaint		Valproate Semisodium (Valproate Semisodium) Risperidone (Risperidone) Levothyroxine Sodium (Levothyroxine Sodium) Guanfacine Hydrochloride (Guanfacine Hydrochloride) Benztropeine (Benztropeine) Quetiapine Fumarate (Quetiapine Fumarate) Diazepam (Diazepam) Vitamins (Vitamins)	C C C C C C C C C C		

Date:03/09/05ISR Number: 4606981-5Report Type:Expedited (15-DaCompany Report #2005018142

Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other (600 MG),		Anger Disease Recurrence Influenza Irritable Bowel Syndrome	Consumer Health Professional	Neurontin (Tablets)(Gabapentin)			ORAL
ORAL		Mood Altered Muscle Spasms Nausea Neuralgia Pain Pain In Extremity Pharmaceutical Product Complaint Spinal Operation Trigeminal Neuralgia Trismus		Oxycodone Hydrochloride (Oxycodone Hydrochloride) Lansoprazole (Lansoprazole) Librax (Chlordiazepoxide Hydrochloride, Clidinium Bromide) Provella-14 (Estrogens Conjugated, Medroxyprogesterone Acetate) Zolpidem Tartrate			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Zolpidem Tartrate0 C
 Prochlorperazine
 Edisylate
 (Prochlorperazine
 Edisylate) C
 Ondasetron
 Hydrochloride
 (Ondansetron
 Hydrochloride0 C
 Oxcarbazepine
 (Oxcarbazepine) C
 Vicodin (Hydrocodone
 Bitartrate,
 Paracetamol) C

Date:03/09/05ISR Number: 4606985-2Report Type:Expedited (15-DaCompany Report #2005038793
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coma Convulsion Encephalopathy Neurotoxicity Respiratory Failure	Literature	Gabapentin (Gabapentin)	PS		

Date:03/09/05ISR Number: 4606986-4Report Type:Expedited (15-DaCompany Report #2005038782
 Age: Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coma Convulsion Encephalopathy Neurotoxicity Respiratory Failure	Literature	Gabapentin (Gabapentin)	PS		

Date:03/09/05ISR Number: 4607199-2Report Type:Expedited (15-DaCompany Report #2005038777
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged 600 MG (300 MG, 2 IN 1 D); UNKNOWN	Abasia Asthenia Dysarthria Dysphemia Myoclonus Neurotoxicity	Literature	Gabapentin (Gabapentin)	PS
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Date:03/09/05ISR Number: 4607205-5Report Type:Expedited (15-DaCompany Report #2004022999
Age: Gender:Male I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Disturbance In Attention	Consumer	Neurontin (Gabapentin)	PS		
		Ejaculation Disorder		Viagra (Sildenafil Citrate)			
		Erectile Dysfunction		(Sildenafil)	SS		
		Lyme Disease		Atorvastatin	C		
		Memory Impairment		Nefazodone			
		Reading Disorder		Hydrochloride	C		
		Triple Vessel Bypass Graft					

Freedom Of Information (FOI) Report

Acetylsalicylic Acid C

Date:03/09/05ISR Number: 4607242-0Report Type:Expedited (15-DaCompany Report #2005037025
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
ORAL		Dyskinesia		(Gabapentin)	PS		ORAL
		Muscle Spasms		Irbesartan	C		
		Nerve Root Compression		Atenolol	C		
		Post Procedural		Lansoprazole	C		
		Complication		Sertraline			
		Spinal Fusion Surgery		Hydrochloride	C		
		Surgical Procedure					
		Repeated					

Date:03/09/05ISR Number: 4608903-XReport Type:Expedited (15-DaCompany Report #2005028496
 Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Asthenia	Foreign	Neurontin (Tablets)			
600 MG (300		Diplopia	Consumer	(Gabapentin)	PS		ORAL
MG, 2 IN 1		Dizziness					
D), ORAL		Drug Withdrawal Syndrome					
		Feeling Abnormal		Carbamazepine			
		Gastric Dilatation		(Carbamazepine)	C		
		Headache		Baclofen (Baclofen)	C		
		Hearing Impaired		Calcium With Vitamin			
		Myalgia		D (Calcium			
		Nausea		Phosphate, Calcium			
		Nervousness		Sodium Lactate,			
		Somnolence		Ergocalciferol)	C		
		Tremor		Thiamine			
		Visual Disturbance		Hydrochloride			
		Vomiting		(Thiamine			
		Weight Increased		Hydrochloride)	C		
				Phenobarbital Sodium			

(Phenobarbital Sodium) C
Phenytoin C
(Phenytoin)

Date:03/10/05ISR Number: 4605709-2Report Type:Direct Company Report #CTU 242627
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required		Rash Pruritic		Gabapentin	PS		ORAL
300 MG TID PO Intervention to Prevent Permanent Impairment/Damage							

Date:03/10/05ISR Number: 4606722-1Report Type:Expedited (15-DaCompany Report #2005022019
Age:29 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
UNKNOWN	600 MG	Cystitis Depression (200	Consumer Health	Neurontin (Gabapentin)	PS		
MG, 3 IN 1		Diarrhoea	Professional				
D), UNKNOWN		Disturbance In Attention					
UNKNOWN	130 MG	Hallucination		Baclofen (Baclofen)	SS		
(UNKNOWN)		Hypotension					
		Insomnia		Paracetamol (Paracetamol)	C		
		Social Avoidant Behaviour		Ibuprofen (Ibuprofen)	C		
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		
				Ciprofloxacin (Ciprofloxacin)	C		

Date:03/10/05ISR Number: 4606809-3Report Type:Expedited (15-DaCompany Report #2005022478
Age:73 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective	Consumer	Neurontin	PS		ORAL
Other		Dyspnoea	Health	(Gabapentin)			
ORAL		Pain In Extremity	Professional	Amitriptyline Hydrochloride			
		Surgical Procedure		(Amitriptyline Hydrochloride)	SS		
		Repeated		Glipizide (Glipizide)	C		
		Triple Vessel Bypass		Amiodarone (Amiodarone)	C		
		Graft		Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Citalopram			

Hydrobromide	
(Citalopram	
Hydrobromide)	C
Simvastatin	
(Simvastatin)	C
Zolpidem Tartrate	
(Zolpidem Tartrate)	C
Furosemide	
(Furosemide)	C
Potassium	
(Potassium)	C

Date:03/10/05ISR Number: 4606982-7Report Type:Expedited (15-DaCompany Report #2005037725

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Aura	Consumer	Neurontin			
		Body Temperature		(Gabapentin)	PS		ORAL
ORAL		Increased		Valproate Semisodium			
		Convulsion		(Valproate			
		Drug Effect Decreased		Semisodium)	C		
				Risperidone			

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D), ORAL

Zolpidem Tartrate	
(Zolpidem Tartrate)	C
Alprazolam	
(Alprazolam)	C
Fluoxetine	
Hydrochloride	
(Fluoxetine	
Hydrochloride)	C
Simvastatin	
(Simvastatin)	C
Amithriptyline	
Hydrochloride	
(Amithriptyline	
Hydrochloride)	C
Ginko Biloba (Kinkgo	
Biloba)	C

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/10/05ISR Number: 4607054-8Report Type:Expedited (15-DaCompany Report #2005036506
 Age:76 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	2400 MG (800 MG, 3 IN 1 D), ORAL; 7-8 YEARS AGO	Blood Cholesterol Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Drug Effect Decreased					
		Gallbladder Operation					
		Inadequate Analgesia					
	2400 MG (800 MG,3 IN 1 D), ORAL	Insomnia Pharmaceutical Product Complaint		Gabapentin (Tablets) (Gabapentin)	SS		ORAL
				Verapamil Hydrochloride (Verapamil Hydrochloride)	C		
				Hydrochlorothiazide (Hydrochlorothiazide)	C		
				Estrogens Conjugated (Estrogens Conjugated)	C		
				Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotinamide, Lekovit Ca (Calcium Carbonate, Colecalciferol)	C		
				Omeprazole (Omeprazole)	C		
				Drug, Unspecified (Drug, Unspecified)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (100 Other MG, 4 IN 1 D), ORAL		Anticonvulsant Drug Level Below Therapeutic Grand Mal Convulsion Medication Error Pharmaceutical Product Complaint	Consumer	Neurontin (Gabapentin) Clopidogrel Sulfate (Clopidogrel Sulfate) Phenytoin Sodium (Phenytoin Sodium)	PS C C		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Gun Shot Wound Hallucination, Auditory Intentional Self-Injury Limb Traumatic Amputation	Consumer Health Professional Company Representative	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/10/05ISR Number: 4607632-6Report Type:Expedited (15-DaCompany Report #2005038107
 Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300 MG (300 MG, 1 IN 1 D), ORAL	Drug Ineffective	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:03/10/05ISR Number: 4607860-XReport Type:Expedited (15-DaCompany Report #2005019189
 Age:18 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 3600 MG (1200 MG, 3 IN 1 D), ORAL	Initial or Prolonged	Drug Rash With Eosinophilia And Systemic Symptoms	Foreign Health Professional Company	Neurontin (Gabapentin)	PS		ORAL
1200 MG (400 MG, 3 IN 1 D), ORAL		Lymphocyte Morphology Abnormal Lymphocyte Transformation Test Positive Skin Test Positive	Representative	Carbamazepine (Carbamazepine) (Carbamazepine)	SS		ORAL

Date:03/11/05ISR Number: 4608267-1Report Type:Expedited (15-DaCompany Report #US014711
 Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
BUCCAL BUCCAL	1600 UG PRN	Communication Disorder Disorientation	Foreign Study	Fentanyl Citrate	PS		

600 MG DAILY	Drug Toxicity	Health	Gabapentin	SS	
80 MG DAILY		Professional	Oxycontin	SS	
30 MG DAILY		Other	Oxynorm	SS	
			Paracetamol	C	
			Dexamethazone-Pix	C	
			Lansoprazole	C	
			Laxoberal	C	Boehringer Ingelheim
			Capecitabine	C	

Date:03/11/05ISR Number: 4608424-4Report Type:Direct Company Report #CTU 242913
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Condition Aggravated		Gabapentin Capsules	PS	Teva	ORAL
600 MG		Drug Effect Decreased					
(2X300MG) PO		Grand Mal Convulsion					
TID		Headache		Glucosamine	C		
		Muscle Spasms		Chondroitin	C		
		Pharmaceutical Product					
		Complaint					
		Tongue Biting					

Date:03/11/05ISR Number: 4608857-6Report Type:Expedited (15-DaCompany Report #2005039751
 Age:83 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Faecal Incontinence
Initial or Prolonged	Loss Of Consciousness
	Oedema Peripheral

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Vomiting

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
THREE TIMES DAILY, ORAL		Foreign Health Professional	Neurontin (Gabepentin)	PS		ORAL
			Furosemide (Furosemide)	C		
			Digoxin (Digoxin)	C		
			Amlodipine Besilate (Amlodipine Besilate)	C		
			Warfarin Sodium (Warfarin Sodium)	C		
			Isosorbide Dinitrate (Isosorbide Dinitrate)	C		
			Valsartan (Valsartan)	C		
			Calcium Carbonate (Calcium Carbonate)	C		
			Selegiline Hydrochloride (Selegiline Hydrochloride)	C		
			Risedronate Sodium (Risedronate Sodium)	C		
			Asasantin (Acetylsalicylic Acid, Dipyridamole)	C		
			Acebutolol Hydrochloride (Acebutolol Hydrochloride)	C		

Date:03/11/05ISR Number: 4608943-0Report Type:Expedited (15-DaCompany Report #2004-121767-NL
Age:19 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 30 MG	2 DAY	Convulsion		Mirtazapine	PS		ORAL

Initial or Prolonged	Eye Rolling	Gabapentin	SS
300 MG BID/			
	Fall		
300 MG QD	180 DAY		
	Foaming At Mouth	Buspirone	
	Loss Of Consciousness	Hydrochloride	C
	Tremor	Fluoxetine	C
		Zopiclone	C

Date:03/11/05ISR Number: 4608988-0Report Type:Expedited (15-DaCompany Report #2004048907
Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abdominal Injury
Initial or Prolonged	Agitation
Other	Anger
	Anxiety
	Arterial Injury
	Atrial Fibrillation
	Biliary Tract Disorder
	Blood Calcium Decreased

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Freedom Of Information (FOI) Report

Dose	Duration	Condition Aggravated	Report Source	Product	Role	Manufacturer	Route
2000 MG (400 MG, 5 IN 1 D), UNKNOWN		Chest Injury Chest Pain Depression Diaphragmatic Injury Drug Screen Positive Gun Shot Wound Haematocrit Decreased Haemoglobin Decreased Haemothorax Hepatic Trauma Hypotension Insomnia Intercostal Neuralgia Nephrogenic Diabetes Insipidus Obsessive-Compulsive Disorder Pain Pericardial Effusion Pleural Effusion Pneumatois Cystoides Intestinalis Pneumonia Pneumothorax Polytraumatism Postoperative Wound Complication Pulmonary Embolism Respiratory Failure Suicide Attempt Traumatic Shock Venous Injury	Consumer	Neurontin (Gabapentin)	PS		
				Cefalexin (Cefalexin)	C		
				Warfarin Sodium (Warfarin Sodium)	C		
				Phenylpropanolamine (Phenylpropanolamine)	C		
				Cyproheptadine Hydrochloride (Cyproheptadine Hydrochloride)	C		
				Yohimbine (Yohimbine)	C		
				Cetirizine Hydrochloride (Cetirizine Hydrochloride)	C		
				Adapalene (Adapalene)	C		
				Meloxicam (Meloxicam)	C		
				Aripiprazole (Aripiprazole)	C		
				Ketorolac Tromethamine (Ketorolac Tromethamine)	C		
				Isocom (Dichloralphenazone, Isometheptene Mucate, Paracetamol)	C		
				Clonazepam (Clonazepam)	C		
				Olanzapine (Olanzapine)	C		
				Zolpidem Tartrate			

(Zolpidem Tartrate)	C
Fluoxetine	
Hydrochloride	
(Fluoxetine	
Hydrochloride)	C
Raloxifene	
Hydrochloride	
(Raloxifene	
Hydrochloride)	C
Lithium Carbonate)	C
Tramadol	
Hydrochloride	
(Tramadol	
Hydrochloride)	C
Atenolol (Atenolol)	C
Trazodone	
(Trazodone)	C
Propacet	

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(Dextropropoxyphene Napsilate, Paracetamol)	C
Estratest Hs (Estrogens Esterified, Methyltestosterone)	C
Estradiol (Estradiol)	C
Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C
Nefazodone Hydrochloride (Nefazodone Hydrochloride)	C
Zolmitriptan (Zolmitriptan)	C
Sodium Fluoride (Sodium Fluoride)	C
Amoxicillin (Amoxicillin)	C
Meperidine W/Promethazine (Pethidine, Promethazine)	C
Azithromycin (Azithromycin)	C
Aquatab C (Dextromethorphan, Guaifenesin Phenylpropanolamine)	C
Prednisone (Prednisone)	C
Pravastatin Sodium (Pravastatin Sodium)	C
Clindamycin (Clindamycin)	C
Chlorhexidine (Chlorhexidine)	C
Vicodin (Hydrocodone Bitartrate, Paracetamol)	C
Docusate Sodium (Docusate Sodium)	C
Risperidone (Risperidone)	C

Mirtazapine (Mirtazapine)	C
Bupropion Hydrochloride (Bupropion Hydrochloride)	C
Propranolol Hydrochloride (Propranolol Hydrochloride)	C
Sumatriptan Succinate (Sumatriptan Succinate)	C

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Methocarbamol	
(Methocarbamol)	C
Naproxen (Naproxen)	C
Narine Repetabs	
(Loratadine,	
Pseudoephedrine	
Sulfate)	C
Temazepam	
(Temazepam)	C
Quetiapine Fumarate	
(Quetiapine	
Fumarate)	C
Robitussin A-C/Old	
Form/ (Codeine	
Phosphate,	
Guaifenesin,	
Pheniramine Maleate)	C
Flurazepam	
Hydrochloride	
(Flurazepam	
Hydrochloride)	C
Mometasone Furoate	
(Mometasone Furoate)	C
Adapalene	
(Adapalene)	C
Ronatic	
(Chlorphenamine	
Tannate, Mepyramine	
Tannate,	
Phenylephrine	C
Modafinil	
(Modafinil)	C
Tizanidine	
Hydrochloride	
(Tizanidine	
Hdyrochloride)	C
Estrogens Conjugated	
(Estrogens	
Conjugated)	C
Estradiol	
(Estradiol)	C
Tiagabine	
Hydrochloride	
(Tiagabine	
Hydrochloride)	C
Sildenafil Citrate	
(Sildenafil Citrate)	C

Butorphanol Tartrate
(Butorphanol
Tartrate) C
Trimethobenzamide
Hydrochloride
(Trimethobenzamide
Hydrochloride) C
Lithium Carbonate
(Lithium Carbonate) C
Valproate Semisodium
(Valproate
Semisodium) C
Beclometasone
Dipropionate

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(Beclometasone
Dipropionate) C
Clavulin
(Amoxicillin
Trihydrate,
Clavulanate
Potassium) C

Date:03/11/05ISR Number: 4609018-7Report Type:Expedited (15-DaCompany Report #2005024130
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Pain	Consumer	Neurontin			
Other		Bedridden		(Gabapentin)	PS		
		Breast Pain		Gabapentin			
(600 MG)		Chest Pain		(Gabapentin)	SS		
		Drug Ineffective					
		Neck Pain					
		Pain					
		Pain In Extremity					
		Paraesthesia					
		Pharmaceutical Product					
		Complaint					

Date:03/11/05ISR Number: 4609036-9Report Type:Expedited (15-DaCompany Report #2004016191
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abdominal Pain	Health	Neurontin			
Initial or Prolonged		Abdominal Pain Upper	Professional	(Gabapentin)			
Disability		Agitation		(Tablets)	PS		ORAL
1800 MG (600		Anxiety					
Other		Ascites					
MG, 1 IN 3		Crying		Diltiazem			
D), ORAL		Diabetes Mellitus		Hydrochloride			
		Dyspnoea		(Diltiazem			
		Emotional Disorder		Hydrochloride)	C		
		Fatigue		Benazepril			

Hepatic Cirrhosis	Hydrochloride	
Hepatic Steatosis	(Benazepril	
Hepatomegaly	Hydrochloride)	C
Herpes Zoster	Hydrochlorothiazide	
Homicidal Ideation	(Hydrochlorothiazide	
Intestinal Obstruction)	C
Nervousness	Insulin (Insulin)	C
Simple Partial Seizures	Insulin Glargine	
Stress	(Insulin Glargine)	C
Tremor		

Date:03/11/05ISR Number: 4609140-5Report Type:Expedited (15-DaCompany Report #2005040445
Age:20 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disease Recurrence	Consumer	Neurontin (Tablets)			
		Petit Mal Epilepsy		(Gabapentin)	PS		ORAL
4800 MG (800							
MG, 6 IN 1 D)							
ORAL							
				Insulin	C		
				Lamotrigine	C		

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Freedom Of Information (FOI) Report

Date:03/11/05ISR Number: 4609141-7Report Type:Expedited (15-DaCompany Report #2005040468

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Neurontin (Gabapentin)	PS		

Date:03/11/05ISR Number: 4609158-2Report Type:Expedited (15-DaCompany Report #2005040440

Age:14 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	400 MG (100 MG, 4 IN 1 D), ORAL	Bipolar Disorder - Epilepsy	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Medication Error					
		Pharmaceutical Product					
		Complaint		Risperidone (Risperidone)	C		
				Clonidine (Clonidine)	C		

Date:03/11/05ISR Number: 4609163-6Report Type:Expedited (15-DaCompany Report #2005039629

Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300 MG (100 MG, 3 IN 1 D), ORAL	Condition Aggravated Pulmonary Hypertension	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Furosemide (Furosemide)	C		
				Warfarin Sodium (Wafarin Sodium)	C		
				Alprazolam (Alprazolam)	C		

Date:03/11/05ISR Number: 4609187-9Report Type:Expedited (15-DaCompany Report #2004067531
Age:41 YR Gender:Male I/FU:F

Outcome	PT
Death	Accidental Death
Hospitalization -	Agitation
Initial or Prolonged	Anxiety
Disability	Arthralgia
Other	Aspiration
	Asthenia
	Atherosclerosis
	Brain Oedema
	Burning Sensation
	Cardio-Respiratory Arrest
	Cardiomegaly
	Coma
	Coronary Artery
	Atherosclerosis
	Discomfort
	Disease Recurrence
	Drug Ineffective
	Drug Interaction
	Drug Level Above

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Dose	Duration	Therapeutic Drug Screen Positive Erectile Dysfunction Excoriation	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, 2 IN 1 D), ORAL		Fall Haematemesis	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Hepatic Steatosis	Professional				
		Hypertensive Heart Disease		Benadryl (Diphenhydramine)	SS		ORAL
ORAL		Hypoaesthesia					
1-3 TABLETS		Intentional Self-Injury Intervertebral Disc Protrusion Joint Injury		Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		ORAL
Q12H, ORAL		Laceration					
1-2 EVERY 4-6 HOURS (10 MG), ORAL		Lumbar Radiculopathy Major Depression		Hydrocodone (Hydrocodone)	SS		ORAL
		Malaise					
		Medical Device Complication		Zolpidem Tartrate (Zolpidem Tartrate)	SS		ORAL
10 MG (10 MG, 1 IN 1 D), ORAL		Mood Swings					
		Muscle Spasms					
		Muscle Spasticity					
80 MG (80 MG, 1 IN 1 D), ORAL		Musculoskeletal Pain Neck Pain Neuropathic Pain Pain		Citalopram Hydrobromide (Citalopram Hydrobromide)	SS		ORAL
		Pain In Extremity					
		Paraesthesia					
		Post Procedural Complication Pseudarthrosis		Amitriptyline Hydrochloride (Amitriptyline			

Pulmonary Oedema	Hydrochloride)	C
Renal Cyst	Fentanyl (Fentanyl)	C
Respiratory Depression	Cyclobenzaprine	
Sleep Disorder	Hydrochloride	
Tearfulness	(Cyclobenzaprine	
Therapy Non-Responder	Hydrochloride)	C
Vomiting	Bupropion	
	Hydrochloride	
	(Bupropion	
	Hydrochloride)	C
	Quinine Sulfate	
	(Quinine Sulfate)	C
	Sertraline	
	Hydrochloride	
	(Sertraline	
	Hydrochloride)	C
	Trazodone	
	(Trazodone)	C
	Hydroxyzine Embonate	
	(Hydroxyzine	
	Embonate)	C
	Carisoprodol	
	(Carisoprodol)	C
	Paroxetine	
	Dhydrochloride	
	(Paroxetine	
	Dhydrochloride)	C

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Freedom Of Information (FOI) Report

Date:03/11/05ISR Number: 4679216-5Report Type:Direct
Age:47 YR Gender:Female I/FU:I

Company Report #CTU 242677

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disturbance In Attention		Neurontin	PS		ORAL
TWO BID PO		Eye Swelling Memory Impairment Sleep Disorder Thinking Abnormal					

Date:03/11/05ISR Number: 4679440-1Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 242995

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Neurontin 300mg	PS		ORAL
300MG ONE PO		Therapeutic Response					
BID		Unexpected With Drug Substitution		Neurontin (Generic)	C		

Date:03/11/05ISR Number: 4679528-5Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 243008

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased		Neurontin (Generic)	PS		ORAL
300 MG TID PO		Pharmaceutical Product Complaint					

Date:03/11/05ISR Number: 4679626-6Report Type:Direct
Age:67 YR Gender:Male I/FU:I

Company Report #CTU 243035

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Urticaria		Gabapentin 300mg	PS		

Date:03/11/05ISR Number: 4679628-XReport Type:Direct Company Report #CTU 243036
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Sleep Disorder		Neurontin	PS		

Date:03/11/05ISR Number: 4693200-7Report Type:Direct Company Report #CTU 243048
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharmaceutical Product		Gabapentin	PS		
300 MG 1 PO		Complaint					
TID AND 3 AT							
BEDTIME							

Date:03/11/05ISR Number: 4693476-6Report Type:Direct Company Report #CTU 242695
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Ineffective		Neurontin 300 Mg /			
Other		Neck Pain		Tid	PS		
1 TID		Pharmaceutical Product					
		Complaint					

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Freedom Of Information (FOI) Report

Date:03/14/05ISR Number: 4606962-1Report Type:Expedited (15-DaCompany Report #GB-JNJFOC-20041007637

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Neutropenia		Risperdal	PS		
OROPHARINGEAL	1.5 MG	bd per					
source							
documents	21	DAY					
OROPHARINGEAL	100 MG	per		Risperdal	SS		
source							
documents	21	DAY					
OROPHARINGEAL				Clozapine	SS		
				Ranitidine	SS		
OROPHARINGEAL				Gabapentin	SS		
				Lamotrigine	SS		
OROPHARINGEAL	25-50 MG						

Date:03/14/05ISR Number: 4609363-5Report Type:Expedited (15-DaCompany Report #2004097068

Age:53 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Agranulocytosis	Foreign	Gabapentin			
Initial or Prolonged		Lymphadenopathy	Health	(Gabapentin)	PS		
UNKNOWN	900 MG	(300					
		Neutropenic Sepsis	Professional				
MG, 3 IN 1		Upper Respiratory Tract					
D), UNKNOWN		Infection		Therapeutic			
		White Blood Cell Count		Radiopharmaceuticals			
		Decreased		(Therapeutic			
				Radiopharmaceuticals			
)	C		
				Lansoprazole			
				(Lansoprazole)	C		
				Morphine Sulfate			

(Morphine Sulfate) C
Senna (Senna) C

Date:03/14/05ISR Number: 4609560-9Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 243178

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness		Gabapentin	PS		
300MG 3X		Headache					
DAILY							
		Pharmaceutical Product					
		Complaint					

Date:03/14/05ISR Number: 4609566-XReport Type:Direct
Age:66 YR Gender:Female I/FU:I

Company Report #CTU 243185

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nausea		Neurontin	PS		ORAL
900 MG TID PO							
		Pharmaceutical Product					
		Complaint					
		Vomiting					

Date:03/14/05ISR Number: 4609589-0Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 243060

Outcome	PT
Other	Feeling Abnormal
	Headache

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		Pharmaceutical Product Complaint	Report Source	Product	Role	Manufacturer	Route
Dose	Duration			Gabapentin 300mg	PS		
3 BID							

Date:03/14/05ISR Number: 4609909-7Report Type:Direct Company Report #CTU 243151
 Age:58 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged	Mental Status Changes		Methadone Gabapentin Cyclobenzaprine Citalopram Buspirone Quetiapine Etodolac Carisoprodol Zocor	PS SS SS C C C C C C		

Date:03/14/05ISR Number: 4610148-4Report Type:Expedited (15-DaCompany Report #2005041869
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Other	Drug Ineffective Pain Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:03/14/05ISR Number: 4610164-2Report Type:Expedited (15-DaCompany Report #001-0945-980292
 Age:51 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization - Initial or Prolonged 1200 MG (300 Other MG, 4 IN 1	Chronic Obstructive Pulmonary Disease Depression	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Required	Dysgeusia		
D), ORAL			
Intervention to	Dysuria	Carisoprodol	
Prevent Permanent	Epididymitis	(Carisoprodol)	C
Impairment/Damage	Headache	Hydrocodone	
	Influenza Like Illness	(Hydrocodone)	C
	Insomnia	Amitriptyline	
	Pain	(Amitriptyline)	C
	Pancreatitis	Fluoxetine	
	Pneumonia Mycoplasmal	Hydrochloride	
	Pneumonia Primary	(Fluoxetine	
	Atypical	Hydrochloride)	C
	Respiratory Failure		
	Swollen Tongue		
	Upper Respiratory Tract		
	Infection		

Date:03/14/05ISR Number: 4610291-XReport Type:Expedited (15-DaCompany Report #2005040143
Age:18 YR Gender:Male I/FU:I

Outcome	PT
Other	Hyperplasia
	Movement Disorder
	Muscle Spasms

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Muscular Weakness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2400 MG (600 MG, 4 IN 1 D), ORAL		Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Atomoxetine Hydrochloride (Atomoxetine Hydrochloride) Methylphenidate Hydrochloride (Methylphenidate Hydrochloride)	C		
				C		

Date:03/14/05ISR Number: 4610293-3Report Type:Expedited (15-DaCompany Report #2005038137
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	900 MG (4 IN 1 D), ORAL	Complex Regional Pain Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Insomnia					
	800 MG (400 MG, 2 IN 1 D), ORAL	Medication Error		Celebrex (Celecoxib)	SS		ORAL
		Pain In Extremity					
		Pharmaceutical Product					
		Complaint Spinal Osteoarthritis Wrist Fracture		Clonazepam (Clonazepam) Donepezil Hydrochloride (Donepezil Hydrochloride) Folic Acid (Folic Acid) Atorvastatin	C		C
					C		
					C		

(Atorvastatin) C
Estradiol
(Estradiol) C

Date:03/14/05ISR Number: 4610299-4Report Type:Expedited (15-DaCompany Report #2004074283
Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Balance Disorder	Consumer	Neurontin			
Initial or Prolonged	Deafness Unilateral	Health	(Gabapentin)	PS		
Other	Ear Pain	Professional				
	Inner Ear Disorder					
	Screaming					
	Sleep Disorder					
	Tinnitus					

Date:03/14/05ISR Number: 4610301-XReport Type:Expedited (15-DaCompany Report #2005014440
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Other	Neuropathy	Consumer	Neurontin			
	Pain In Extremity		(Gabapentin)	PS		
			Venlafaxine			

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Hydrochloride (Venlafaxine Hydrochloride)	SS
Oxycocet (Oxycodone Hydrochloride, Paracetamol)	SS
All Other Therapeutic Products (All Other Therapeutic Products)	C

Date:03/14/05ISR Number: 4610303-3Report Type:Expedited (15-DaCompany Report #2005040451
Age:65 YR Gender:Female I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG (100 MG, 1 IN 1 D), ORAL		Atrial Fibrillation Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG (100 MG, 1 IN 1 D), ORAL		Lung Neoplasm Malignant		Gabapentin (Gabapentin)	SS		ORAL
				Clonazepam (Clonazepam)	C		
				Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
				Glycopyrronium Bromide (Glycopyrronium Bromide)	C		
				Paracetamol (Paracetamol)	C		

Date:03/15/05ISR Number: 4610939-XReport Type:Direct Company Report #CTU 243246
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Amnesia Somnolence		Oxycontin Neurontin	PS SS		

Date:03/15/05ISR Number: 4610988-1Report Type:Direct Company Report #CTU 243245
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness Drug Intolerance		Gabapentin Generic Brand	PS		ORAL
100MG TID		Nausea					
ORAL		Pharmaceutical Product Complaint					

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Date:03/15/05ISR Number: 4613213-0Report Type:Expedited (15-DaCompany Report #2005040668
 Age:82 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 900 MG (300 MG, 3 IN 1 D), UNKNOWN	Agitation Confusional State Drug Interaction Hallucination Nightmare	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Amitriptyline Hydrochloride (Amitriptyline Hydrochloride) Fluindione (Fluindione) Cibenzoline Succinate (Cibenzoline Succinate) Simvastatin (Simvastatin)	PS SS C C C		

Date:03/15/05ISR Number: 4614168-5Report Type:Expedited (15-DaCompany Report #2005042034
 Age:69 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 200 MG (100 MG, 2 IN 1 Other D), ORAL	Apnoea Blood Disorder Brain Death Haematochezia Infection Multi-Organ Failure Pneumonia	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:03/15/05ISR Number: 4614170-3Report Type:Expedited (15-DaCompany Report #2005042017
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain	Consumer	Neurontin			
200 MG (200		Initial Insomnia		(Gabapentin)	PS		ORAL
MG, 1 IN 1		Intervertebral Disc					
D), ORAL		Degeneration					
		Spinal Disorder		Trazodone			
				(Trazodone)	C		
				Clonazepam			
				(Clonazepam)	C		
				Morphine (Morphine)	C		
				All Other			
				Therapeutic Products	C		

Date:03/15/05ISR Number: 4614174-0Report Type:Expedited (15-DaCompany Report #2005041264
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Consumer	Neurontin			
Other		Drug Ineffective		(Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:03/15/05ISR Number: 4614195-8Report Type:Expedited (15-DaCompany Report #2005040464

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (400 MG, 3 IN 1 D), ORAL		Abdominal Distension Amnesia Balance Disorder Chest Pain Confusional State Constipation Contusion Deafness Dizziness Dry Mouth Dyspnoea Ecchymosis Fatigue Haemorrhage Heart Rate Increased Hepatocellular Damage Hyperventilation Insomnia Mood Altered Nausea Oedema Peripheral Renal Disorder Somnolence Throat Tightness Tremor Vision Blurred	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Carbamazepine (Carbamazepine)	C		

Date:03/15/05ISR Number: 4614207-1Report Type:Expedited (15-DaCompany Report #2005014989

Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 900 MG (300 MG, 3 IN 1		Amnesia Balance Disorder Cervical Spinal Stenosis	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

D), ORAL	Depression			
	Erectile Dysfunction		Vicodin (Hydrocodone	
	Hypertension		Bitartrate,	
	Hypoaesthesia		Paracetamol)	SS
ORAL				ORAL
	Pain		Pantoprazole	
	Spinal Fusion Surgery		(Pantoprazole)	C
			Nabumetone	
			(Nabumetone)	C
			Potassium Chloride	
			(Potassium Chloride)	C
			Furosemide	
			(Furosemide)	C
			Tizanidine	
			Hydrochloride	
			(Tizanidine	
			Hydrochloride)	C

Other 900MG TID PO Nausea Neurontin 900mg PS ORAL

Pharmaceutical Product
Complaint
Vomiting

Date:03/16/05ISR Number: 4612572-2Report Type:Direct Company Report #CTU 243373
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Pharmaceutical Product Complaint Urticaria		Neurontin Generic	PS		

Date:03/16/05ISR Number: 4612658-2Report Type:Expedited (15-DaCompany Report #2005041846
Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Meningitis Chemical Rash	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other				Metoprolol (Metoprolol)	C		
				Carbamazepine (Carbamazepine)	C		

1000 MG (2 IN

1 DS), ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/16/05ISR Number: 4612660-0Report Type:Expedited (15-DaCompany Report #2005040889
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG (200 MG, 1 IN 1 D), ORAL		Back Disorder	Health	Celebrex (Celecoxib)	PS		ORAL
		Convulsion	Professional				
		Dysstasia					
900 MG (300 MG, 3 IN 1 D), ORAL		Muscle Spasms Oedema Peripheral		Neurontin (Gabapentin)	SS		ORAL
		Pharmaceutical Product					
		Complaint					
(300 MG), ORAL		Stool Analysis Abnormal Tremor		Gabapentin (Gabapentin)	SS		ORAL
		Urinary Incontinence					
		Weight Decreased		Esomeprazole (Esomeprazole)	C		
				Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
				Oxycocet (Oxycodone Hydrochloride, Paracetamol)	C		
				Furosemide (Furosemide)	C		
				Clonazepam (Clonazepam)	C		
				Diclofenac Sodium (Diclofenanc Sodium)	C		

Date:03/16/05ISR Number: 4612661-2Report Type:Expedited (15-DaCompany Report #2005040437
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Complex Regional Pain	Consumer	Neurontin			

300 MG (100	Syndrome	(Gabapentin)	PS	ORAL
MG, 3 TIMES	Condition Aggravated			
DAILY AS	Drug Ineffective			
NEEDED), ORAL	Gallbladder Operation			
	Pharmaceutical Product Complaint	Vitamin B (Vitamin B)	C	

Date:03/16/05ISR Number: 4612756-3Report Type:Expedited (15-DaCompany Report #2005032411
Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arterial Disorder	Consumer	Celebrex (Celecoxib)	PS		ORAL
Other		Drug Effect Decreased		Bextra (Valdecoxib)	SS		ORAL
ORAL		Myocardial Infarction		Neurontin (Gabapentin)	SS		ORAL
ORAL				Cyclobenzaprine Hydrochloride (Cyclobenzaprine Hydrochloride)	C		
				Lidocaine Hydrochloride (Lidocaine Hydrochloride)	C		
				Oxycodone Hydrochloride			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Oxycodone
Hydrochloride) C

Date:03/16/05ISR Number: 4612757-5Report Type:Expedited (15-DaCompany Report #2005041242
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin (Gabapentin)	PS		

Date:03/16/05ISR Number: 4613177-XReport Type:Expedited (15-DaCompany Report #2005042305
Age:79 YR Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 200 MG (100 MG, 2 IN 1 D), ORAL		Depression Drug Effect Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL

Diltiazem
Hydrochloride
(Diltiazem
Hydrochloride) C
Furosemide
(Furosemide) C
Spironolactone
(Spironolactone) C
Atenolol (Atenolol) C
Warfarin Sodium
(Warfarin Sodium) C
Tolterodine
L-Tartrate
(Tolterodine
L-Tartrate) C
Potassium Chloride
(Potassium Chloride) C
Calcium Carbonate
(Calcium Carbonate) C
All Other
Therapeutic Products

(All Other
 Therapeutic
 Products) C
 Multivitamins
 (Ascorbic Acid,
 Ergocalciferol,
 Folic Acid,
 Nicotinamide, C

Date:03/16/05ISR Number: 4613178-1Report Type:Expedited (15-DaCompany Report #2005042101
 Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Brain Operation Convulsion	Consumer	Neurontin (Gabapentin)	PS		
600 MG (100 MG, 6 IN 1 D)		Ill-Defined Disorder		All Other Therapeutic Products (All Other Therapeutic			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Products) C
 Primidone
 (Primidone) C
 Lamotrigine
 (Lamotrigine) C

Date:03/16/05ISR Number: 4613188-4Report Type:Expedited (15-DaCompany Report #2005042048
 Age:63 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	200 MG (100 Other MG, 2 IN 1	Drug Ineffective Fatigue	Consumer	Neurontin (Gabapentin)	PS		ORAL
D), ORAL		Vascular Occlusion					

Metformin
 Hydrochloride
 (Metformin
 Hydrochloride) C
 Pioglitazone
 (Pioglitazone) C
 Clopidogrel Sulfate
 (Clopidogrel
 Sulfate) C
 Trazodone
 (Trazodone) C
 Atenolol (Atenolol) C
 Vitamins (Vitamins) C
 Prednisone
 (Prednisone) C

Date:03/16/05ISR Number: 4613343-3Report Type:Expedited (15-DaCompany Report #2005042029
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	100 MG (100 MG, 1 IN 1	Hyperchlorhydria Mood Altered	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
		Neutropenia	Professional				

D), ORAL

Clozapine
(Clozapine) SS ORAL

ORAL

Ranitidine
Hydrochloride
(Ranitidine
Hydrochloride) SS

150 MG (150
MG, 1 IN 1 D)

Risperidone
(Risperidone) SS ORAL

3 MG (1.5 MG,
2 IN 1

D), ORAL

Lamotrigine
(Lamotrigine) SS ORAL

ORAL

Date:03/16/05ISR Number: 4614208-3Report Type:Expedited (15-DaCompany Report #2005PK00415
Age:85 YR Gender:Female I/FU:I

Outcome	PT	Report Source
Hospitalization -	Drug Interaction	Foreign
Initial or Prolonged	Overdose	Health
	Stupor	Professional

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Other

Dose	Duration	Product	Role	Manufacturer	Route
100 MG DAILY		Beloc Zok	PS		ORAL
PO					
10 MG BID PO		Mst Continus	SS		ORAL
75 MG QID PO		Valoron	SS		ORAL
25 MG QD PO		Surmontil	SS		ORAL
600 MG TID PO		Neurontin	SS		ORAL
600 MG PO		Neurontin	SS		ORAL
30 MG QD PO		Agopton	SS		ORAL
		Marcoumar	C		
		Triatec	C		
		Norvasc	C		
		Selipran	C		
		Corvaton	C		
		Paracetamol	C		
		Calcium	C		
		Fosamax	C		
		Nitroderm	C		
		Torem	C		
		Heparin	C		
		Prednison	C		

Date:03/16/05ISR Number: 4624013-XReport Type:Direct
 Age:55 YR Gender:Female I/FU:I

Company Report #CTU 243452

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Gabapentin (Teva Pharm)	PS	(Teva Pharm)	ORAL
600MG ONE PO							
TID							

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Literature	Gabapentin			
		Intentional Misuse	Health	(Gabapentin)	PS		
		Respiratory Arrest	Professional	Cocaine (Cocaine)	SS		
				Ethanol (Ethanol)	SS		
				All Other Therapeutic Products			
				(All Other Therapeutic Products)	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Abuser	Literature	Sertraline			
		Intentional Misuse	Health	(Sertraline)	PS		
		Multiple Drug Overdose	Professional	Methadone Hydrochloride			
				(Methadone Hydrochloride)	SS		
				Gabapentin			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Gabapentin)
 (Gabapentin) SS
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) SS

Date:03/17/05ISR Number: 4615273-XReport Type:Expedited (15-DaCompany Report #2004070289
 Age:38 YR Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Methadone Hydrochloride (Methadone) (Methadone Hydrochloride)	SS		ORAL
ORAL				All Other Therapeutic Products (All Other Therapeutic Products)	SS		ORAL
ORAL				Fluoxetine (Fluoxetine)	SS		ORAL

Date:03/17/05ISR Number: 4615286-8Report Type:Expedited (15-DaCompany Report #2004070286
 Age:23 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest Drug Toxicity	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL		Intentional Misuse	Professional	Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		ORAL
ORAL							

Date:03/17/05ISR Number: 4615292-3Report Type:Expedited (15-DaCompany Report #2005042251

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Medication Error	Consumer	Neurontin			
		Pharmaceutical Product		(Gabapentin)	PS		ORAL
ORAL		Complaint					

Date:03/17/05ISR Number: 4615293-5Report Type:Expedited (15-DaCompany Report #2005042140

Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Bone Neoplasm Malignant	Consumer	Neurontin			
		No Therapeutic Response		(Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/17/05ISR Number: 4615299-6Report Type:Expedited (15-DaCompany Report #2005042021

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Neurontin			
(2 IN 1 D),		Coordination Abnormal		(Gabapentin)	PS		ORAL
ORAL		Depression					
				Valproate Semisodium			
				(Valproate			
				Semisodium)	SS		
				Bupropion			
				(Bupropion)	C		
				Quinapril			
				Hydrochloride			
				(Quinapril			
				Hydrochloride)	C		
				Venlafaxine			
				Hydrochloride			
				(Venlafaxine			
				Hydrochloride)	C		
				Estrogens Conjugated			
				(Estrogens			
				Conjugated)	C		
				Vitamins (Vitamins)	C		

Date:03/17/05ISR Number: 4615396-5Report Type:Expedited (15-DaCompany Report #2004070512

Age:92 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Medication Error	Literature	Gabapentin			
ORAL		Multiple Drug Overdose	Health	(Gabapentin)	PS		ORAL
		Accidental	Professional	Clozapine			
ORAL				(Clozapine)	SS		ORAL

Date:03/17/05ISR Number: 4615398-9Report Type:Expedited (15-DaCompany Report #2004070485

Age:19 YR Gender: I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide Intentional Misuse	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL				Professional	Baclofen (Baclofen)	SS		ORAL
ORAL					Oxybutynin (Oxybutynin)	SS		ORAL

Date:03/17/05ISR Number: 4615401-6Report Type:Expedited (15-DaCompany Report #2004070297
Age:44 YR Gender: I/FU:F

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death			Completed Suicide Intentional Misuse	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL				Professional	Valproic Acid (Valproic Acid)	SS		ORAL
ORAL					Levothyroxine (Levothyroxine)	SS		ORAL
ORAL					All Other Therapeutic Products (All Other			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

ORAL

Therapeutic Products) SS ORAL

Date:03/17/05ISR Number: 4615417-XReport Type:Expedited (15-DaCompany Report #2005042343
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG (100		Uterine Cyst					
MG, 2 IN 1		Uterine Leiomyoma					
D), ORAL		Weight Increased					
				Alprazolam (Alprazolam)	C		
				Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		

Date:03/17/05ISR Number: 4615419-3Report Type:Expedited (15-DaCompany Report #2005043855
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia	Consumer	Neurontin (Gabapentin)	PS		
		Eye Disorder					
		Pain					

Date:03/17/05ISR Number: 4615422-3Report Type:Expedited (15-DaCompany Report #2005042128
 Age:74 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blister	Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG (100		Blood Disorder					
MG), ORAL		Drug Ineffective					

200 MG (100	Metastases To Lung	Celebrex (Celecoxib)	SS	ORAL
MG, 2 IN 1	Neoplasm Malignant			
D), ORAL	Pain In Extremity			
	Pharmaceutical Product	Losartan Potassium		
	Complaint	(Losartan Potassium)	C	
	Post Procedural	Levothyroxine Sodium		
	Complication	(Levothyroxine		
	Radiation Skin Injury	Sodium)	C	
	Sarcoma	Amlodipine Besilate		
		(Amlodipine		
		Besilate)	C	
		Ranitidine		
		Hydrochloride		
		(Ranitidine		
		Hydrochloride)	C	
		Atenolol (Atenolol)	C	
		Furosemide		
		(Furosemide)	C	
		Pravastatin Sodium		
		(Pravastatin Sodium)	C	
		Ultracet		
		(Paracetamol,		
		Tramadol		
		Hydrochloride)	C	
		Clorazepate		
		Dipotassium		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Clorazepate
Dipotassium) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C

Date:03/17/05ISR Number: 4615425-9Report Type:Expedited (15-DaCompany Report #2004070293
Age:35 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardio-Respiratory Arrest	Literature	Gabapentin			
ORAL		Completed Suicide	Health	(Gabapentin)	PS		ORAL
ORAL		Multiple Drug Overdose	Professional	Tramadol (Tramadol)	SS		ORAL
ORAL				Citalopram (Citalopram)	SS		ORAL
ORAL				All Other Therapeutic Products	SS		ORAL

Date:03/17/05ISR Number: 4615426-0Report Type:Expedited (15-DaCompany Report #2005040472
Age:85 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchitis	Consumer	Neurontin(Gabapentin			
Hospitalization -		Diarrhoea)	PS		ORAL
ORAL		Urinary Incontinence		Propacet (Dextropropoxyphene Napsilate, Paracetamol)	C		

Date:03/17/05ISR Number: 4615427-2Report Type:Expedited (15-DaCompany Report #2005042008
Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Hospitalization - Initial or Prolonged ORAL	Compression Fracture Dyskinesia Hot Flush Hyperhidrosis Prostatic Disorder	Consumer	Neurontin (Gabapentin)	PS	ORAL
			Oxycodone (Oxycodone)	C	
			Zolpidem Tartrate (Zolpidem Tartrate)	C	
			Antibiotics (Antibiotics)	C	

Date:03/17/05ISR Number: 4615428-4Report Type:Expedited (15-DaCompany Report #2004070288
Age:45 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Multiple Drug Overdose	Literature Health	Gabapentin (Gabapentin)	PS		ORAL
ORAL			Professional	Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		ORAL
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/17/05ISR Number: 4615500-9Report Type:Expedited (15-DaCompany Report #2004080583
Age:88 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Fall	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
Life-Threatening		Hypotension					
300 MG (300 Hospitalization - MG, 1 IN 1 Initial or Prolonged D), ORAL		Renal Failure Acute Respiratory Failure	Professional Company Representative	Amiodarone (Amiodarone)	C		
				Carbocisteine (Carbocisteine)	C		
				Tiotropium Bromide (Tiotropium Bromide)	C		
				Candesartan (Candesartan)	C		
				Salbutamol (Salbutamol)	C		
				Beclometasone (Beclometasone)	C		
				Furosemide (Furosemide)	C		

Date:03/17/05ISR Number: 4615503-4Report Type:Expedited (15-DaCompany Report #2004056068
Age:25 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Exposure During Pregnancy	Foreign Health	Gabapentin (Gabapentin)	PS		
600 MG (600 MG, 1 IN 1 D)		Drug Interaction	Professional				
		Pregnancy With Injectable Contraceptive Unintended Pregnancy		Depo-Provera Suspension, Sterile (Medroxyprogesterone Acetate)			
INTRAMUSCULAR	(150 MG, 1 IN			(Medroxyprogesterone	SS		

INTRAMUSCULAR

C

Date:03/17/05ISR Number: 4615521-6Report Type:Expedited (15-DaCompany Report #2004070292
Age:49 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Literature	Gabapentin			
ORAL		Intentional Misuse	Health	(Gabapentin)	PS		ORAL
		Respiratory Arrest	Professional	Methadone			
				Hydrochloride			
				(Methadone)			
ORAL				(Methadone			
				Hydrochloride)	SS		ORAL
				Methocarbamol			
ORAL				(Methocarbamol)	SS		ORAL

Date:03/17/05ISR Number: 4615556-3Report Type:Expedited (15-DaCompany Report #2004070501
Age:50 YR Gender:Unknown I/FU:F

Outcome	PT	Report Source
Death	Completed Suicide	Literature
	Intentional Misuse	Health

Freedom Of Information (FOI) Report

Professional

Dose	Duration	Product	Role	Manufacturer	Route
ORAL		Gabapentin (Gabapentin)	PS		ORAL
ORAL		Chlorpromazine (Chlorpromazine)	SS		ORAL
ORAL		Quetiapine (Quetiapine)	SS		ORAL
ORAL		All Other Therapeutic Products (All Other Therapeutic Products)	SS		ORAL

Date:03/17/05ISR Number: 4615628-3Report Type:Expedited (15-DaCompany Report #L04-USA-07403-02
Age:53 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anoxic Encephalopathy Cerebral Infarction Coma Completed Suicide Dizziness Intentional Misuse Loss Of Consciousness Metabolic Acidosis Mydriasis	Literature Health Professional	Levothyroxine Ethylene Glycol (Polyethylene Glycol) Gabapentin Paroxetine Rofecoxib Trazodone Lovastatin Clonazepam Lorazepam Benazepril Lansoprazole Medroxyprogesterone Aspirin	PS SS SS SS SS SS SS SS SS SS SS SS SS		

Date:03/17/05ISR Number: 4615635-0Report Type:Expedited (15-DaCompany Report #L04-USA-07403-20
Age:44 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Multiple Drug Overdose	Literature Health Professional	Levothyroxine Valproic Acid Gabapentin	PS SS SS		

Date:03/17/05ISR Number: 4619591-0Report Type:Direct Company Report #CTU 243475
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening 600 MG QID Hospitalization - Initial or Prolonged		Convulsion		Neurontin	PS		

Date:03/17/05ISR Number: 4621015-4Report Type:Periodic Company Report #2005030999
 Age: Gender:Male I/FU:I

Outcome	PT
Other	Difficulty In Walking Pain In Extremity

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Polyneuropathy Tenderness					
Dose	Duration		Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, 2 IN 1 D)			Health Professional	Vfend (Voriconazole)	PS		
ORAL				Neurontin (Gabapentin)	SS		ORAL
				Vancomycin	C		
				Oxycodone	C		
				Amitriptyline	C		
Date:03/17/05 Age:70 YR Gender:Female I/FU:I			ISR Number: 4622579-7 Report Type:Periodic Company Report #PHEH2004US11279				

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dizziness Fall	Health Professional	Trileptal (Oxcarbazepine)	PS		ORAL
450 MG, BID, ORAL		Syncope		Neurontin (Gabapentin)	SS		
				Amitriptyline (Amitriptyline)	C		
				Atorvastatin (Atorvastatin)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Methotrexate (Methotrexate)	C		

Date:03/18/05 Age: Gender:Female I/FU:I			ISR Number: 4613430-X Report Type:Direct Company Report #CTU 243705				
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Drug Effect Decreased Gabapentin (Generic) PS
 400 TID AND Pain
 800 QHS Pharmaceutical Product
 Complaint

Date:03/18/05ISR Number: 4613635-8Report Type:Direct Company Report #CTU 243628
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - PO PRIOR Initial or Prolonged TO ADMISSION		Catatonia		Sinequan	PS		ORAL
PO PRIOR TO ADMISSION		Dysarthria					
		Fatigue		Klonopin	SS		ORAL
PO PRIOR TO ADMISSION		Staring					
		Treatment Noncompliance		Neurontin	SS		ORAL
PO PRIOR TO ADMISSION				Fioricet	SS		ORAL
				Ultram	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/18/05ISR Number: 4615026-2Report Type:Direct
 Age:42 YR Gender:Male I/FU:I

Company Report #CTU 243779

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
PO TID		Difficulty In Walking		Gabapentin 500mg	PS		ORAL
		Hypoaesthesia		Percocet	C		
				Restoril	C		

Date:03/18/05ISR Number: 4616427-9Report Type:Expedited (15-DaCompany Report #2005042083
 Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (100 Disability MG, 3 IN 1 Other D), ORAL		Cerebrovascular Accident Hemiparesis	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Clonidine (Clonidine)	C		
				Propranolol (Propranolol)	C		
				Glyceryl Trinitrate (Glyceryl Trinitrate)	C		
				Ezetimibe (Ezetimibe)	C		
				Alendronate Sodium (Alendronate Sodium)	C		
				Rosuvastatin (Rosuvastatin)	C		
				Aporex (Dextropropoxyphene Hydrochloride, Paracetamol)	C		
				Valproate Semisodium (Valproate Semisodium)	C		

Date:03/18/05ISR Number: 4616440-1Report Type:Expedited (15-DaCompany Report #2005041871
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin (Gabapentin)	PS		

Date:03/18/05ISR Number: 4616451-6Report Type:Expedited (15-DaCompany Report #2005042159
Age:94 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Abasia	Consumer	Neurontin (Gabapentin)	PS		ORAL
200 MG (100 MG, 2 DAILY), ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/18/05ISR Number: 4616495-4Report Type:Expedited (15-DaCompany Report #2005040445
 Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Petit Mal Epilepsy Pharmaceutical Product	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
4800 MG (800 MG, 6 IN 1 D), ORAL		Complaint					
				Insulin (Insulin) Lamotrigine (Lamotrigine)	C C		

Date:03/18/05ISR Number: 4616513-3Report Type:Expedited (15-DaCompany Report #2005042723
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death ORAL		Adverse Event Pneumonia	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:03/18/05ISR Number: 4616603-5Report Type:Expedited (15-DaCompany Report #2005042258
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Complex Partial Seizures Pharmaceutical Product	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Complaint		Phenytoin Sodium (Phenytoin Sodium) Lamotrigine (Lamotrigine) Lorazepam (Lorazepam) Fexofenadine Hydrochloride (Fexofenadine Hydrochloride)	C C C C		

Ipratropium Bromide
(Ipratropium
Bromide) C
Fluticasone
Propionate
(Fluticasone
Propionate) C
Pirbuterol Acetate
(Pirbuterol Acetate) C
Epinephrine
(Epinephrine) C

Date:03/18/05ISR Number: 4616605-9Report Type:Expedited (15-DaCompany Report #2005042471

Age:71 YR Gender:Female I/FU:I

Outcome PT
Other Deafness
Drug Ineffective
Endolymphatic Hydrops
Fall
Impaired Driving Ability
Pharmaceutical Product
Complaint
Post Procedural

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Complication

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, 3 IN 1 D), ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
			Levothyroxine Sodium (Levothyroxine Sodium)	C		
			Triamterene (Triamterene)	C		

Date:03/18/05ISR Number: 4616606-0Report Type:Expedited (15-DaCompany Report #2005042541
 Age:90 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	70 MG (3 IN 1 D), ORAL	Abasia Asthenia Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Dysstasia Essential Tremor Fall Movement Disorder		Terazosin (Terazosin)	C		
				Lisinopril (Lisinopril)	C		
				Simvastatin (Simvastatin)	C		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:03/18/05ISR Number: 4616611-4Report Type:Expedited (15-DaCompany Report #2005038011
 Age:75 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 200 MG (200 MG, 1 IN 1 D), ORAL		Buttock Pain	Consumer	Celebrex (Celecoxib)	PS		ORAL
		Condition Aggravated					
		Diabetic Neuropathy					
600 MG (1 IN 11D), ORAL		Diarrhoea Drug Ineffective		Neurontin (Gabapentin)	SS		ORAL
		Localised Oedema					
		Myalgia Post Procedural Pain		Pioglitazone (Pioglitazone)	C		

Date:03/18/05ISR Number: 4617233-1Report Type:Expedited (15-DaCompany Report #L04-USA-07403-12
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Arrest	Literature	Citalopram	PS		
		Coma	Health	Tramadol	SS		
		Completed Suicide	Professional	Gabapentin	SS		
		Convulsion		Metoclopramide	SS		
		Hypotension		Omeprazole	SS		
		Hypoxic Encephalopathy					
		Intentional Misuse					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/21/05ISR Number: 4616198-6Report Type:Expedited (15-DaCompany Report #2005041857
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		

Date:03/21/05ISR Number: 4616303-1Report Type:Expedited (15-DaCompany Report #2005042423
 Age:64 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Fear Feeling Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG (100 MG, 1 IN 1 D), ORAL		Headache Hypersomnia Ill-Defined Disorder Pharmaceutical Product Complaint Road Traffic Accident Thinking Abnormal		Clonazepam (Clonazepam) Quetiapine Fumarate (Quetiapine Fumarate)	C C		

Date:03/21/05ISR Number: 4616307-9Report Type:Expedited (15-DaCompany Report #2005042488
 Age:84 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aphagia Aphasia	Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL		Confusional State Disease Recurrence Drug Ineffective Inadequate Analgesia Medical Device Complication Neuralgia		Oxycocet (Oxycodone Hydrochloride, Paracetamol) Diltiazem (Diltiazem)	SS C		

Pharmaceutical Product
Complaint
Weight Decreased

Centrum (Minerals
Nos, Vitamins Nos) C

Date:03/21/05ISR Number: 4616310-9Report Type:Expedited (15-DaCompany Report #2005042482
Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL Other	Blindness Cerebrovascular Accident Drug Hypersensitivity Gastrointestinal Disorder Haemorrhage Hallucination, Visual Macular Degeneration Myocardial Infarction Photopsia Suicidal Ideation Visual Field Defect	Consumer	Neurontin (Gabapentin) Sertraline Hydrochloride (Sertraline Hydrochloride) Flurazepam (Flurazepam)	PS C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/21/05ISR Number: 4616390-0Report Type:Expedited (15-DaCompany Report #2005042679
 Age:74 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG (100 Other MG, 3 IN 1 D), ORAL	Cardiac Disorder Cardiac Failure Congestive Cardiac Valve Disease	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Methadone (Methadone)	C		
			Isosorbide (Isosorbide)	C		
			Digoxin (Digoxin)	C		
			Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		

Date:03/21/05ISR Number: 4616391-2Report Type:Expedited (15-DaCompany Report #2005042994
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG, ORAL	Gastric Cancer	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:03/21/05ISR Number: 4616392-4Report Type:Expedited (15-DaCompany Report #2005042717
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Lung Neoplasm Malignant Neoplasm Malignant Neoplasm Recurrence	Consumer	Neurontin (Gabapentin)	PS		

Date:03/21/05ISR Number: 4616402-4Report Type:Expedited (15-DaCompany Report #2005042984
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Literature	Gabapentin (Tablets)			
Other		Depressed Level Of Consciousness		(Gabapentin)	PS		
		Drug Toxicity		Metaxalone			
				(Metaxalone)	SS		
				Paracetamol			
				(Paracetamol)	C		
				Citalopram			
				(Citalopram)	C		
				Flurazepam			
				(Flurazepam)	C		

Date:03/21/05ISR Number: 4616408-5Report Type:Expedited (15-DaCompany Report #2005042698
Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Anaemia
Initial or Prolonged	Arterial Disorder
Other	Asthenia
	Coronary Artery Occlusion
	Dysplasia
	Gastric Haemorrhage
	Headache

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Nausea

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
100 MG (100 MG, 1 IN 1 D), ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL			Acetylsalicylic Acid (Acetylsalicylic Acid)	SS		ORAL
			Lansoprazole (Lansoprazole)	C		
			Diltiazem Hydrochloride (Diltiazem Hydrochloride)	C		
			Alprazolam (Alprazolam)	C		
			Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C		
			Tolterodine L-Tartrate (Tolterodine L-Tartrate)	C		
			Furosemide (Furosemide)	C		
			Atorvastatin (Atorvastatin)	C		
			Thiamine Hydrochloride (Thiamine Hydrochloride)	C		
			Paracetamol (Paracetamol)	C		
			Calcium (Calcium)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Choking Sensation Neuropathy Peripheral	Consumer	Gabapentin (Gabapentin)	PS		ORAL
200 MG (100 MG, 2 IN 1 D), ORAL		Throat Irritation		Vicodine (Hydrochloride Bitartrate, Paracetamol)	C		

Date:03/21/05ISR Number: 4616413-9Report Type:Expedited (15-DaCompany Report #2005043617
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Eye Disorder Eye Swelling	Consumer	Neurontin (Gabapentin)	PS		ORAL
500 MG (3 IN 1 D), ORAL		Vascular Rupture		Gabapentin	SS		ORAL
500 MG (3 IN		Visual Acuity Reduced					

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Freedom Of Information (FOI) Report

1 D), ORAL

Fexofenadine Hydrochloride (Fexofenadine Hydrochloride)	C
Diclofenac (Diclofenac)	C
Cetirizine Hydrochloride (Cetirizine Hydrochloride)	C
Benzonatate (Benzonatate)	C
Donepezil Hydrochloride (Donepezil Hydrochloride)	C
Propranolol Hydrochloride (Propranolol Hydrochloride)	C
Tiagabine Hydrochloride (Tiagabine Hydrochloride)	C
Risperidone (Risperidone)	C
Valsartan (Valsartan)	C
Rosiglitazone Maleate (Rosiglitazone Maleate)	C
Nortriptyline (Nortriptyline)	C
Insulin (Insulin)	C

Date:03/21/05ISR Number: 4616414-0Report Type:Expedited (15-DaCompany Report #2005042704

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Anaemia Cataract Operation	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (100							

MG, 6 IN 1

Dyspnoea

D), ORAL

Fatigue

Feeling Abnormal
Pharmaceutical Product
Complaint
Visual Disturbance

Furosemide
(Furosemide) C
Asasantin
(Acetylsalicylic
Acid, Dipyrindamole) C
Sertraline
Hydrochloride
(Sertraline
Hydrochloride) C
Amlodipine Besilate
(Amlodipine
Besilate) C
Zonisamide
(Zonisamide) C
Levothyroxine Sodium
(Levothyroxine

Freedom Of Information (FOI) Report

Sodium) C
 Losartan Potassium
 (Losartan Potassium) C
 Centrum Silver
 (Ascorbic Acid,
 Calcium, Minerals
 Nos, Retinol,
 Tocopheryl Acetate, C
 Multivitamins
 (Ascorbic Acid,
 Ergocalciferol,
 Folic Acid,
 Nicotinamide, C
 Calcium Carbonate
 (Calcium Carbonate) C
 Dicycloverine
 (Dicycloverine) C

Date:03/21/05ISR Number: 4616416-4Report Type:Expedited (15-DaCompany Report #2005042689
 Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion	Consumer	Neurontin			
Other		Headache		(Gabapentin)	PS		ORAL
100 MG (100		Stress					
MG, 1 IN 1							
D), ORAL				Carbamazepine (Carbamazepine)	C		

Date:03/21/05ISR Number: 4616417-6Report Type:Expedited (15-DaCompany Report #2005042638
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anxiety	Consumer	Neurontin			
Hospitalization -		Drug Effect Decreased		(Gabapentin)	PS		ORAL
Initial or Prolonged							
400 MG (100		Eczema					
Other							
MG, 4 IN 1							

D), ORAL	Feeling Abnormal				
	Insomnia		Xanax Tablet		
0.25 MG, ORAL	Mental Disorder		(Alprazolam)	SS	ORAL
	Post-Traumatic Stress Disorder		Hydroxyzine Pamoate (Caps) (Hydroxyzine Pamoate)	SS	ORAL
ORAL	Weight Fluctuation		Quetiapine Fumarate (Quetiapine Fumarte)	SS	
			Propylthiouracil (Propylthiouracil)	C	
			Lisinopril (Lisinopril)	C	
			Lekovit Ca (Calcium Carbonate, Colecalciferol)	C	

Date:03/21/05ISR Number: 4616890-3Report Type:Expedited (15-DaCompany Report #2005042664
Age:52 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Hepatitis	Foreign Health Professional	Neurontin (Gabapentin) Heparin-Fraction,	PS		ORAL

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Freedom Of Information (FOI) Report

SUBCUTANEOUS 40 MG (40 MG,
 1 IN 1 D),
 SUBCUTANEOUS

Sodium Salt
 (Heparin-Fraction,
 Sodium Salt) SS

Hydroxyzine
 Hydrochloride
 (Hydroxyzine
 Hydrochloride) C
 Clonazepam
 (Clonazepam) C
 Prednisone
 (Prednisone) C

Date:03/21/05ISR Number: 4616899-XReport Type:Expedited (15-DaCompany Report #2005043004
 Age:14 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Aggression	Foreign	Neurontin			
Other		Eye Disorder	Health	(Gabapentin)	PS		ORAL
900 MG (300		Feeling Abnormal	Professional				
MG, 3 IN 1		Social Avoidant Behaviour					
D), ORAL				Gabapentin			
				(Gabapentin)	SS		ORAL
900 MG (300							
MG, 3 IN 1							
D), ORAL				Valproate Sodium			
				(Valproate Sodium)	C		

Date:03/21/05ISR Number: 4627731-2Report Type:Direct Company Report #CTU 243817
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other 600 MG PO TID	Drug Intolerance	Gabapentin 600 Mg	PS	ORAL
	Nausea	Neurontin	C	

Date:03/22/05ISR Number: 4615226-1Report Type:Expedited (15-DaCompany Report #PHEH2005US03386
Age:18 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Balance Disorder		Trileptal	PS	Novartis Sector:	
Initial or Prolonged	False Negative Laboratory				Pharma	ORAL
Other	Result		Neurontin	SS		
	Optic Neuritis		Urbanyl	SS		
	Status Epilepticus					
	Visual Acuity Reduced					
	Visual Field Defect					

Date:03/22/05ISR Number: 4617572-4Report Type:Expedited (15-DaCompany Report #2005042988
Age:39 YR Gender:Female I/FU:I

Outcome	PT
Other	Activities Of Daily
	Living Impaired
	Arteriovenous
	Malformation
	Condition Aggravated
	Medication Error

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Spinal Disorder Suicidal Ideation Tremor	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Neurontin (Gabapentin)	PS		ORAL
1600 MG							
DAILY, ORAL				Clonazepam (Clonazepam)	C		

Date:03/22/05ISR Number: 4617574-8Report Type:Expedited (15-DaCompany Report #2005042991
Age: Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Consumer	Neurontin (Gabapentin)	PS		ORAL
Dose		Amnesia Condition Aggravated					
Hospitalization - Initial or Prolonged							
300 MG (100 Other MG, 3 IN 1		Fall					
D), ORAL		Hypoaesthesia					
		Impaired Driving Ability Insomnia Loss Of Consciousness Pain In Extremity Pharmaceutical Product Complaint		Paracetamol (Paracetamol) All Other Therapeutic Products	C C		

Date:03/22/05ISR Number: 4617767-XReport Type:Expedited (15-DaCompany Report #DEU-2005-0001316
Age:84 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Foreign Health Professional	Mst 10 Mg Mundipharma(Morphine Sulfate) Cr Tablet	PS		ORAL
Dose		Drug Interaction Glasgow Coma Scale Abnormal					
Hospitalization - Initial or Prolonged							
10 MG, BID, ORAL		Overdose	Other				
		Stupor		Valoron N (Tilidine			

75 MG, QID, ORAL	Hydrochloride) Oral Drops	SS	ORAL
25 MG, DAILY, ORAL	Surmontil (Trimipramine) Tablet	SS	ORAL
600 MG, TID, ORAL	Neurontin (Gabapentin) Coated Tablet	SS	ORAL
30 MG, DAILY. ORAL	Agopton (Lansoprazole)	SS	
100 MG, DAILY, ORAL	Beloc Zok Tablet (Metoprolol Succinate) Tablet	SS	
	Marcoumar (Phenprocoumon)	C	
	Triatec (Ramipril)	C	
	Norvasc (Amlodipine Besilate)	C	
	Selipran (Pravastatin Sodium)	C	
	Corvaton "Hoechst"		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Molsidomine)	C
Paracetamol	C
Calcium	C
Fosamax (Alendronate Sodium)	C
Nitroderm (Glyceryl Trinitrate)	C
Torem (Torasemide)	C

Date:03/22/05ISR Number: 4617803-0Report Type:Expedited (15-DaCompany Report #2005043001
Age:54 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (300 MG 3 IN 1 D), ORAL	Vitamin B12 Deficiency	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
10 DROP, ORAL			Clonazepam (Clonazepam)	SS		ORAL
			Insulin (Insulin)	C		

Date:03/22/05ISR Number: 4617973-4Report Type:Expedited (15-DaCompany Report #2005028590
Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Death Other ORAL	Dysphagia Parkinson'S Disease	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Sinemet (Carbidopa, Levodopa)	C		
			Carbamazepine (Carbamazepine)	C		

Date:03/22/05ISR Number: 4618048-0Report Type:Expedited (15-DaCompany Report #2005045983
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Circulatory Collapse	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL			Professional Company Representative				
				Ethanol (Ethanol) Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS C		

Date:03/22/05ISR Number: 4618060-1Report Type:Expedited (15-DaCompany Report #2005042344
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - (200 MG, Initial or Prolonged ORAL		Chest Pain Diabetes Mellitus Hiatus Hernia Intervertebral Disc Protrusion Micturition Disorder	Consumer	Celebrex (Celecoxib)	PS		ORAL
4 MG (4 MG, 1 IN 1 D), ORAL				Detrol La Capsule, Prolonged Release (Tolterodine L-Tartrate)	SS		ORAL
				Neurontin			

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(Gabapentin) SS
 Lisinopril
 (Lisinopril) C

Date:03/22/05ISR Number: 4618077-7Report Type:Expedited (15-DaCompany Report #001-0945-980292
 Age:51 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (300 Other MG, 4 IN 1 Required D), ORAL Intervention to Prevent Permanent Impairment/Damage	Chronic Obstructive Pulmonary Disease Depression Drug Effect Decreased Dysgeusia Dysuria Epididymitis Feeling Abnormal Headache Influenza Like Illness Insomnia Neoplasm Pain Pancreatitis Pneumonia Mycoplasmal Respiratory Failure Suicidal Ideation Swollen Tongue Upper Respiratory Tract Infection	Consumer Health Professional	Neurontin (Gabapentin) Carisprodol (Carisprodol) Hydrocodone (Hydrocodone) Amitriptyline (Amitriptyline) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	PS C C C C		ORAL

Date:03/22/05ISR Number: 4618112-6Report Type:Expedited (15-DaCompany Report #2005044386
 Age:67 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG (100 MG, 1 IN 1	Dysphagia Throat Tightness Vomiting	Consumer	Neurontin (Gabapentin)	PS		ORAL

D), ORAL

Date:03/22/05ISR Number: 4618118-7Report Type:Expedited (15-DaCompany Report #2005043864

Age:49 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Amnesia	Consumer	Neurontin			
Initial or Prolonged	Depression		(Gabapentin)	PS		ORAL
ORAL						
Other	Suicidal Ideation		All Other			
			Non-Therapeutic			
			Products (All Other			
			Non-Therapeutic			
			Products)	SS		
			Levothyroxine Sodium			
			(Levothyroxine			
			Sodium)	C		
			Rosuvastatin			
			(Rosuvastatin)	C		
			Verapamil			
			(Verapamil)	C		

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FDA - Adverse Event Reporting System (AERS)

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Date:03/22/05ISR Number: 4619974-9Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 243920

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased		Gabapentin 400 Mg	PS		ORAL
2 TABLETS PO		Pharmaceutical Product					
BID		Complaint					

Date:03/23/05ISR Number: 4618122-9Report Type:Expedited (15-DaCompany Report #TPG2005A00081
 Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Drug Interaction - Herpes Zoster Meningitis	Foreign Health Professional	Agopton (Lansoprazole) (30 Milligram, Capsules)	PS		ORAL
30 MG, (30 MG, 1 IN 1 D) PER ORAL		Neuralgia	Other				
		Overdose		Mst Continus (Morphine Sulfate) (10 Milligram, Tablets)	SS		ORAL
20 MG (10 MG, 2 IN 1 D) PER ORAL	2 DAY	Stupor					
300 MG (75 MG, 4 IN 1 D) PER ORAL	3 DAY			Valoron (Tilidine Hydrochloride) (Drops)	SS		ORAL
				Surmontil (Trimipramine Maleate) (25			

25 MG (25 MG, 1 IN 1 D) PER ORAL	2 DAY	Milligram, Tablets)	SS	ORAL
1800 (600 MG, 3 IN 1 D) PER ORAL	3 DAY	Neurontin (Gabapentin) (600 Milligram, Tablets)	SS	ORAL
100 MG (100 MG, 1 IN 1 D) PER ORAL		Beloc Zok (Metoprolol Succinate) (100 Milligram, Tablets)	SS	ORAL
		Marcoumar (Phenprocoumon) (3 Milligram, Tablets)	C	
		Triatec (Ramipril) (5 Milligram, Tablets)	C	
		Norvasc (Amlodipine Besilate) (5 Milligram, Tablets)	C	
		Selipran (Pravastatin Sodium) (40 Milligram, Tablets)	C	
		Corvaton (Molsidomine) (4 Milligram, Tablets)	C	

Freedom Of Information (FOI) Report

Paracetamol
 (Paracetamol) (1
 Gram, Tablets) C
 Calcium (Calcium
 /N/A/) C
 Fosamax (Alendronate
 Sodium) (70
 Milligram, Tablets) C
 Nitroderm Tts
 (Glyceryl
 Trinitrate) (50
 Milligram, Poultice
 Or Patch) C
 Torem (Torasemide
 (10 Milligram,
 Tablets) C

Date:03/23/05ISR Number: 4618558-6Report Type:Expedited (15-DaCompany Report #2005042797
 Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG (800 Other MG, 3 IN 1 D)		Cytolytic Hepatitis Intracranial Pressure Increased	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Levetiracetam (Levetiracetam)	PS C		

Date:03/23/05ISR Number: 4618762-7Report Type:Expedited (15-DaCompany Report #2005045468
 Age:72 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG (100 MG, 3 IN 1 D), ORAL		Pain In Extremity Peripheral Coldness Prostate Cancer	Consumer	Neurontin (Gabapentin) Glipizide	PS		ORAL

9glipizide0 C
 Metformin
 Hydrochloride
 (Metfromin
 Hydrochloride0 C
 Atorvastatin
 (Atorvasttin) C
 Enalapril
 (Enalapril) C

Date:03/23/05ISR Number: 4618763-9Report Type:Expedited (15-DaCompany Report #2005044705
 Age:66 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Bedridden Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG , 3 IN 1 D), ORAL		Drug Ineffective Immobile Inflammation		Dyazide (Hydrochlorothiazide , Triamterene) Atenolol (Atenolol)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Estrogen Nos
 (Estrogen Nos) C
 Vitamins (Vitamins) C

Date:03/23/05ISR Number: 4618764-0Report Type:Expedited (15-DaCompany Report #2005043174
 Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 MG, 2 IN 1 D), ORAL		Gastric Disorder - Paraesthesia Pharmaceutical Product Complaint	Consumer	Neurontin (Gabapentin)	PS		ORAL

Allopurinol
 (Allopurinol) C
 Levothyroxine Sodium
 (Levothyroxine
 Sodium) C
 Oemprazole
 (Omeprazole0) C

Date:03/23/05ISR Number: 4618771-8Report Type:Expedited (15-DaCompany Report #2005044427
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 1800 MG (600 MG, 3 IN 1 D), ORAL		Bite Drug Dependence Drug Withdrawal Syndrome Generalised Oedema Nasopharyngitis Personality Change Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Estrogens Conjugated (Estrogens Conjugated) All Other Therapeutic Products (All Other Therapeutic	C		

Products) C
Levothyroxine Sodium
(Levothyroxine
Sodium) C
Lotrel (Amlodipine,
Benazepril
Hydrochloride) C
Biselect (Bisoprolol
Fumarate,
Hydrochlorothiazide) C

Date:03/23/05ISR Number: 4618772-XReport Type:Expedited (15-DaCompany Report #2005044792

Age: Gender:Female I/FU:I

Outcome PT
Disability Accident
Other Balance Disorder
Chest Discomfort
Disorientation
Drug Ineffective
Fatigue
Feeling Abnormal
Fibromyalgia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG (200 MG, 3 IN 1 D), ORAL		Heart Rate Increased Hypersensitivity Intervertebral Disc Protrusion Loss Of Consciousness Pain Pharmaceutical Product	Consumer	Neurontin (Gabapentin)	PS		ORAL
90 MG, 1 IN 1 D, ORAL		Complaint Respiratory Distress Sedation		Morphine Sulfate (Morphine Sulfate)	SS		ORAL
				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	SS		
				Carisoprodol (Carisoprodol)	C		
				Tizanidine Hydrochloride (Tizanidine Hydrochloride)	C		

Date:03/23/05ISR Number: 4618777-9Report Type:Expedited (15-DaCompany Report #2005044422
Age:70 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Other		Condition Aggravated Convulsion Feeling Abnormal Overdose Pain Pharmaceutical Product Complaint	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Irbesartan (Irbesartan)	C		
				Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
				Carvedilol (Carvedilol)	C		
				Acetylsalicylic Acid (Acetylsalicylic			

Acid) C
Ezetimibe C
(Ezetimibe)
Rosiglitazone
Maleate
(Rosiglitazone
Maleate) C

Date:03/23/05ISR Number: 4618848-7Report Type:Expedited (15-DaCompany Report #2004048907
Age:51 YR Gender:Female I/FU:F

Outcome PT
Hospitalization - Abdominal Distension
Initial or Prolonged Abdominal Injury
Other Abdominal Pain
Agitation
Anger
Anxiety
Arterial Injury
Atrial Fibrillation
Biliary Tract Disorder
Blood Calcium Decreased

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Freedom Of Information (FOI) Report

Dose	Duration	Condition Aggravated	Report Source	Product	Role	Manufacturer	Route
2000 MG (440 MG, 5 IN 1 D)		Body Dysmorphic Disorder Compulsions	Consumer	Neurontrn (Gabapentin)	PS		
		Confusional State Depression					
		Diaphragmatic Injury					
		Drug Screen Positive		Cefalexin (Cefalexin)	C		
		Dysphagia		Warfarin Sodium (Warfarin Sodium)	C		
		Economic Problem		Phenylpropanolamine (Phenylpropanolamine	C		
		Faecaloma)	C		
		Feeling Abnormal		Cyproheptadine Hydrochloride (Cyproheptadine	C		
		Feeling Guilty		Hydrochloride)	C		
		Gun Shot Wound		Yohimbine (Yohimbine)	C		
		Haematocrit Decreased		Cetirizine Hydrochloride	C		
		Haemoglobin Decreased		(Cetirizine	C		
		Haemothorax		Hydrochloride)	C		
		Hepatic Trauma		Adapalene (Adapalene)	C		
		Hypernatraemia		Meloxicam (Meloxicam)	C		
		Hypotension		Aripiprazole (Aripiprazole)	C		
		Insomnia		Ketorolac Tromethamine (Ketorolac	C		
		Intercostal Neuralgia		Tromethamine)	C		
		Laceration		Isocom (Dichloralphenazone,	C		
		Malnutrition		Isometheptene	C		
		Mental Impairment		Mucate, Paracetamol)	C		
		Mood Altered		Clonazepam (Clonazepam)	C		
		Nephrogenic Diabetes		Olanzapine (Olanzapine)	C		
		Insipidus		Zolpidem Tartrate (Zolpidem Tartrate)	C		
		Pain		Fluoxetine			
		Pericardial Disease					
		Pericardial Effusion					
		Pleuritic Pain					
		Pneumatois Cystoides					
		Intestinalis					
		Pneumonia					
		Pneumothorax					
		Polytraumatism					
		Psychiatric Symptom					
		Pulmonary Embolism					
		Respiratory Failure					
		Scar					
		Sedation					
		Self Esteem Decreased					
		Self Injurious Behaviour					
		Suicide Attempt					

Surgery
Tachypnoea
Traumatic Shock

Hydrochloride
(Fluoxetine
Hydrochloride) C
Raloxifene
Hydrochloride
(Raloxifene
Hydrochloride) C
Lithium Carbonate
(Lithium Carbonate) C
Tramadol
Hydrochloride
(Tramadol
Hydrochloride) C
Atenolol (Atenolol) C
Trazodone
(Trazodone) C
Propacet

Freedom Of Information (FOI) Report

(Dextropropoxyphene Napsilate, Paracetamol)	C
Estratest Hs (Estrogens Esterified, Methyltestosterone)	C
Estradiol (Estradiol)	C
Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C
Nefazodone Hydrochloride (Nefazodone Hydrochloride)	C
Zolmitriptan (Zolmitriptan)	C
Sodium Fluoride (Sodium Fluoride)	C
Amoxicillin (Amoxicillin)	C
Meperidine W/Promethazine (Pethidine, Promethazine)	C
Azithromycin (Azithromycin)	C
Aquatab C (Dextromethorpan, Guaifenesin, Phenylpropanolamine)	C
Prednisone (Prednisone)	C
Pravastatin Sodium (Pravastatin Sodium)	C
Clindamycin (Clindamycin)	C
Chlorhexidine (Chlorhexidine)	C
Vicodin (Hydrocodone Bitartrate, Paracetamol)	C
Docusate Sodium (Docusate Sodium)	C
Risperidone (Risperidone)	C

Mirtazapine (Mirtazapine)	C
Bupropion Hydrochloride (Bupropion Hydrochloride)	C
Propranolol Hydrochloride (Propranolol Hydrochloride)	C
Sumatriptan Succinate (Sumatriptan Succinate)	C

Freedom Of Information (FOI) Report

Methocarbamol	
(Methocarbamol)	C
Naproxen (Naproxen)	C
Narine Repetabs	
(Loratadine,	
Pseudoephedrine	
Sulfate)	C
Temazepam	
(Temazepam)	C
Quetiapine Fumarate	
(Quetiapine	
Fumarate)	C
Robitussin A-C /Old	
Form/ (Codeine	
Phosphate,	
Guaifenesin,	
Pheniramine Maleate)	C
Flurazepam	
Hydrochloride	
(Flurazepam	
Hydrochloride)	C
Mometasone Furoate	
(Mometasone Furoate)	C
Adapalene	
(Adapalene)	C
Ronatic	
(Chlorphenamine	
Tannate, Mepyramine	
Tannate,	
Phenylephrine	C
Modafinil	
(Modafinil)	C
Tizanidine	
Hydrochloride	
(Tizanidine	
Hydrochloride)	C
Estrogens Conjugated	
(Estrogens	
Conjugated)	C
Estradiol	
(Estradiol)	C
Tiagabine	
Hydrochloride	
(Tiagabine	
Hydrochloride)	C
Sildenafil Citrate	
(Sildenafil Citrate)	C

Butorphanol Tartrate
(Butorphanol
Tartrate) C
Trimethobenzamide
Hydrochloride
(Trimethobenzamide
Hydrochloride) C
Lithium Carbonate
(Lithium Carbonate) C
Valproate Semisodium
(Valproate
Semisodium) C
Beclometasone
Dipropionate

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Freedom Of Information (FOI) Report

(Beclometasone
Dipropionate) C
Clavulin
(Amoxicillin
Trihydrate,
Clavulanate
Potassium) C

Date:03/23/05ISR Number: 4618851-7Report Type:Expedited (15-DaCompany Report #2004059024
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthralgia	Consumer	Neurontin			
		Arthritis		(Gabapentin)	PS		
1800 MG (600		Knee Arthroplasty					
MG, 3 IN 1 D)		Osteoporosis		Fentanyl (Fentanyl)	C		
				Ibuprofen			
				(Ibuprofen)	C		
				Rofecoxib			
				(Rofecoxib)	C		
				Heliantus Tuberosus			
				(Heliantus			
				Tuberosus)	C		
				All Other			
				Therapeutic Products	C		

Date:03/23/05ISR Number: 4618857-8Report Type:Expedited (15-DaCompany Report #2005044232
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alopecia	Consumer	Neurontin			
		Antipsychotic Drug Level		(Gabapentin)	PS		ORAL
800 MG (1 D),		Increased					
ORAL		Bipolar Disorder		Lithium (Lithium)	SS		
		Bladder Prolapse		Cyclobenzaprine			
		Breath Odour		(Cyclobenzaprine)	C		
		Dental Necrosis		Thyroid (Thyroid)	C		
		Drug Ineffective		All Other			

Hypoaesthesia
Medication Error
Panic Attack
Tooth Discolouration
Tooth Loss

Therapeutic Products
(All Other
Therapeutic
Products) C
Clonazepam
(Clonazepam) C
Rosiglitazone
Maleate
(Rosiglitazone
Maleate) C
Naproxen (Naproxen) C
Simvastatin
(Simvastatin) C

Date:03/23/05ISR Number: 4618895-5Report Type:Expedited (15-DaCompany Report #2005042258
Age:45 YR Gender:Female I/FU:F

Outcome PT
Other Complex Partial Seizures
Drug Ineffective
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
ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
			Phenytoin Sodium (Phenytoin Sodium)	C		
			Lamotrigine (Lamotrigine)	C		
			Lorazepam (Lorazepam)	C		
			Fexofenadine Hydrochloride (Fexofenadine Hydrochloride)	C		
			Ipratropium Bromide (Ipratropium Bromide)	C		
			Fluticasone Propionate (Fluticasone Propionate)	C		
			Pirbuterol Acetate (Pirbuterol Acetate)	C		
			Epinephrine (Epinephrine)	C		

Date:03/23/05ISR Number: 4618898-0Report Type:Expedited (15-DaCompany Report #2005045432
Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Death	Consumer	Nurontin (Gabapentin)	PS		
				Antihypertensives (Antihypertensives)	C		

Date:03/23/05ISR Number: 4621267-0Report Type:Direct
Age:37 YR Gender:Female I/FU:I

Company Report #CTU 244028

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG QID		Condition Aggravated Pain In Extremity Pharmaceutical Product Complaint		Gabapentin	PS		

Date:03/24/05ISR Number: 4620368-0Report Type:Expedited (15-DaCompany Report #2005044794
Age:85 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (100 MG, 3 IN 1 D), ORAL ORAL		Blood Glucose Increased Hypoaesthesia Pain Paraplegia Pharmaceutical Product Complaint	Consumer	Neurontin (Gabapentin) Gabapentin (Gabapentin) All Other Therapeutic Products (All Other	PS SS		ORAL ORAL

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Therapeutic Products)	C
Levothyroxine Sodium (Levothyroxine Sodium)	C
Candesartan Cilexetil (Candesartan Cilexetil)	C
Amiodarone (Amiodarone)	C
Torasemide (Torasemide)	C
Vitamins, Other Combinations (Vitamins, Other Combinations)	C
Pentoxifylline (Pentoxifylline)	C
Amlodipine Besilate (Amlodipine Besilate)	C
Potassium Chloride (Potassium Chloride)	C
Prednisone (Prednisone)	C
Warfarin (Warfarin)	C
Atorvastatin (Atorvastatin)	C
Glimepiride (Glimepiride)	C

Date:03/24/05ISR Number: 4620374-6Report Type:Expedited (15-DaCompany Report #2005044740

Age:74 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Fatigue	Consumer	Neurontin			
Other		Goitre		(Gabapentin)	PS		ORAL
900 MG (300		Somnolence					
MG, 3 IN 1							
D), ORAL				Omeprazole			
				(Omeprazole)	C		

Tocopherol
(Tocopherol) C
Calcium (Calcium) C
Centrum Silver
(Ascorbic Acid,
Calcium, Minerals
Nos, Retinol,
Tocopheryl Acetate, C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C

Date:03/24/05ISR Number: 4620386-2Report Type:Expedited (15-DaCompany Report #2005044442
Age:35 YR Gender:Male I/FU:I

Outcome PT
Other Abdominal Distension
Blood Cholesterol

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (300 MG, 1 IN 1 D), ORAL		Increased Bowel Sounds Abnormal Convulsion Dysgeusia Haematemesis Haemorrhage Hair Colour Changes Melaena Muscle Disorder Muscle Twitching Pharmaceutical Product Complaint Vomiting Weight Fluctuation	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Dextropropoxyphene Hydrochloride (Dextropropoxyphene Hydrochloride) Propacet (Dextropropoxyphene Napsilate, Paracetamol) Tramadol Hydrochloride (Tramadol Hydrochloride) Esomeprazole (Esomeprazole) Lansoprazole (Lansoprazole) Rabeprazole Sodium (Rabeprazole Sodium) Pantoprazole (Pantoprazole) Acetylsalicylic Acid (Acetylsalicylic Acid)	C C C C C		

Date:03/24/05ISR Number: 4620388-6Report Type:Expedited (15-DaCompany Report #2005044743
Age:55 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, 3 IN 1 D)		Blindness Convulsion Coordination Abnormal	Consumer	Neurontin (Gabapentin)	PS		

Dizziness	Estratest Hs	
Feeling Abnormal	(Estrogens	
Illiteracy	Esterified,	
Impaired Driving Ability	Methyltestosterone)	C
Pain	Medroxyprogesterone	
Pharmaceutical Product	Acetate	
Complaint	(Medroxyprogesterone	
Vertigo	Acetate)	C
	Atorvastatin	
	(Atorvastatin)	C
	Clonazepam	
	(Clonazepam)	C
	Furosemide	
	(Furosemide)	C
	Tizanidine	
	(Tizanidine)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/24/05ISR Number: 4620424-7Report Type:Expedited (15-DaCompany Report #2005045559
 Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNK (100 MG, Other QOD), ORAL		Condition Aggravated Depression Drug Ineffective Hallucination Medication Error Pain Pharmaceutical Product Complaint	Consumer	Neurontin (Gabapentin) Nortriptyline (Nortriptyline)	PS C		ORAL

Date:03/24/05ISR Number: 4620476-4Report Type:Expedited (15-DaCompany Report #2005044700
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 800 MG (400 MG, 2 IN 1 D), ORAL 3600 MG (1200 MG, 3 IN 1 D)		Acne Chromaturia Depressed Level Of Consciousness Diabetes Mellitus Dysuria Fatigue Feeling Abnormal Hallucination, Auditory Memory Impairment Pain Paranoia Skin Atrophy Speech Disorder Urine Analysis Abnormal	Consumer	Neurontin (Gabapentin) Gabapentin (Gabapentin) Tocopherol (Tocopherol) Baclofen (Baclofen) Furosemide (Furosemide) Potassium (Potassium) Lekovit Ca (Calcium Carbonae, Colecalciferol) Prenatal Vitamins (Ascorbic Acid,	PS SS C C C C C		ORAL

Biotin, Minerals
 Nos, Nicotinic Acid,
 Retinol, Tocopherol, C

Date:03/24/05ISR Number: 4620569-1Report Type:Expedited (15-DaCompany Report #2005045811
 Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 100 MG (100 MG, 1 IN 1 D), ORAL		Back Pain	Consumer	Zoloft (Sertraline)	PS		ORAL
		Condition Aggravated					
		Intervertebral Disc					
		Degeneration		Xanax Tablet			
3 MG (1 MG, 3 IN 1 D), ORAL		Intervertebral Disc		(Alprazolam)	SS		ORAL
		Protrusion					
		Neuropathy		Neurontin			
		Treatment Noncompliance		(Gabapentin)	SS		
				Allegra-D			
				(Fexofenadine, Pseudoephedrine Hydrochloride)	C		
				Prinzide			
				(Hydrochlorothiazide , Lisinopril)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/24/05ISR Number: 4624105-5Report Type:Direct
Age:50 YR Gender:Male I/FU:I

Company Report #CTU 244252

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
900 MG, DAILY, ORAL		Overdose					

Fentanyl C

Date:03/24/05ISR Number: 4624106-7Report Type:Direct
Age:41 YR Gender:Male I/FU:I

Company Report #CTU 244216

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1800MG DAILY, ORAL		Gun Shot Wound					

Remoron C

Date:03/24/05ISR Number: 4624107-9Report Type:Direct
Age:63 YR Gender:Male I/FU:I

Company Report #CTU 244217

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2400MG DAILY, ORAL		Gun Shot Wound					

Bextra C
Oxycontin C
Altace C
Warfarin C

Date:03/24/05ISR Number: 4624108-0Report Type:Direct
Age:30 YR Gender:Male I/FU:I

Company Report #CTU 244218

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death 1800MG DAILY	Completed Suicide Gun Shot Wound		Neurontin	PS		
Date:03/24/05ISR Number: 4624109-2Report Type:Direct		Company Report #CTU 244219				
Age:49 YR Gender:Male	I/FU:I					
Outcome Dose Death 3300MG DAILY, ORAL	PT Completed Suicide Gun Shot Wound	Report Source	Product Neurontin Elavil Senexon	Role PS C C	Manufacturer	Route ORAL
Date:03/24/05ISR Number: 4624110-9Report Type:Direct		Company Report #CTU 244220				
Age:51 YR Gender:Male	I/FU:I					
Outcome Dose Death	PT Completed Suicide Gun Shot Wound	Report Source	Product Neurontin	Role PS	Manufacturer	Route
Date:03/24/05ISR Number: 4624111-0Report Type:Direct		Company Report #CTU 244221				
Age:46 YR Gender:Male	I/FU:I					
Outcome Dose Death 2700MG	PT Completed Suicide Overdose	Report Source	Product Neurontin Zoloft	Role PS C	Manufacturer	Route
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Seroquel C
Klonopin C

Date:03/24/05ISR Number: 4624112-2Report Type:Direct
Age:45 YR Gender:Female I/FU:I

Company Report #CTU 244222

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY	Completed Suicide		Neurontin	PS		
		Overdose		Oxycontin	C		
				Clonazepam	C		
				Percocet	C		
				Keppra	C		

Date:03/24/05ISR Number: 4624113-4Report Type:Direct
Age:41 YR Gender:Male I/FU:I

Company Report #CTU 244244

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	4800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose		Clonazepam	C		
				Duragesic	C		
				Effexor	C		

Date:03/24/05ISR Number: 4624114-6Report Type:Direct
Age:44 YR Gender:Male I/FU:I

Company Report #CTU 244245

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose		Paxil	C		
				Seroquel	C		
				Pepakote	C		

Date:03/24/05ISR Number: 4624115-8Report Type:Direct Company Report #CTU 244246
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
300 MG DAILY		Intentional Misuse					

Date:03/24/05ISR Number: 4624116-XReport Type:Direct Company Report #CTU 244247
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/24/05ISR Number: 4624117-1Report Type:Direct Company Report #CTU 244248
Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1800MG DAILY,		Overdose					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/24/05ISR Number: 4624118-3Report Type:Direct
Age:45 YR Gender:Female I/FU:I

Company Report #CTU 244249

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	3000MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose		Paxil	C		
				Zyprexa	C		

Date:03/24/05ISR Number: 4624119-5Report Type:Direct
Age:35 YR Gender:Male I/FU:I

Company Report #CTU 244250

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose					

Date:03/24/05ISR Number: 4624120-1Report Type:Direct
Age:59 YR Gender:Female I/FU:I

Company Report #CTU 244251

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	600MG, DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose					

Date:03/24/05ISR Number: 4624121-3Report Type:Direct
Age:56 YR Gender:Male I/FU:I

Company Report #CTU 244236

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	3000MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose					

Celebrex C
Novolin C
Paxil C
Nexium C

Date:03/24/05ISR Number: 4624123-7Report Type:Direct Company Report #CTU 244237
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2400MG DAILY, ORAL		Overdose					
				Klonopin	C		
				Trezodone	C		
				Celebrex	C		
				Prevacid	C		

Date:03/24/05ISR Number: 4624124-9Report Type:Direct Company Report #CTU 244238
Age:39 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
300 MG 3X DAILY		Overdose					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/24/05ISR Number: 4624125-0Report Type:Direct
Age:47 YR Gender:Male I/FU:I

Company Report #CTU 244239

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1600 MG	Completed Suicide		Neurontin	PS		ORAL
	DAILY, ORAL	Overdose					
				Paxil	C		
				Clonazepam	C		
				Ambien	C		

Date:03/24/05ISR Number: 4624126-2Report Type:Direct
Age:38 YR Gender:Male I/FU:I

Company Report #CTU 244240

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/24/05ISR Number: 4624127-4Report Type:Direct
Age:17 YR Gender:Male I/FU:I

Company Report #CTU 244241

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY	Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/24/05ISR Number: 4624128-6Report Type:Direct
Age:43 YR Gender:Female I/FU:I

Company Report #CTU 244242

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/24/05ISR Number: 4624129-8Report Type:Direct Company Report #CTU 244243
Age:58 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		

Date:03/24/05ISR Number: 4624130-4Report Type:Direct Company Report #CTU 244228
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		ORAL
2400MG DAILY, ORAL							

Date:03/24/05ISR Number: 4624131-6Report Type:Direct Company Report #CTU 244229
Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		ORAL
1200 MG DAILY, ORAL							
				Elavil	C		
				Methadone	C		
				Ambien	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/24/05ISR Number: 4624132-8Report Type:Direct
Age:60 YR Gender:Male I/FU:I

Company Report #CTU 244230

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose					
				Klonopin	C		
				Ultram	C		
				Celexa	C		

Date:03/24/05ISR Number: 4624133-XReport Type:Direct
Age:43 YR Gender:Female I/FU:I

Company Report #CTU 244231

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose					

Date:03/24/05ISR Number: 4624140-7Report Type:Direct
Age:55 YR Gender:Female I/FU:I

Company Report #CTU 244232

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Intentional Misuse					

Date:03/24/05ISR Number: 4624149-3Report Type:Direct
Age:58 YR Gender:Male I/FU:I

Company Report #CTU 244233

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/24/05ISR Number: 4624159-6Report Type:Direct
Age:61 YR Gender:Female I/FU:I

Company Report #CTU 244234

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2400MG DAILY,		Overdose					
ORAL				Ambien	C		
				Clonidine	C		
				Nexium	C		
				Diazepam	C		

Date:03/24/05ISR Number: 4624168-7Report Type:Direct
Age:51 YR Gender:Female I/FU:I

Company Report #CTU 244235

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1200 MG DAILY		Overdose					
ORAL				Sonata	C		
				Actonel	C		
				Mofic	C		
				Imitrix	C		
				Indomethacin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/24/05ISR Number: 4624208-5Report Type:Direct
Age:35 YR Gender:Male I/FU:I

Company Report #CTU 244223

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose					

Date:03/24/05ISR Number: 4624209-7Report Type:Direct
Age:57 YR Gender:Female I/FU:I

Company Report #CTU 244224

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2700MG DAILY	Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/24/05ISR Number: 4624210-3Report Type:Direct
Age:36 YR Gender:Female I/FU:I

Company Report #CTU 244225

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2400MG DAILY	Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/24/05ISR Number: 4624211-5Report Type:Direct
Age:49 YR Gender:Female I/FU:I

Company Report #CTU 244226

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	600MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose					

Date:03/24/05ISR Number: 4624212-7Report Type:Direct Company Report #CTU 244227
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1800MG DAILY, ORAL		Injury Asphyxiation					

Date:03/25/05ISR Number: 4620706-9Report Type:Expedited (15-DaCompany Report #2005044389
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Ineffective Fatigue	Consumer	Neurontin (Gabapentin)	PS		ORAL
1200 MG, ORAL		Feeling Abnormal Hypersomnia Road Traffic Accident					

Date:03/25/05ISR Number: 4620722-7Report Type:Expedited (15-DaCompany Report #2005045397
Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Constipation Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL
(300 MG, 1 IN 1 D), ORAL		Feeling Abnormal Genital Burning Sensation Muscle Spasms Pharmaceutical Product Complaint		Estradiol (Estradiol) Verapamil (Verapamil)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Ibuprofen
 (Ibuprofen) C
 Lorazepam
 (Lorazepam) C
 Cyclobenzaprine
 Hydrochloride
 (Cyclobenzaprine
 Hydrochloride) C

Date:03/25/05ISR Number: 4620799-9Report Type:Expedited (15-DaCompany Report #2005044694
 Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (600 MG, 1 IN 1 D), ORAL		Abasia Feeling Abnormal Hypoaesthesia Pain	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Paracetamol (Paracetamol)	C		

Date:03/25/05ISR Number: 4620857-9Report Type:Expedited (15-DaCompany Report #2005045808
 Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (100 MG, 2 IN 1 D), ORAL		Cerebrovascular Accident Dialysis Pharmaceutical Product Complaint	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Insulin Glargine (Insulin Glargine) Glimepiride (Glimepiride) Triobe (Cyanocobalamin, Folic Acid,	C C		

Pyridoxine)	C
Furosemide	
(Furosemide)	C
Amlodipine Besilate	
(Amlodipine	
Besilate)	C
Metoprolol	
(Metoprolol)	C
Oxybutynin	
Hydrochloride	
(Oxybutynin	
Hydrochloride)	C
Acetylsalicylic Acid	
(Acetylsalicylic	
Acid)	C
Metoclopramide	
(Metoclopramide)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/25/05ISR Number: 4620860-9Report Type:Expedited (15-DaCompany Report #2005045392

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Ineffective Renal Disorder	Consumer	Neurontin (Gabapentin)	PS		
300 MG (100 MG, 3 IN 1 D)				Folic Acid (Folic Acid) Udramil (Trandolapril, Verapamil Hydrochloride)	C C		

Date:03/25/05ISR Number: 4620861-0Report Type:Expedited (15-DaCompany Report #2005045571

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Blood Pressure Decreased Pharmaceutical Product Complaint Sepsis Transplant Rejection	Consumer Health Professional	Neurontin (Gabapentin) Mycophenolate Mofetil (Mycophenolate Mofetil) All Other Therapeutic Products (All Other Therapeutic Products) Ciclosporin (Ciclosporin) Clonidine (Clonidine)	PS C C C C		

Date:03/25/05ISR Number: 4620885-3Report Type:Expedited (15-DaCompany Report #2005045395

Age:77 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged 500 MG (100 Other MG, 5 IN 1 D); ORAL	Blood Test Abnormal Feeling Abnormal Grand Mal Convulsion Pancreatitis	Consumer	Neurontin (Gabapentin)	PS	ORAL
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Pharmaceutical Product Complaint	Ranitidine Hydrochloride Methylphenobarbital Nabumetone Atenolol Paroxetine Hydrochloride	C C C C
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Date:03/25/05ISR Number: 4621047-6Report Type:Expedited (15-DaCompany Report #2005045460
Age:92 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 1800 MG (1 D) Initial or Prolonged		Cardiac Failure	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/25/05ISR Number: 4622489-5Report Type:Direct
Age:81 YR Gender:Female I/FU:I

Company Report #CTU 244328

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ONE TID	1 YR	Feeling Abnormal Gastric Disorder Pharmaceutical Product Complaint		Gabapentin 30 Mg Cap	PS		

Date:03/25/05ISR Number: 4622494-9Report Type:Direct
Age:62 YR Gender:Male I/FU:I

Company Report #CTU 244379

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death		Completed Suicide Overdose		Neurontin	PS		

Date:03/25/05ISR Number: 4622496-2Report Type:Direct
Age:30 YR Gender:Female I/FU:I

Company Report #CTU 244336

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 2700MG DAILY, ORAL		Completed Suicide Gun Shot Wound		Neurontin Remeron Ultram	PS C C		ORAL

Date:03/25/05ISR Number: 4622497-4Report Type:Direct
Age:57 YR Gender:Male I/FU:I

Company Report #CTU 244335

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 1200MG DAILY		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/25/05ISR Number: 4622498-6Report Type:Direct
Age:41 YR Gender:Female I/FU:I

Company Report #CTU 244334

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
2400MG DAILY		Gun Shot Wound					

Date:03/25/05ISR Number: 4622528-1Report Type:Direct
Age:49 YR Gender:Female I/FU:I

Company Report #CTU 244349

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
2400MG DAILY		Overdose					

Date:03/25/05ISR Number: 4622531-1Report Type:Direct
Age:67 YR Gender:Female I/FU:I

Company Report #CTU 244351

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2400MG DAILY		Intentional Misuse					
, ORAL				Klor-Con	C		
				Proboxy	C		
				Nifedipine	C		
				Tricor	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/25/05ISR Number: 4622533-5Report Type:Direct
Age:20 YR Gender:Female I/FU:I

Company Report #CTU 244352

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY	Completed Suicide		Neurontin	PS		ORAL
		Overdose					
				Zithromax	C		
				Baclofen	C		
				Benzonatate	C		
				Trazodone	C		

Date:03/25/05ISR Number: 4622536-0Report Type:Direct
Age:50 YR Gender:Female I/FU:I

Company Report #CTU 244355

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200MG DAILY	Completed Suicide		Neurontin	PS		
		Overdose					
				Klonzopam	C		
				Xanax	C		

Date:03/25/05ISR Number: 4622537-2Report Type:Direct
Age:48 YR Gender:Female I/FU:I

Company Report #CTU 244357

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800 MG	Completed Suicide		Neurontin	PS		ORAL
		Overdose					
	DAILY, ORAL						

Date:03/25/05ISR Number: 4622538-4Report Type:Direct
Age:25 YR Gender:Male I/FU:I

Company Report #CTU 244358

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/25/05ISR Number: 4622539-6Report Type:Direct
Age:46 YR Gender:Female I/FU:I

Company Report #CTU 244362

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
2600 MG		Overdose					

Date:03/25/05ISR Number: 4622540-2Report Type:Direct
Age:42 YR Gender:Female I/FU:I

Company Report #CTU 244364

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Intentional Misuse		Neurontin	PS		

Date:03/25/05ISR Number: 4622541-4Report Type:Direct
Age:77 YR Gender:Male I/FU:I

Company Report #CTU 244337

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
800MG		Gun Shot Wound					
DAILY, ORAL				Tequim	C		
				Leuthroid	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/25/05ISR Number: 4622542-6Report Type:Direct
Age:52 YR Gender:Male I/FU:I

Company Report #CTU 244338

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Self Injurious Behaviour		Neurontin	PS		ORAL
	ORAL						

Date:03/25/05ISR Number: 4622543-8Report Type:Direct
Age:58 YR Gender:Male I/FU:I

Company Report #CTU 244339

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Gun Shot Wound					

Date:03/25/05ISR Number: 4622544-XReport Type:Direct
Age:86 YR Gender:Male I/FU:I

Company Report #CTU 244342

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL						

Date:03/25/05ISR Number: 4622545-1Report Type:Direct
Age:38 YR Gender:Male I/FU:I

Company Report #CTU 244343

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2800MG	Completed Suicide		Neurontin	PS		
		Intentional Self-Injury					

Date:03/25/05ISR Number: 4622546-3Report Type:Direct Company Report #CTU 244344
Age:32 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1800MG							

Date:03/25/05ISR Number: 4622547-5Report Type:Direct Company Report #CTU 244345
Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
900MG DAILY,		Overdose					
ORAL				Topomax	C		
				Seroquel	C		
				Gabitril	C		
				Synarel	C		

Date:03/25/05ISR Number: 4622548-7Report Type:Direct Company Report #CTU 244348
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1600MG DAILY,		Overdose					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/28/05ISR Number: 4621147-0Report Type:Expedited (15-DaCompany Report #2005038107
 Age:31 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Disturbance In Attention Drug Ineffective	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (300 MG, 1 IN 1 D), ORAL		Epilepsy Somnolence					

Date:03/28/05ISR Number: 4621613-8Report Type:Expedited (15-DaCompany Report #2005045412
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Upper Alopecia	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG UP TO 3 CAPSULES (3 IN 1 D), ORAL		Back Pain Dysgeusia Muscle Spasms Renal Disorder Renal Pain Skin Disorder Swelling Tooth Disorder					

Date:03/28/05ISR Number: 4621615-1Report Type:Expedited (15-DaCompany Report #2005046005
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Life-Threatening		Hypotension Myocardial Infarction	Consumer	Neurontin (Neurontin)	PS		ORAL
200 MG (100 MG, 2 IN 1							

D), ORAL

Metoprolol
(Metoprolol) SS ORAL

ORAL

Allopurinol
(Allopurinol) C

Date:03/28/05ISR Number: 4621677-1Report Type:Expedited (15-DaCompany Report #2005045999

Age:68 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - 2400 MG (800 Initial or Prolonged MG, 3 IN 1 D) Other 400 MG (200 MG, 2 IN 1 D)	Coronary Artery Occlusion Depression Dyspnoea Facial Bones Fracture Fall Homeless Joint Injury Suicidal Ideation Suicide Attempt	Consumer	Neurontin (Gabapentin) Celebrex (Celecoxib) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Alprazolam (Alprazolam) Thyroid Hormones (Thyroid Hormones) Estrogens Conjugated (Estrogens Conjugated) Antihypertensives (Antihypertensives)	PS SS SS C C C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/28/05ISR Number: 4621679-5Report Type:Expedited (15-DaCompany Report #2005045777

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 300 MG (100 Other MG, 3 IN 1 D)	Back Pain Cataract Operation Condition Aggravated Drug Effect Decreased Eye Laser Surgery Herpes Zoster Intestinal Obstruction Pain In Extremity Radiculotomy	Consumer	Neurontin (Gabapentin)	PS		
			Amlodipine Besilate (Amlodipine Besilate)	C		
			Valsartan (Valsartan)	C		
			Carbamazepine (Carbamazepine)	C		
			Levothyroxine (Levothyroxine)	C		
			Lansoprazole (Lansoprazole)	C		
			Montelukast Sodium (Montelukast Sodium)	C		
			Clindamycin (Clindamycin)	C		
			Hydrocodone (Hydrocodone)	C		
			Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)	C		
			Combivent (Ipratropium Bromide, Salbutamol Sulfate)	C		
			Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
			Cyclobenzaprine (Cyclobenzaprine)	C		
			Risedronate Sodium (Risedronate Sodium)	C		
			Tolterodine L-Tartrate (Tolterodine			

L-Tartrate)

C

Date:03/28/05ISR Number: 4622028-9Report Type:Expedited (15-DaCompany Report #2005046048

Age:57 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Feeling Drunk	Consumer	Neurontin			
Initial or Prolonged	Neuralgia		(Gabapentin)	PS		ORAL
600 OR 800MG						
Other	Treatment Noncompliance					
(1 IN 1 D),						
ORAL	Ulcer Haemorrhage					
			Etodolac (Etodolac)	SS		ORAL
ORAL			Sertraline			
			Hydrochloride			
			(Sertraline			
			Hydrochloride)	C		
			Amitriptyline			
			(Amitriptyline)	C		

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Freedom Of Information (FOI) Report

Carvedilol
 (Carvedilol) C
 Spironolactone
 (Spironolactone) C
 Olmesartan Medoxomil
 (Olmesartan
 Medoxomil) C

Date:03/28/05ISR Number: 4622123-4Report Type:Expedited (15-DaCompany Report #US014711
 Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Communication Disorder	Foreign	Fentanyl Citrate	PS		
BUCCAL	1600 UG PRN	Disorientation	Study				
BUCCAL		Drug Interaction	Health	Gabapentin	SS		
600 MG DAILY		Drug Toxicity	Professional	Oxycontin	SS		
80 MG DAILY			Other	Oxynorm	SS		
30 MG DAILY				Co-Proxamol	SS		
8 TAB/CAP							
DAILY							

Paracetamol C
 Dexamethazone-Pix C
 Lansoprazole
 "Boehringer
 Ingelheim" C
 Capecitabine C
 Movicol C
 Senna C
 Spironolactone C
 Omeprazole C

Date:03/28/05ISR Number: 4622325-7Report Type:Expedited (15-DaCompany Report #2005043843
 Age:85 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Freedom Of Information (FOI) Report

Ramipril C
 Amlodipine Besilate C
 Pravastatin Sodium C
 Molsidomine C
 Paracetamol C
 Calcium D3 "Stada"
 (Calcium,
 Colecalciferol) C
 Alendronate Sodium C
 Glyceryl Trinitrate C
 Torasemide C
 Prednisone C
 Heparin C

Date:03/28/05ISR Number: 4622368-3Report Type:Expedited (15-DaCompany Report #2005045081
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Arteriosclerosis	Foreign	Norvasc (Amlodipine			
Other		Cerebral Haemorrhage	Health	Besilate)			
ORAL		Chest Pain	Professional	(Amlodipine)	PS		ORAL
ORAL		Duodenal Ulcer		Neurontin			
		Perforation		(Gabapentin)			
		Feeling Abnormal		(Gabapentin)	SS		ORAL
		Gastrointestinal		Trombyl Tabletter			
		Haemorrhage		(Acetylsalicylic			
		Pallor		Acid)	SS		ORAL
	75 MG (75 MG,	Shock					
	1 IN 1 D)						
ORAL				Diclofenac Sodium			
				(Diclofenac Sodium)	SS		ORAL
	150 MG (1 D)						
ORAL				Citalopram			
				Hydrobromide			
				(Citalopram			
				Hydrobromide)	SS		ORAL
	20 MG (20 MG,						

1 IN 1 D)

ORAL

Metoprolol Succinate C
Ramipril C

Date:03/28/05ISR Number: 4622686-9Report Type:Direct
Age:47 YR Gender:Female I/FU:I

Company Report #CTU 244491

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Inadequate Analgesia		Gabapentin -Generic			
		Pharmaceutical Product		400 Mg Caps	PS		ORAL
1200 MG PO		Complaint					
TID							

Date:03/28/05ISR Number: 4623079-0Report Type:Direct
Age:53 YR Gender:Female I/FU:I

Company Report #CTU 244419

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Convulsion		Neurontin 100 Mg			
Initial or Prolonged		Medication Error		Pfizer	PS	Pfizer	ORAL
100 MG BID		Pharmaceutical Product					
ORAL		Complaint					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4621567-4Report Type:Expedited (15-DaCompany Report #2005042689
 Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Headache Medication Error	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG (100 MG, 1 IN 1 D), ORAL		Petit Mal Epilepsy Stress		Carbamazepine (Carbamazepine)	C		

Date:03/29/05ISR Number: 4621570-4Report Type:Expedited (15-DaCompany Report #2005046040
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Cardiac Valve Disease Cardiac Ventricular Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
100 MG (100 MG, 1 IN 1 D), ORAL		Heart Rate Irregular					

Date:03/29/05ISR Number: 4621594-7Report Type:Expedited (15-DaCompany Report #2005045701
 Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dry Mouth	Consumer	Detrol La Capsule, Prolonged Release (Tolterodine L-Tartrate)	PS		ORAL
4 MG (4 MG, QD), ORAL				Neurontin (Gabapentin)	SS		ORAL
ORAL							

12.5 MG (12.5	Hydrochlorothiazide (Hydrochlorothiazide)	SS	ORAL
MG , QD),			
ORAL			
	Fenofibrate (Fenofibrate0	SS	ORAL
ORAL			
	Ramipril (Ramipril)	SS	ORAL
ORAL			
	Metoprolol Succinate (Metoprolol Succinate0	C	ORAL
ORAL			

Date:03/29/05ISR Number: 4622017-4Report Type:Expedited (15-DaCompany Report #2005046003
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Diarrhoea	Consumer	Phenytoin Suspension (Phenytoin Sodium)	PS		
300 MG		Nausea Transient Ischaemic Attack		Gabapentin (Gabapentin) Lipitor (Atorvastatin)	SS SS		

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4622106-4Report Type:Expedited (15-DaCompany Report #2004009355

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG (600 Other MG, 3 IN 1 D), ORAL	Abdominal Injury Abdominal Pain	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
20 MG (20 MG, 1 IN 1 D), ORAL	Back Disorder Blood Cholesterol Increased Chest Pain	Professional	Lipitor (Atorvastatin)	SS		ORAL
1000 MG (1 IN 2 D), ORAL	Contusion Coordination Abnormal Difficulty In Walking Disorientation Drug Ineffective Erectile Dysfunction Facial Pain Feeling Cold Gastrooesophageal Reflux Disease General Physical Health Deterioration Glossodynia Hostility Hyporeflexia Impaired Healing Injury Irritability Limb Injury Movement Disorder Muscle Spasms Myalgia Neck Pain Oesophageal Spasm Pain In Extremity Pain In Jaw Road Traffic Accident Sedation		Lithium (Lithium) (Lithium) Naproxen (Naproxen) Amoxicillin (Amoxicillin) (Amoxicillin)	SS SS SS		ORAL
			Rofecoxib (Rofecoxib) Levothyroxine Sodium (Levothyroxine Sodium) Vitamins (Vitamins) Diltiazem Hydrochloride (Diltiazem Hydrochloride)	SS C C C		

Sleep Disorder
Somnolence
Tenderness
Tooth Abscess
Tremor
Vertigo

Date:03/29/05ISR Number: 4622120-9Report Type:Expedited (15-DaCompany Report #001-0945-980292
Age:51 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Acute Respiratory
Initial or Prolonged	Distress Syndrome
Other	Adenoma Benign
Required	Anaemia
Intervention to	Anaesthetic Complication
Prevent Permanent	Neurological
Impairment/Damage	Cardiovascular
	Deconditioning
	Chest Pain
	Chronic Obstructive
	Pulmonary Disease
	Coagulopathy

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1200 MG (300 MG, 4 IN 1 D), ORAL		Delirium Depression Dysgeusia Dysuria Encephalopathy Epididymitis Feeling Abnormal Headache Hemiparesis Hypernatraemia Hypoglycaemia Hypokalaemia Influenza Like Illness Insomnia Malnutrition Muscle Atrophy Muscular Weakness Pain Pancreatitis Acute Pneumonia Mycoplasma Pneumonitis Renal Failure Respiratory Failure Sepsis Suicidal Ideation Swollen Tongue Upper Respiratory Tract Infection Viral Infection	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				All Other Therapeutic Products (All Other Therapeutic Products)	SS		
				Carisoprodol (Carisoprodol)	C		
				Hydrocodone (Hydrocodone)	C		
				Amitriptyline (Amitriptyline)	C		
				Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C		
				Doxepin (Doxepin)	C		
				Bromfenac Sodium (Bromfenac Sodium)	C		

Date:03/29/05ISR Number: 4622142-8Report Type:Expedited (15-DaCompany Report #2005045322
Age:71 YR Gender:Female I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, 3 IN 1)	Hospitalization - Initial or Prolonged	Blood Bilirubin Increased Cardio-Respiratory Arrest Hypersensitivity	Consumer	Neurontin (Gabapentin)	PS		ORAL

D), ORAL	Hypotension			
	Lung Disorder	Lipitor		
10 MG (10 MG,	Pain In Extremity	(Atorvastatin)	SS	ORAL
1 IN 1 D),	Pharmaceutical Product			
ORAL	Complaint			
	Thrombosis	Allopurinol		
		(Allopurinol)	C	
		Potassium		
		(Potassium)	C	
		Atenolol (Atenolol)	C	
		Rosiglitazone		
		Maleate		
		(Rosiglitazone		
		Maleate)	C	
		Magnesium		
		(Magnesium)	C	
		Insulin (Insulin)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4622169-6Report Type:Expedited (15-DaCompany Report #2005046470
 Age:56 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Agitation Confusional State	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
1200 MG (1200 MG, 1 IN 1 D) ORAL		Drug Toxicity Movement Disorder	Professional				
4 GRAM (4 GRAM, 1 IN 1 D) ORAL		Petit Mal Epilepsy Posture Abnormal		Capecitabine (Capecitabine)	SS		ORAL
				Omeprazole	C		
				Dexamethasone	C		
				Oxycodone			
				Hydrochloride	C		
				Fentanyl Citrate	C		

Date:03/29/05ISR Number: 4622170-2Report Type:Expedited (15-DaCompany Report #2005046333
 Age:50 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyspnoea Ear Pain	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
300 MG ORAL		Nasal Congestion Sleep Disorder	Professional	Valeriana Officinalis Melatonin	C C		

Date:03/29/05ISR Number: 4622789-9Report Type:Direct Company Report #CTU 244531
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Convulsion		Neurontin 800mg			

800 MG TID Pharmaceutical Product (Generic) PS
 Complaint

Date:03/29/05ISR Number: 4623069-8Report Type:Expedited (15-DaCompany Report #2005046321
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin			
		Nerve Injury		(Gabapentin)	PS		ORAL
600 MG (100		Pain					
MG, 3 IN 1		Pharmaceutical Product					
D), ORAL		Complaint		Vitamins (Vitamins)	C		

Date:03/29/05ISR Number: 4623255-7Report Type:Expedited (15-DaCompany Report #2005045559
 Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Condition Aggravated	Consumer	Neurontin			
Initial or Prolonged		Depression		(Gabapentin)	PS		ORAL
(100 MG, QOD)		Drug Ineffective					
Other		Hallucination		Nortriptyline			
ORAL		Pain		(Nortriptyline)	C		
		Pharmaceutical Product					
		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4623257-0Report Type:Expedited (15-DaCompany Report #2005014989
 Age:44 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	900 MG (300 MG, 3 IN 1 D), ORAL	Amnesia Balance Disorder Cervical Spinal Stenosis Depression Erectile Dysfunction Hypertension Hypoaesthesia	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL		Loss Of Employment Pain Pharmaceutical Product Complaint		Vicodin (Hydrocodone Bitartrate, Paracetamol) Pantoprazole (Pantoprazole) Nabumetone (Nabumetone) Potassium Chloride (Potassium Chloride) Furosemide (Furosemide) Tizanidine Hydrochloride (Tizanidine Hydrochloride)	SS C C C C C		ORAL

Date:03/29/05ISR Number: 4623267-3Report Type:Expedited (15-DaCompany Report #2005046480
 Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	100 MG (100 MG, 1 IN 1 D), ORAL	Amnesia Disorientation Drug Ineffective Physical Examination Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL				Gabapentin (Gabapentin) Lansprazole	SS		ORAL

(Lansprozole)	C
Lisinopril	
(Lisinopril)	C
Paracetamol	
(Paracetamol)	C
Atorvastatin	
(Atorvastatin)	C
Ibuprofen	
(Ibuprofen)	C
Fluoxetine	
Hydrochloride	
(Fluoxetine	
Hydrochloride)	C

Date:03/29/05ISR Number: 4623802-5Report Type:Direct
 Age:48 YR Gender:Female I/FU:I

Company Report #CTU 244640

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1200MG DAILY		Gun Shot Wound					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4623803-7Report Type:Direct
Age:43 YR Gender:Male I/FU:I

Company Report #CTU 244641

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury Asphyxiation		Neurontin	PS		

Date:03/29/05ISR Number: 4623815-3Report Type:Direct
Age:52 YR Gender:Male I/FU:I

Company Report #CTU 244642

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/29/05ISR Number: 4623816-5Report Type:Direct
Age:16 YR Gender:Male I/FU:I

Company Report #CTU 244643

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 100MG 3X DAILY		Completed Suicide Injury Asphyxiation		Neurontin	PS		

Date:03/29/05ISR Number: 4623817-7Report Type:Direct
Age:36 YR Gender:Male I/FU:I

Company Report #CTU 244644

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 300MG 3X DAILY		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/29/05ISR Number: 4623818-9Report Type:Direct
Age:59 YR Gender:Female I/FU:I

Company Report #CTU 244645

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG 3X	Completed Suicide		Neurontin	PS		
	DAILY	Gun Shot Wound					
				Ultracet Tablet	C		
				Amitriptyline Hcl	C		
				Sucralfate	C		
				Prevacid	C		
				Promethazine	C		

Date:03/29/05ISR Number: 4623819-0Report Type:Direct Company Report #CTU 244646
 Age:37 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Overdose		Prevacid	C		
				Zoloft	C		
				Claritan	C		

Date:03/29/05ISR Number: 4623820-7Report Type:Direct Company Report #CTU 244647
 Age:40 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	3200MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4623821-9Report Type:Direct
Age:43 YR Gender:Male I/FU:I

Company Report #CTU 244648

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200MG DAILY	Completed Suicide		Neurontin	PS		
		Gun Shot Wound		Naproxen	C		
				Cyclobenzaprine	C		

Date:03/29/05ISR Number: 4623822-0Report Type:Direct
Age:45 YR Gender:Male I/FU:I

Company Report #CTU 244649

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200MG, DAILY	Completed Suicide		Neurontin	PS		
		Injury Asphyxiation		Paxil	C		
				Trazodone	C		
				Humaloy	C		

Date:03/29/05ISR Number: 4623823-2Report Type:Direct
Age:44 YR Gender:Female I/FU:I

Company Report #CTU 244650

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		

Date:03/29/05ISR Number: 4623824-4Report Type:Direct
Age:66 YR Gender:Female I/FU:I

Company Report #CTU 244651

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG, DAILY, ORAL	Completed Suicide		Neurontin	PS		ORAL
		Gun Shot Wound		Duragesic	C		
				Mirtazapin	C		
				Aprazola	C		

Date:03/29/05ISR Number: 4623825-6Report Type:Direct
 Age:50 YR Gender:Male I/FU:I

Company Report #CTU 244652

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY, ORAL	Blood Carbon Monoxide Abnormal Completed Suicide		Neurontin Daypro Clonazepam	PS C C		ORAL

Date:03/29/05ISR Number: 4623826-8Report Type:Direct
 Age:34 YR Gender:Male I/FU:I

Company Report #CTU 244653

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury Asphyxiation		Neurontin	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4623827-XReport Type:Direct
 Age:47 YR Gender:Male I/FU:I

Company Report #CTU 244654

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
600 MG AM,		Gun Shot Wound					
600 MG PM 900		Head Injury					
MG BEDTIME							

Date:03/29/05ISR Number: 4623828-1Report Type:Direct
 Age:50 YR Gender:Male I/FU:I

Company Report #CTU 244655

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
400MG		Injury Asphyxiation					

Date:03/29/05ISR Number: 4623829-3Report Type:Direct
 Age:32 YR Gender:Male I/FU:I

Company Report #CTU 244656

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Injury Asphyxiation					

Date:03/29/05ISR Number: 4623830-XReport Type:Direct
 Age:51 YR Gender:Male I/FU:I

Company Report #CTU 244639

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
300MG, DAILY,		Gun Shot Wound					
ORAL							

Date:03/29/05ISR Number: 4623831-1Report Type:Direct Company Report #CTU 244638
Age:69 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1200MG DAILY		Gun Shot Wound					

Date:03/29/05ISR Number: 4623832-3Report Type:Direct Company Report #CTU 244637
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
900MG DAILY,		Gun Shot Wound					
ORAL				Bextra	C		

Date:03/29/05ISR Number: 4623833-5Report Type:Direct Company Report #CTU 244636
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1800MG DAILY,		Gun Shot Wound					
ORAL				Ultram	C		
				Oxaprozin	C		
				Mirapex	C		
				Lexpro	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4623834-7Report Type:Direct
 Age:34 YR Gender:Male I/FU:I

Company Report #CTU 244635

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
ORAL		Injury Asphyxiation					
				Valium	C		
				Zoloft	C		
				Lortab	C		
				Oxycotin	C		
				Flexeril	C		

Date:03/29/05ISR Number: 4623835-9Report Type:Direct
 Age:39 YR Gender:Female I/FU:I

Company Report #CTU 244634

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
ORAL		Gun Shot Wound					
				Propoxyphene	C		
				Allegra	C		
				Nortriptylin	C		
				Clonazepam	C		
				Amithriptycline	C		

Date:03/29/05ISR Number: 4623836-0Report Type:Direct
 Age:49 YR Gender:Male I/FU:I

Company Report #CTU 244633

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
ORAL		Injury Asphyxiation					
				Cardvra	C		
				Vicodin	C		
				Clonzepam	C		

Date:03/29/05ISR Number: 4623837-2Report Type:Direct
Age:52 YR Gender:Male I/FU:I

Company Report #CTU 244632

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
600MG DAILY,		Injury Asphyxiation					
ORAL				Xanax	C		
				Wellbutrum	C		
				Zestrill	C		
				Pravachol	C		

Date:03/29/05ISR Number: 4623838-4Report Type:Direct
Age:20 YR Gender:Male I/FU:I

Company Report #CTU 244631

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2400MG DAILY,							
ORAL				Celexra	C		
				Bupropion	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4623856-6Report Type:Direct
Age:34 YR Gender:Female I/FU:I

Company Report #CTU 244630

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		

Date:03/29/05ISR Number: 4623857-8Report Type:Direct
Age:40 YR Gender:Male I/FU:I

Company Report #CTU 244629

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG.	Completed Suicide		Neurontin	PS		ORAL
	DAILY, ORAL	Gun Shot Wound					

Date:03/29/05ISR Number: 4624190-0Report Type:Direct
Age:42 YR Gender:Female I/FU:I

Company Report #CTU 244627

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		
				Paxil	C		
				Zoloft	C		
				Methodone	C		

Date:03/29/05ISR Number: 4624191-2Report Type:Direct
Age:48 YR Gender:Male I/FU:I

Company Report #CTU 244628

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		

Date:03/29/05ISR Number: 4624205-XReport Type:Direct
Age:39 YR Gender:Female I/FU:I

Company Report #CTU 244624

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose		Zoloft	C		
				Zanx	C		
				Valuim	C		

Date:03/29/05ISR Number: 4624206-1Report Type:Direct Company Report #CTU 244625
Age:61 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	900 MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose		Ambien	C		
				Risperdal	C		
				Zonegran	C		

Date:03/29/05ISR Number: 4624207-3Report Type:Direct Company Report #CTU 244626
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Overdose					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4624231-0Report Type:Direct
Age:35 YR Gender:Male I/FU:I

Company Report #CTU 244620

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200MG DAILY, ORAL	Completed Suicide Overdose		Neurontin	PS		ORAL
				Zoloft	C		
				Zyprexa	C		
				Remeron	C		

Date:03/29/05ISR Number: 4624232-2Report Type:Direct
Age:67 YR Gender:I/FU:I

Company Report #CTU 244621

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG 3X DAILY	Completed Suicide Overdose		Neurontin	PS		
				Celebrex	C		
				Oxycodent	C		
				Tempazopan	C		
				Nortriptyline	C		

Date:03/29/05ISR Number: 4624233-4Report Type:Direct
Age:34 YR Gender:Female I/FU:I

Company Report #CTU 244622

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	900MG	Completed Suicide Overdose		Neurontin	PS		
				Topomax	C		
				Seroquel	C		
				Gabitril	C		
				Synarel	C		

Date:03/29/05ISR Number: 4624234-6Report Type:Direct
Age:39 YR Gender:Male I/FU:I

Company Report #CTU 244623

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Neurontin	PS		
1800 MG,		Overdose					
DAILY							

Date:03/29/05ISR Number: 4624236-XReport Type:Direct Company Report #CTU 244612
Age:36 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/29/05ISR Number: 4624237-1Report Type:Direct Company Report #CTU 244613
Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Neurontin	PS		ORAL
1200MG DAILY,		Overdose					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4624238-3Report Type:Direct
Age:47 YR Gender:Male I/FU:I

Company Report #CTU 244614

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300 MG 3X A	Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/29/05ISR Number: 4624239-5Report Type:Direct
Age:49 YR Gender:Female I/FU:I

Company Report #CTU 244615

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/29/05ISR Number: 4624240-1Report Type:Direct
Age:29 YR Gender:Male I/FU:I

Company Report #CTU 244616

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Overdose					
				Lithobid	C		
				Seroquel	C		
				Navaine	C		
				Klonopin	C		
				Cogentin	C		

Date:03/29/05ISR Number: 4624241-3Report Type:Direct
Age:51 YR Gender:Male I/FU:I

Company Report #CTU 244617

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2400MG DAILY	Completed Suicide		Neurontin	PS		
		Overdose		Mysoline	C		

Kepra C
Methadone C
Prevacid C

Date:03/29/05ISR Number: 4624242-5Report Type:Direct Company Report #CTU 244618
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY	Completed Suicide		Neurontin	PS		
		Overdose		Zeprexa	C		
				Effexor Xr	C		
				Trazodone	C		

Date:03/29/05ISR Number: 4624243-7Report Type:Direct Company Report #CTU 244619
Age:24 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800 MG	Completed Suicide		Neurontin	PS		ORAL
	DAILY, ORAL	Overdose		Prevacid	C		
				Effexor	C		
				Baclofen	C		
				Duragesic	C		
				Morphine Sulphate	C		
				Zonegran	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/29/05ISR Number: 4624247-4Report Type:Direct
Age:55 YR Gender:Female I/FU:I

Company Report #CTU 244607

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		

Date:03/29/05ISR Number: 4624248-6Report Type:Direct
Age:49 YR Gender:Female I/FU:I

Company Report #CTU 244608

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1500MG DAILY	Completed Suicide Overdose		Neurontin	PS		

Date:03/29/05ISR Number: 4624249-8Report Type:Direct
Age:38 YR Gender:Female I/FU:I

Company Report #CTU 244609

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG 3X DAILY	Completed Suicide Overdose		Neurontin	PS		
				Sulindac	C		
				Darvocet-N	C		
				Alprazolam	C		
				Bupropion Hcl	C		
				Phentermine	C		
				Acyclovir	C		
				Ranitidine	C		
				Levoxyl	C		

Date:03/29/05ISR Number: 4624250-4Report Type:Direct
Age:33 YR Gender:Female I/FU:I

Company Report #CTU 244610

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		

Date:03/29/05ISR Number: 4624251-6Report Type:Direct
 Age:32 YR Gender:Female I/FU:I

Company Report #CTU 244611

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound Head Injury		Neurontin	PS		

Date:03/29/05ISR Number: 4625975-7Report Type:Expedited (15-DaCompany Report #2005047536
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 4800 MG (800 MG, 6 IN 1 D), ORAL		Abdominal Pain Blood Alkaline Phosphatase Increased Blood Creatine Increased Diarrhoea Medication Error Pneumonia	Health Professional	Neurontin (Tablets) (Gabapentin) Hydromorphone Hydrochloride (Hydsromorphone Hydrochloride) Methadone	PS C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Methadone) C
 Citalopram
 Hydrobromide
 (Citalopram
 Hydrobromide) C
 Prochlorperazine
 Edisylate
 (Prochlorperazine
 Edisylate) C
 Levofloxacin
 (Levofloxacin) C
 Megestrol Acetate
 (Megestrol Acetate) C

Date:03/30/05ISR Number: 4623272-7Report Type:Expedited (15-DaCompany Report #2005040472
 Age:85 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Bronchitis	Consumer	Neurontin			
Hospitalization - ORAL		Diarrhoea	Health	(Gabapentin)	PS		ORAL
Initial or Prolonged		Gastrointestinal Haemorrhage Pharmaceutical Product Complaint Urinary Incontinence	Professional	Propacet (Dextropropoxyphene Napsilate, Paracetamol)	C		

Date:03/30/05ISR Number: 4623278-8Report Type:Expedited (15-DaCompany Report #2005042048
 Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (100 Other MG, 2 IN 1		Drug Ineffective Fatigue	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Vascular Occlusion					
D), ORAL				Metformin Hydrochloride (Metformin Hydrochloride)	C		

Pioglitazone
(Pioglitazone) C
Clopidogrel Sulfate
(Clopidogrel
Sulfate) C
Trazadone
(Trazadone) C
Atenolol (Atenolol) C
Vitamins (Vitamins) C
Prednisone
(Prednisone) C

Date:03/30/05ISR Number: 4623420-9Report Type:Expedited (15-DaCompany Report #2005042471

Age:71 YR Gender:Female I/FU:F

Outcome PT
Other Activities Of Daily
Living Impaired
Balance Disorder
Condition Aggravated
Deafness

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, 3 IN 1 D), ORAL		Drug Ineffective Headache Impaired Driving Ability	Consumer	Meurontin (Gabapentin)	PS		ORAL
		Medication Error Meniere'S Disease					
		Pharmaceutical Product Complaint					
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Triamterene (Triamterene)	C		

Date:03/30/05ISR Number: 4623432-5Report Type:Expedited (15-DaCompany Report #2005046314

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other ORAL		Abasia Arthritis	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Condition Aggravated		Bextra (Valdecoxib)	SS		ORAL
		Convulsion Decreased Activity Drug Ineffective Treatment Noncompliance Weight Increased		Gabapentin (Gabapentin)	SS		
				Omeprazole (Omeprazole)	C		
				Hyzaar (Hydrochlorothiazide , Losartan Potassium)	C		
				Raloxifene Hydrochloride (Raloxifene Hydrochloride)	C		
				Desipramine (Desipramine)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1800 MG (600 Other MG, 3 IN 1 D), ORAL	Coronary Artery Occlusion Drug Effect Decreased Gastric Disorder Gastrooesophageal Reflux Disease Lung Disorder Myocardial Infarction Nausea Urine Output Decreased	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Simvastatin (Simvastatin)	C		
			Metoprolol (Metoprolol)	C		
			Clopidogrel Sulfate (Clopidogrel Sulfate)	C		
			Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
			Dexamethasone (Dexamethasone)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4623783-4Report Type:Direct
Age:59 YR Gender:Female I/FU:I

Company Report #CTU 244687

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury Asphyxiation		Neurontin	PS		

Date:03/30/05ISR Number: 4623784-6Report Type:Direct
Age:26 YR Gender:Female I/FU:I

Company Report #CTU 244686

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4623785-8Report Type:Direct
Age:50 YR Gender:Male I/FU:I

Company Report #CTU 244685

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4623786-XReport Type:Direct
Age:25 YR Gender:Female I/FU:I

Company Report #CTU 244684

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	900-1800MG, DAILY, ORAL	Completed Suicide Gun Shot Wound		Neurontin	PS		ORAL
				Paxil	C		
				Zyprexa	C		

Date:03/30/05ISR Number: 4623787-1Report Type:Direct
Age:43 YR Gender:Male I/FU:I

Company Report #CTU 244683

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Completed Suicide Intentional Self-Injury		Neurontin	PS		

Date:03/30/05ISR Number: 4623789-5Report Type:Direct Company Report #CTU 244682
Age:51 YR Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Completed Suicide Intentional Self-Injury		Neurontin	PS		

Date:03/30/05ISR Number: 4623790-1Report Type:Direct Company Report #CTU 244681
Age:56 YR Gender:Male I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Completed Suicide		Neurontin	PS		ORAL
600 MG DAILY,		Gun Shot Wound					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4623791-3Report Type:Direct
Age:58 YR Gender:Male I/FU:I

Company Report #CTU 244680

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Carbon Monoxide Poisoning Completed Suicide		Neurontin	PS		

Date:03/30/05ISR Number: 4623792-5Report Type:Direct
Age:49 YR Gender:Male I/FU:I

Company Report #CTU 244679

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2700MG DAILY, ORAL	Completed Suicide Gun Shot Wound		Neurontin	PS		ORAL

Date:03/30/05ISR Number: 4623793-7Report Type:Direct
Age:55 YR Gender:Male I/FU:I

Company Report #CTU 244678

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2700MG	Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4623794-9Report Type:Direct
Age:37 YR Gender:Male I/FU:I

Company Report #CTU 244677

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1500MG	Completed Suicide Injury Asphyxiation		Neurontin	PS		

Date:03/30/05ISR Number: 4623795-0Report Type:Direct
Age:38 YR Gender:Male I/FU:I

Company Report #CTU 244676

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Completed Suicide		Neurontin	PS		
100MG DAILY		Gun Shot Wound					
Date:03/30/05ISR Number: 4623799-8Report Type:Direct			Company Report #CTU 244675				
Age:49 YR	Gender:Male	I/FU:I					
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Completed Suicide		Neurontin	PS		
		Gun Shot Wound					
Date:03/30/05ISR Number: 4623800-1Report Type:Direct			Company Report #CTU 244674				
Age:34 YR	Gender:Male	I/FU:I					
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Completed Suicide		Neurontin	PS		
		Road Traffic Accident					
Date:03/30/05ISR Number: 4623801-3Report Type:Direct			Company Report #CTU 244673				
Age:57 YR	Gender:Male	I/FU:I					
Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Death		Completed Suicide		Neurontin	PS		
		Gun Shot Wound					
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4623804-9Report Type:Direct
Age:28 YR Gender:Female I/FU:I

Company Report #CTU 244748

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Carbon Monoxide Poisoning		Neurontin	PS		
		Completed Suicide					

Date:03/30/05ISR Number: 4623805-0Report Type:Direct
Age:24 YR Gender:Male I/FU:I

Company Report #CTU 244749

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1800MG		Injury Asphyxiation					

Date:03/30/05ISR Number: 4623806-2Report Type:Direct
Age:45 YR Gender:Male I/FU:I

Company Report #CTU 244750

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
2400MG		Gun Shot Wound					

Date:03/30/05ISR Number: 4623807-4Report Type:Direct
Age:40 YR Gender:Female I/FU:I

Company Report #CTU 244751

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Carbon Monoxide Poisoning		Neurontin	PS		
2400MG		Completed Suicide					

Date:03/30/05ISR Number: 4623808-6Report Type:Direct
Age:58 YR Gender:Male I/FU:I

Company Report #CTU 244752

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
600MG DAILY,		Intentional Misuse					
ORAL				Pravachol	C		
				Oxazepam	C		

Date:03/30/05ISR Number: 4623809-8Report Type:Direct Company Report #CTU 244695
 Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2700MG DAILY,		Gun Shot Wound					
ORAL							

Date:03/30/05ISR Number: 4623810-4Report Type:Direct Company Report #CTU 244694
 Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
900MG DAILY,		Injury Asphyxiation					
ORAL				Diclofeuc	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4623811-6Report Type:Direct
Age:42 YR Gender:Female I/FU:I

Company Report #CTU 244877

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1600 MG, DAILY		Gun Shot Wound					
				Symbyaz	C		
				Zithromax	C		
				Nexium	C		
				Alprazolam	C		

Date:03/30/05ISR Number: 4623812-8Report Type:Direct
Age:53 YR Gender:Male I/FU:I

Company Report #CTU 244869

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1500MG DAILY, ORAL		Gun Shot Wound					
				Diovan	C		
				Starlex	C		
				Propoxx	C		

Date:03/30/05ISR Number: 4623813-XReport Type:Direct
Age:49 YR Gender:Male I/FU:I

Company Report #CTU 244868

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1800MG DAILY, ORAL		Gun Shot Wound					

Date:03/30/05ISR Number: 4623814-1Report Type:Direct
Age:54 YR Gender:Female I/FU:I

Company Report #CTU 244865

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
240MG		Injury Asphyxiation		Synthroid Klonopen	C C		

Date:03/30/05ISR Number: 4623858-XReport Type:Direct Company Report #CTU 244698
 Age:7 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
400 MG, DAILY, ORALY		Injury Asphyxiation		Zoloft	C		

Date:03/30/05ISR Number: 4623859-1Report Type:Direct Company Report #CTU 244699
 Age:29 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
600MG DAILY, ORAL		Gun Shot Wound		Vioxx Enbrel Vicodin Methotrexate	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4623860-8Report Type:Direct
Age:57 YR Gender:Female I/FU:I

Company Report #CTU 244700

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	3600MG DAILY,	Chest Injury		Neurontin	PS		ORAL
ORAL		Completed Suicide					
		Gun Shot Wound		Procardia	C		
				Celebrex	C		
				Vioxx	C		
				Robaxin	C		
				Ambien	C		
				Vicodin	C		
				K-Dur	C		

Date:03/30/05ISR Number: 4623861-XReport Type:Direct
Age:44 YR Gender:Male I/FU:I

Company Report #CTU 244708

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Gun Shot Wound					

Date:03/30/05ISR Number: 4623862-1Report Type:Direct
Age:18 YR Gender:Male I/FU:I

Company Report #CTU 244709

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
ORAL		Gun Shot Wound					
				Welbutrin	C		
				Alprazolam	C		
				Fluxetine	C		

Date:03/30/05ISR Number: 4623863-3Report Type:Direct
Age:39 YR Gender:Male I/FU:I

Company Report #CTU 244711

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1600MG DAILY,		Gun Shot Wound					
ORAL							

Date:03/30/05ISR Number: 4623875-XReport Type:Direct Company Report #CTU 244712
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2400MG DAILY,		Gun Shot Wound					
ORAL							

Date:03/30/05ISR Number: 4623876-1Report Type:Direct Company Report #CTU 244714
 Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1800MG DAILY,		Gun Shot Wound					
ORAL							

- Platal C
- Accupril C
- Prozac C
- Imdur C
- Lanoxin C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4623877-3Report Type:Direct
Age:52 YR Gender:Female I/FU:I

Company Report #CTU 244717

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2700MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Gun Shot Wound					

Date:03/30/05ISR Number: 4623878-5Report Type:Direct
Age:54 YR Gender:Male I/FU:I

Company Report #CTU 244719

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	900MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Gun Shot Wound		Phenytoin	C		

Date:03/30/05ISR Number: 4623889-XReport Type:Direct
Age:51 YR Gender:Female I/FU:I

Company Report #CTU 244723

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY	Completed Suicide		Neurontin	PS		
		Gun Shot Wound					

Date:03/30/05ISR Number: 4623896-7Report Type:Direct
Age:51 YR Gender:Female I/FU:I

Company Report #CTU 244724

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2400MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Gun Shot Wound					

Date:03/30/05ISR Number: 4623899-2Report Type:Direct Company Report #CTU 244726
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
900MG, DAILY,		Gun Shot Wound					
ORAL				Novolin	C		
				Lipitor	C		
				Ambien	C		

Date:03/30/05ISR Number: 4623906-7Report Type:Direct Company Report #CTU 244729
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2400MG DAILY,		Gun Shot Wound					
ORAL							

Date:03/30/05ISR Number: 4623916-XReport Type:Direct Company Report #CTU 244731
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1800MG DAILY,		Gun Shot Wound					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4623917-1Report Type:Direct
Age:49 YR Gender:Female I/FU:I

Company Report #CTU 244733

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2400MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Gun Shot Wound					

Date:03/30/05ISR Number: 4623918-3Report Type:Direct
Age:49 YR Gender:Male I/FU:I

Company Report #CTU 244738

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	3000MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Gun Shot Wound					
				Elevil	C		
				Promethazine	C		
				Macrobid	C		

Date:03/30/05ISR Number: 4623932-8Report Type:Direct
Age:31 YR Gender:Male I/FU:I

Company Report #CTU 244740

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY	Completed Suicide		Neurontin	PS		
		Gun Shot Wound					

Date:03/30/05ISR Number: 4623933-XReport Type:Direct
Age:60 YR Gender:Male I/FU:I

Company Report #CTU 244742

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Gun Shot Wound					

Date:03/30/05ISR Number: 4623934-1Report Type:Direct Company Report #CTU 244745
Age:25 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
3600MG DAILY,		Intentional Misuse					
ORAL							

Date:03/30/05ISR Number: 4623957-2Report Type:Direct Company Report #CTU 244746
Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2400MG DAILY,		Gun Shot Wound					
ORAL				Vioxx	C		
				Codeine	C		
				Diazepam	C		

Date:03/30/05ISR Number: 4623958-4Report Type:Direct Company Report #CTU 244747
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2400MG DAILY,		Gun Shot Wound					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4623959-6Report Type:Direct
Age:44 YR Gender:Female I/FU:I

Company Report #CTU 244909

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	900MG DAILY, ORAL	Completed Suicide Gun Shot Wound		Neurontin	PS		ORAL
				Clonazepam	C		
				Risperoal	C		

Date:03/30/05ISR Number: 4623960-2Report Type:Direct
Age:39 YR Gender:Male I/FU:I

Company Report #CTU 244908

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2400 MG DAILY, ORAL	Completed Suicide Injury Asphyxiation		Neurontin	PS		ORAL
				Clonazepam	C		
				Paxil	C		

Date:03/30/05ISR Number: 4623969-9Report Type:Direct
Age:59 YR Gender:Male I/FU:I

Company Report #CTU 244907

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	3200MG DAILY, ORAL	Completed Suicide Gun Shot Wound		Neurontin	PS		ORAL
				Zoloft	C		
				Trileptal	C		

Date:03/30/05ISR Number: 4623970-5Report Type:Direct
Age:33 YR Gender:Male I/FU:I

Company Report #CTU 244906

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Outcome	Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2700MG DAILY,		Completed Suicide		Neurontin	PS		ORAL
ORAL			Gun Shot Wound					
Date:03/30/05ISR Number: 4623971-7Report Type:Direct				Company Report #CTU 244905				
Age:44 YR Gender:Male I/FU:I								
Death	2700MG DAILY,		Completed Suicide		Neurontin	PS		ORAL
ORAL								
Date:03/30/05ISR Number: 4623972-9Report Type:Direct				Company Report #CTU 244904				
Age:44 YR Gender:Male I/FU:I								
Death	2700MG DAILY,		Completed Suicide		Neurontin	PS		ORAL
ORAL			Gun Shot Wound					
Date:03/30/05ISR Number: 4623973-0Report Type:Direct				Company Report #CTU 244903				
Age:42 YR Gender:Male I/FU:I								
Death	1800MG DAILY,		Completed Suicide		Neurontin	PS		ORAL
ORAL			Gun Shot Wound					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4623974-2Report Type:Direct
Age:40 YR Gender:Male I/FU:I

Company Report #CTU 244902

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Injury Asphyxiation					

Date:03/30/05ISR Number: 4623989-4Report Type:Direct
Age:27 YR Gender:Male I/FU:I

Company Report #CTU 244693

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL						

Date:03/30/05ISR Number: 4623990-0Report Type:Direct
Age:56 YR Gender:Male I/FU:I

Company Report #CTU 244692

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL	Injury Asphyxiation					

Date:03/30/05ISR Number: 4623991-2Report Type:Direct
Age:67 YR Gender:Male I/FU:I

Company Report #CTU 244691

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG	Completed Suicide		Neurontin	PS		
		Self Injurious Behaviour		Lamictal	C		
				Carizem	C		
				Seroquel	C		
				Wellbutrin	C		

Date:03/30/05ISR Number: 4623995-XReport Type:Direct
Age:46 YR Gender:Male I/FU:I

Company Report #CTU 244690

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4623996-1Report Type:Direct
Age:48 YR Gender:Male I/FU:I

Company Report #CTU 244689

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4623997-3Report Type:Direct
Age:42 YR Gender:Female I/FU:I

Company Report #CTU 244688

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4623998-5Report Type:Direct
 Age:63 YR Gender:Male I/FU:I

Company Report #CTU 244919

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
3000MG, DAILY, ORAL		Gun Shot Wound					
				Remoron	C		
				Celexra	C		
				Kloropin	C		

Date:03/30/05ISR Number: 4623999-7Report Type:Direct
 Age:70 YR Gender:Male I/FU:I

Company Report #CTU 244918

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Gun Shot Wound					

Date:03/30/05ISR Number: 4624072-4Report Type:Direct
 Age:24 YR Gender:Male I/FU:I

Company Report #CTU 244917

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Gun Shot Wound		Zoloft	C		
				Prolixin	C		

Date:03/30/05ISR Number: 4624073-6Report Type:Direct
 Age:48 YR Gender:Male I/FU:I

Company Report #CTU 244916

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		

Date:03/30/05ISR Number: 4624074-8Report Type:Direct
 Age:64 YR Gender:Male I/FU:I

Company Report #CTU 244915

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
900 MG DAILY,		Drowning					
ORAL							

Date:03/30/05ISR Number: 4624075-XReport Type:Direct Company Report #CTU 244914
Age:71 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1200MG DAILY,		Gun Shot Wound					
ORAL							

Date:03/30/05ISR Number: 4624076-1Report Type:Direct Company Report #CTU 244913
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2400MG DAILY,		Gun Shot Wound					
ORAL							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624077-3Report Type:Direct
Age:38 YR Gender:Male I/FU:I

Company Report #CTU 244912

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1800MG		Injury Asphyxiation					

Date:03/30/05ISR Number: 4624078-5Report Type:Direct
Age:75 YR Gender:Female I/FU:I

Company Report #CTU 244911

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1200MG DAILY		Gun Shot Wound		Labetalol	C		
				Triam	C		
				Amiticiptyline	C		

Date:03/30/05ISR Number: 4624079-7Report Type:Direct
Age:42 YR Gender:Male I/FU:I

Company Report #CTU 244910

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
2400MG		Injury Asphyxiation					

Date:03/30/05ISR Number: 4624081-5Report Type:Direct
Age:20 YR Gender:Male I/FU:I

Company Report #CTU 244901

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1800 MG		Intentional Self-Injury					
DAILY, ORAL							

Date:03/30/05ISR Number: 4624082-7Report Type:Direct Company Report #CTU 244900
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
2400MG DAILY		Injury Asphyxiation					

Date:03/30/05ISR Number: 4624083-9Report Type:Direct Company Report #CTU 244899
Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Gun Shot Wound					

Date:03/30/05ISR Number: 4624084-0Report Type:Direct Company Report #CTU 244898
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Gun Shot Wound					
		Head Injury					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624085-2Report Type:Direct
 Age:49 YR Gender:Male I/FU:I

Company Report #CTU 244897

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
3000 MG		Gun Shot Wound					
DAILY, ORAL							
0/2/02-07/21/							
02							

Date:03/30/05ISR Number: 4624086-4Report Type:Direct
 Age:56 YR Gender:Male I/FU:I

Company Report #CTU 244896

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
600 MG DAILY,		Gun Shot Wound					
ORAL							

Date:03/30/05ISR Number: 4624087-6Report Type:Direct
 Age:44 YR Gender:Male I/FU:I

Company Report #CTU 244895

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Gun Shot Wound					

Date:03/30/05ISR Number: 4624088-8Report Type:Direct
 Age:46 YR Gender:Male I/FU:I

Company Report #CTU 244894

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Gun Shot Wound					

Date:03/30/05ISR Number: 4624089-XReport Type:Direct
Age:72 YR Gender:Male I/FU:I

Company Report #CTU 244893

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Gun Shot Wound					

Date:03/30/05ISR Number: 4624090-6Report Type:Direct
Age:59 YR Gender:Male I/FU:I

Company Report #CTU 244892

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
900MG		Gun Shot Wound		Plavix	C		
				Micardis	C		
				Zocor	C		
				St. Joseph Aspirin	C		
				Alprazolam	C		
				Temazepam	C		
				Serzone	C		
				Effexor	C		
				Androgel	C		
				Niaspan	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624091-8Report Type:Direct
Age:37 YR Gender:Female I/FU:I

Company Report #CTU 244891

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4624092-XReport Type:Direct
Age:36 YR Gender:Male I/FU:I

Company Report #CTU 244890

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2400 DAILY, ORAL	Completed Suicide Injury Asphyxiation		Neurontin	PS		ORAL
				Zyprexa	C		
				Syntxsid	C		
				Serezone	C		

Date:03/30/05ISR Number: 4624093-1Report Type:Direct
Age:67 YR Gender:Male I/FU:I

Company Report #CTU 244889

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2400MG DAILY, ORAL	Completed Suicide Gun Shot Wound		Neurontin	PS		ORAL

Date:03/30/05ISR Number: 4624094-3Report Type:Direct
Age:39 YR Gender:Female I/FU:I

Company Report #CTU 244888

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY, ORAL	Completed Suicide		Neurontin	PS		ORAL
				Methadone	C		

Oxycontin C
Klonopin C
Effexor Xr C

Date:03/30/05ISR Number: 4624095-5Report Type:Direct Company Report #CTU 244887
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1800MG DAILY,		Gun Shot Wound					
ORAL							

Date:03/30/05ISR Number: 4624096-7Report Type:Direct Company Report #CTU 244886
Age:26 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1600MG DAILY,		Gun Shot Wound					
ORAL				Tizanidine	C		
				Ketorolac	C		
				Percocet	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624097-9Report Type:Direct
Age:40 YR Gender:Male I/FU:I

Company Report #CTU 244885

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	3000MG DAILY, ORAL	Completed Suicide Gun Shot Wound		Neurontin	PS		ORAL

Date:03/30/05ISR Number: 4624098-0Report Type:Direct
Age:41 YR Gender:Female I/FU:I

Company Report #CTU 244884

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		

Date:03/30/05ISR Number: 4624099-2Report Type:Direct
Age:53 YR Gender:Male I/FU:I

Company Report #CTU 244883

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4624100-6Report Type:Direct
Age:56 YR Gender:Male I/FU:I

Company Report #CTU 244882

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	900-1200MG, DAILY, ORAL	Completed Suicide Gun Shot Wound		Neurontin	PS		ORAL

Date:03/30/05ISR Number: 4624101-8Report Type:Direct
Age:48 YR Gender:Female I/FU:I

Company Report #CTU 244881

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
SEE IMAGE		Gun Shot Wound					

Date:03/30/05ISR Number: 4624102-XReport Type:Direct Company Report #CTU 244880
Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
900MG DAILY, ORAL		Gun Shot Wound					
				Celebrex	C		
				Enalapril	C		

Date:03/30/05ISR Number: 4624103-1Report Type:Direct Company Report #CTU 244879
Age:52 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
2700MG		Gun Shot Wound					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624104-3Report Type:Direct
Age:53 YR Gender:Male I/FU:I

Company Report #CTU 244878

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800 MG,	Completed Suicide		Neurontin	PS		
	DAILY	Gun Shot Wound		Ultracet	C		
				Glyburide	C		

Date:03/30/05ISR Number: 4624192-4Report Type:Direct
Age:39 YR Gender:Female I/FU:I

Company Report #CTU 244657

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		

Date:03/30/05ISR Number: 4624193-6Report Type:Direct
Age:44 YR Gender:Female I/FU:I

Company Report #CTU 244658

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		

Date:03/30/05ISR Number: 4624194-8Report Type:Direct
Age:37 YR Gender:Male I/FU:I

Company Report #CTU 244659

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG	Convulsion		Neurontin	PS		

Date:03/30/05ISR Number: 4624195-XReport Type:Direct
Age:43 YR Gender:Female I/FU:I

Company Report #CTU 244660

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1800 MG,		Overdose					
DAILY ORAL							

Date:03/30/05ISR Number: 4624198-5Report Type:Direct Company Report #CTU 244661
Age:43 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
2400MG		Overdose					

Date:03/30/05ISR Number: 4624199-7Report Type:Direct Company Report #CTU 244662
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1800MG		Overdose					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624200-0Report Type:Direct
Age:38 YR Gender:Male I/FU:I

Company Report #CTU 244668

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2400MG DAILY	Completed Suicide		Neurontin	PS		
		Overdose		Oxycontin	C		
				Relefen	C		

Date:03/30/05ISR Number: 4624201-2Report Type:Direct
Age:49 YR Gender:Male I/FU:I

Company Report #CTU 244669

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/30/05ISR Number: 4624202-4Report Type:Direct
Age:34 YR Gender:Male I/FU:I

Company Report #CTU 244670

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1800MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORA02L	Overdose		Vioxx	C		
				Carisoprodol	C		
				Ferrous	C		

Date:03/30/05ISR Number: 4624203-6Report Type:Direct
Age:63 YR Gender:Female I/FU:I

Company Report #CTU 244671

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200MG	Completed Suicide		Neurontin	PS		
		Overdose					

Date:03/30/05ISR Number: 4624204-8Report Type:Direct
Age:43 YR Gender:Male I/FU:I

Company Report #CTU 244672

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		

Date:03/30/05ISR Number: 4624213-9Report Type:Direct
Age:19 YR Gender:Female I/FU:I

Company Report #CTU 244665

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 2400MG 3X DAILY		Completed Suicide Overdose		Neurontin	PS		

Date:03/30/05ISR Number: 4624214-0Report Type:Direct
Age:42 YR Gender:Male I/FU:I

Company Report #CTU 244666

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 2000MG, DAILY, ORAL		Completed Suicide Overdose		Neurontin	PS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624215-2Report Type:Direct
Age:41 YR Gender:Male I/FU:I

Company Report #CTU 244667

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1800MG		Overdose					

Date:03/30/05ISR Number: 4624244-9Report Type:Direct
Age:32 YR Gender:Male I/FU:I

Company Report #CTU 244663

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1600 MG DAILY		Overdose					

Date:03/30/05ISR Number: 4624245-0Report Type:Direct
Age:52 YR Gender:Female I/FU:I

Company Report #CTU 244664

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1800MG		Overdose					

Date:03/30/05ISR Number: 4624252-8Report Type:Direct
Age:48 YR Gender:Female I/FU:I

Company Report #CTU 244876

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
400 MG DAILY, ORAL		Injury Asphyxiation					

Date:03/30/05ISR Number: 4624260-7Report Type:Direct
Age:33 YR Gender:Male I/FU:I

Company Report #CTU 244866

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
2400MG,		Intentional Misuse					
DAILY, ORAL				Inderal	C		
				Imipramine	C		

Date:03/30/05ISR Number: 4624261-9Report Type:Direct Company Report #CTU 244867
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
1200 MG		Intentional Misuse					
DAILY, ORAL				Lorazepam	C		
				Zithromax	C		
				Prevacid	C		
				Promethazine	C		

Date:03/30/05ISR Number: 4624262-0Report Type:Direct Company Report #CTU 244870
 Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624263-2Report Type:Direct
Age:56 YR Gender:Male I/FU:I

Company Report #CTU 244871

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	3600 MG	Completed Suicide		Neurontin	PS		ORAL
	DAILY, ORAL	Gun Shot Wound					

Date:03/30/05ISR Number: 4624264-4Report Type:Direct
Age:14 YR Gender:Male I/FU:I

Company Report #CTU 244872

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	900 MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
	ORAL						

Date:03/30/05ISR Number: 4624265-6Report Type:Direct
Age:17 YR Gender:Male I/FU:I

Company Report #CTU 244873

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1600MG DAILY	Completed Suicide		Neurontin	PS		
				Desoxyn	C		
				Remeron	C		
				Wellbutrin	C		
				Ambien	C		
				Effexer	C		

Date:03/30/05ISR Number: 4624266-8Report Type:Direct
Age:44 YR Gender:Male I/FU:I

Company Report #CTU 244874

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Gun Shot Wound		Carbatrol	C		
				Baclofen	C		

Date:03/30/05ISR Number: 4624267-XReport Type:Direct
 Age:21 YR Gender:Male I/FU:I

Company Report #CTU 244875

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4624268-1Report Type:Direct
 Age:47 YR Gender:Female I/FU:I

Company Report #CTU 244856

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL

1800MG,
DAILY, ORAL

Date:03/30/05ISR Number: 4624269-3Report Type:Direct
 Age:41 YR Gender:Female I/FU:I

Company Report #CTU 244857

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL

ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624270-XReport Type:Direct
Age:34 YR Gender: I/FU:I

Company Report #CTU 244858

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4624271-1Report Type:Direct
Age:36 YR Gender:Male I/FU:I

Company Report #CTU 244859

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4624272-3Report Type:Direct
Age:54 YR Gender:Male I/FU:I

Company Report #CTU 244860

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4624273-5Report Type:Direct
Age:39 YR Gender:Male I/FU:I

Company Report #CTU 244861

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1500MG DAILY	Completed Suicide Injury Asphyxiation		Neurontin	PS		

Date:03/30/05ISR Number: 4624274-7Report Type:Direct
Age:39 YR Gender:Female I/FU:I

Company Report #CTU 244863

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death Completed Suicide Neurontin PS ORAL
3200MG DAILY,
Overdose
ORAL

Date:03/30/05ISR Number: 4624275-9Report Type:Direct Company Report #CTU 244864
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
900MG DAILY, ORAL		Overdose					

Date:03/30/05ISR Number: 4624276-0Report Type:Direct Company Report #CTU 244848
Age:22 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
4800MG DAILY		Injury Asphyxiation		Tequin	C		
				Albuteral	C		
				Remeron	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624277-2Report Type:Direct
Age:55 YR Gender:Male I/FU:I

Company Report #CTU 244849

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
1200MG		Gun Shot Wound		Atenol	C		
				Lamotrigine	C		
				Trazodone	C		

Date:03/30/05ISR Number: 4624278-4Report Type:Direct
Age:36 YR Gender:Male I/FU:I

Company Report #CTU 244850

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
2400MG DAILY		Gun Shot Wound		Celexa	C		
				Lorazepam	C		
				Buspar	C		
				Flomax	C		
				Lorcet	C		

Date:03/30/05ISR Number: 4624279-6Report Type:Direct
Age:32 YR Gender:Female I/FU:I

Company Report #CTU 244851

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
2400MG DAILY		Gun Shot Wound					
		Head Injury					

Date:03/30/05ISR Number: 4624280-2Report Type:Direct
Age:47 YR Gender:Male I/FU:I

Company Report #CTU 244852

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Injury Asphyxiation					

Date:03/30/05ISR Number: 4624281-4Report Type:Direct
Age:25 YR Gender:Male I/FU:I

Company Report #CTU 244853

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury Asphyxiation		Neurontin	PS		

Date:03/30/05ISR Number: 4624282-6Report Type:Direct
Age:44 YR Gender:Male I/FU:I

Company Report #CTU 244854

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2400MG DAILY	Completed Suicide Gun Shot Wound Head Injury		Neurontin	PS		

Date:03/30/05ISR Number: 4624283-8Report Type:Direct
Age:44 YR Gender:Male I/FU:I

Company Report #CTU 244855

Outcome	PT
Death	Completed Suicide Gun Shot Wound

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Head Injury

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2400MG DAILY			Neurontin	PS		

Date:03/30/05ISR Number: 4624284-XReport Type:Direct
 Age:48 YR Gender:Male I/FU:I Company Report #CTU 244840

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury Asphyxiation		Neurontin	PS		

Date:03/30/05ISR Number: 4624285-1Report Type:Direct
 Age:22 YR Gender: I/FU:I Company Report #CTU 244841

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury Asphyxiation		Neurontin	PS		

Date:03/30/05ISR Number: 4624286-3Report Type:Direct
 Age:33 YR Gender:Male I/FU:I Company Report #CTU 244842

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4624287-5Report Type:Direct
 Age:48 YR Gender:Male I/FU:I Company Report #CTU 244843

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Gun Shot Wound		Neurontin	PS		

Date:03/30/05ISR Number: 4624288-7Report Type:Direct Company Report #CTU 244844
Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Self Injurious Behaviour		Neurontin	PS		

Date:03/30/05ISR Number: 4624289-9Report Type:Direct Company Report #CTU 244845
Age:13 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Injury Asphyxiation		Neurontin	PS		

Date:03/30/05ISR Number: 4624290-5Report Type:Direct Company Report #CTU 244846
Age:48 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	300MG 3X	Completed Suicide		Neurontin	PS		
DAILY		Gun Shot Wound		Oxycontin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624291-7Report Type:Direct
Age:56 YR Gender:Male I/FU:I

Company Report #CTU 244847

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		ORAL
3600MG, DAILY, ORAL		Gun Shot Wound		Prozac	C		
				Baclofen	C		
				Klonopin	C		

Date:03/30/05ISR Number: 4624345-5Report Type:Direct
Age:47 YR Gender:Female I/FU:I

Company Report #CTU 244783

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Feeling Abnormal		Gabapentin 600 Mg			
2 TID		Gastric Disorder		Teva	PS	Teva	

Date:03/30/05ISR Number: 4624380-7Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 244791

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Dizziness		Gabapentine 400 Mg			
400 MG PO TID		Feeling Abnormal		Tid	PS		ORAL

Date:03/30/05ISR Number: 4624427-8Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 244720

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Activities Of Daily		Neurontin 800mg Tid	PS		ORAL
800MG TID PO		Living Impaired					
		Pharmaceutical Product					
		Complaint					

Sedation

Date:03/30/05ISR Number: 4624439-4Report Type:Direct
Age:58 YR Gender:Male I/FU:I

Company Report #CTU 244832

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Overdose		Neurontin	PS		

Date:03/30/05ISR Number: 4624446-1Report Type:Direct
Age:53 YR Gender:Female I/FU:I

Company Report #CTU 244834

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200 MG DAILY, ORAL	Completed Suicide Overdose		Neurontin	PS		ORAL

Date:03/30/05ISR Number: 4624448-5Report Type:Direct
Age:52 YR Gender:Male I/FU:I

Company Report #CTU 244836

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	600MG DAILY, ORAL	Completed Suicide Overdose		Neurontin	PS		ORAL
				Contin Baclofen	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4624450-3Report Type:Direct
Age:44 YR Gender:Female I/FU:I

Company Report #CTU 244837

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	2400MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
ORAL		Intentional Misuse					
				Celebrex	C		
				Vioxx	C		
				Premarin	C		
				Meperidine	C		

Date:03/30/05ISR Number: 4624451-5Report Type:Direct
Age:42 YR Gender:Male I/FU:I

Company Report #CTU 244838

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	1200MG DAILY,	Completed Suicide		Neurontin	PS		ORAL
ORAL		Overdose					
				Clonazepam	C		
				Coumadin	C		
				Prevacid	C		

Date:03/30/05ISR Number: 4624452-7Report Type:Direct
Age:31 YR Gender:Male I/FU:I

Company Report #CTU 244839

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin	PS		
		Intentional Misuse					

Date:03/30/05ISR Number: 4625904-6Report Type:Expedited (15-DaCompany Report #2005044232
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Alopecia	Consumer	Neurontin			

800 MG (1 D),	Bladder Prolapse	(Gabapentin)	PS	ORAL
ORAL	Breath Odour			
	Drug Ineffective	Cyclobenzaprine		
	Dysgeusia	(Cyclobenzaprine)	C	
	Gangrene	Thyroid (Thyroid)	C	
	Hair Disorder	All Other		
	Hypoaesthesia	Therapeutic Products	C	
	Malaise	Clonazepam		
	Medication Error	(Clonazepam)	C	
	Oral Mucosal Disorder	Rosiglitazone		
	Panic Attack	Maleate		
	Tooth Discolouration	(Rosiglitazone		
	Tooth Loss	Maleate)	C	
		Naproxen (Naproxen0	C	
		Simvastatin		
		(Simvastatin)	C	

Date:03/30/05ISR Number: 4626072-7Report Type:Expedited (15-DaCompany Report #2005049540
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Brain Operation	Foreign	Neurontin			
		Hot Flush	Health	(Gabapentin)	PS		ORAL
1600 MG (800			Professional				
MG, 2 IN 1							
D), ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/30/05ISR Number: 4626124-1Report Type:Expedited (15-DaCompany Report #2005047806
Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Road Traffic Accident	Consumer	Neurontin (Gabapentin)	PS		

Date:03/31/05ISR Number: 4626951-0Report Type:Expedited (15-DaCompany Report #2004030610
Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1500 MG (3 IN Disability 1 D), ORAL Other 200 MG (200 MG, 1 IN 1 D), ORAL		Arthralgia Muscular Weakness Myalgia Myopathy Nuclear Magnetic Resonance Imaging Abnormal Oedema Peripheral Pain In Extremity Walking Aid User	Foreign Consumer Health Professional	Neurontin (Gabapentin) Celebrex (Celecoxib) Diclofenac Sodium (Diclofenac Sodium) Venostasin Forte (Horse Chestnut Extract Thiamine Hydrochloride) All Other Therapeutic Products (All Other Therapeutic Products) Biomagnesium (Citric Acid, Magnesium Citrate, Magnesium Phosphate Dibasic)	PS SS C C C C		ORAL ORAL

Date:03/31/05ISR Number: 4627001-2Report Type:Expedited (15-DaCompany Report #2005049630
Age:59 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation Pain	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
900 MG (300 MG 3 IN 1 D)		Paraesthesia	Professional				
ORAL			Other				

Date:03/31/05ISR Number: 4627074-7Report Type:Expedited (15-DaCompany Report #2005048413
Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Abdominal Pain Upper Amnesia	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG (600 Other MG, 3 IN 1 D), ORAL		Deafness Unilateral Dizziness					
ORAL		Tinnitus Transient Ischaemic Attack		Sinemet (Carbidopa, Levodopa)	SS		ORAL
				Antihypertensives (Antihypertensives)	C		
				Anti-Diabetics (Anti-Diabetics)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/31/05ISR Number: 4627102-9Report Type:Expedited (15-DaCompany Report #2005043174
Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 MG, 2 IN 1 D), ORAL		Gastric Disorder - Paraesthesia Pharmaceutical Product Complaint	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Allopurinol (Allopurinol)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Omeprazole (Omeprazole)	C		

Date:03/31/05ISR Number: 4627104-2Report Type:Expedited (15-DaCompany Report #2005016291
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Memory Impairment	Consumer	Neurontin (Gabapentin)	PS		
				Diazepam (Diazepam)	SS		
				Ibuprofen (Ibuprofen)	SS		
				Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		
				Panadeine Co (Codeine Phosphate, Paracetamol)	SS		
				Atorvastatin (Atorvastatin)	C		
				Bupropion Hydrochloride (Bupropion Hydrochloride)	C		
				Fluconazole (Fluconazole)	C		

Date:03/31/05ISR Number: 4627108-XReport Type:Expedited (15-DaCompany Report #2005049636
Age:18 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Clonic Convulsion Partial Seizures	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG, 2 OR							
3 CAPSULES							
EVERY 8							
HOURS, ORAL							

Date:03/31/05ISR Number: 4627113-3Report Type:Expedited (15-DaCompany Report #2005048434
Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide		Neurontin			
Other		Drug Ineffective		(Gabapentin)	PS		
UNKNOWN	UNKNOWN						

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:03/31/05ISR Number: 4627116-9Report Type:Expedited (15-DaCompany Report #2005046944

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 20 MG (20 MG, Initial or Prolonged 1 IN 1 D), Other ORAL		Aggression	Consumer	Bextra (Valdecoxib)	PS		ORAL
UNK (1 D), ORAL		Anticonvulsant Drug Level Decreased		Xanax Tablet (Alprazolam)	SS		ORAL
500 MG (1 D), ORAL		Blood Pressure Increased Chest Pain		Dilantin (Phenytoin Sodium)	SS		ORAL
ORAL		Choking Sensation					
UNKNOWN, ORAL		Drug Effect Decreased Drug Interaction		Neurontin (Gabapentin)	SS		ORAL
UNKNOWN	UNKNOWN	Flatulence		Methadone (Methadone)	SS		
UNKNOWN	UNKNOWN	Gastritis Throat Tightness		Topiramate (Topiramate)	SS		
				Axotal (Old Form) (Butalbital, Caffeine, Paracetamol)	C		

Date:03/31/05ISR Number: 4627123-6Report Type:Expedited (15-DaCompany Report #2005048409

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other UNKNOWN		Anxiety Completed Suicide	Consumer	Neurontin (Gabapentin)	PS		
	UNKNOWN	Drug Ineffective Suicidal Ideation					

Suicide Attempt

Date:04/01/05ISR Number: 4626822-XReport Type:Direct
 Age:44 YR Gender:Male I/FU:I

Company Report #CTU 245011

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Death		Neurontin	PS		
2 MON		Circulatory Collapse		Oxycontin	SS		
		Dehydration		Oxycodone	SS		
		Drug Interaction		Effexor	SS		
		Drug Toxicity		Alprazolam	SS		
		Incoherent					
		Loss Of Consciousness					

Date:04/01/05ISR Number: 4626890-5Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 245087

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion		Neurontin (Generic)			
Other		Pharmaceutical Product		600 Mg Two Po Qhs	PS		ORAL
60 MG TWO PO		Complaint					
QHS							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/01/05ISR Number: 4627046-2Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 245063

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Nightmare		Generic Neurontin			
600MG	TID	Pharmaceutical Product		600mg	PS		ORAL
PO		Complaint					

Date:04/01/05ISR Number: 4627632-XReport Type:Expedited (15-DaCompany Report #2005049181
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Death	Consumer	Neurontin			
Other		Circulatory Collapse		(Gabapentin)	PS		ORAL
1 IN 1 D,		Confusional State					
ORAL		Drug Toxicity		Oxycodone			
		Hepatic Steatosis		(Oxycodone)	SS		
		Hypotonia		Oxycontin (Oxycodone			
		Lung Disorder		Hydrochloride)	SS		
		Unevaluable Event		Effexor (Venlafaxine			
		Vision Blurred		Hydrochloride)	SS		
				Alprazolam			
				(Alprazolam)	C		

Date:04/01/05ISR Number: 4627962-1Report Type:Expedited (15-DaCompany Report #2005045983
 Age:30 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Circulatory Collapse	Foreign	Neurontin			
900 MG (300		Loss Of Consciousness	Health	(Gabapentin)	PS		ORAL
MG, 3 IN 1		Pharmaceutical Product	Professional				
D), ORAL		Complaint	Company				
			Representative	Ethanol			

Other

(Ethanol) SS
Amitriptyline
Hydrochloride
(Amitriptyline
Hydrochloride) C

Date:04/01/05ISR Number: 4628210-9Report Type:Expedited (15-DaCompany Report #2005038390

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged	Carpal Tunnel Syndrome	Consumer	Celebrex (Celecoxib)	PS		
Other 100 MG; 200 MG	Cataract	Health	Neurontin			
	Knee Operation	Professional	(Gabapentin)	SS		
	Neuropathy Peripheral					
	Poor Peripheral Circulation		Bextra (Valdecoxib)	SS		
	Post Procedural Complication		Bengay (Menthol)	SS		
	Staphylococcal Infection		Paracetamol			
	Streptococcal Infection		(Paracetamol)	C		
	Surgery		Analgesics			
			(Analgesics)	C		

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Freedom Of Information (FOI) Report

Date:04/01/05ISR Number: 4628226-2Report Type:Expedited (15-DaCompany Report #2005042488
Age:84 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG, 3 IN 1 D), ORAL		Activities Of Daily Living Impaired Confusional State Drug Ineffective Eating Disorder Neuralgia Pain In Jaw Pharmaceutical Product Complaint Speech Disorder Weight Decreased	Consumer	Neurontin (Gapapentin) Oxycocet (Oxycodone Hydrochloride, Paracetamol) Diltiazem (Diltiazem) Centrum (Minerals Nos, Vitamins Nos)	PS SS C C		ORAL

Date:04/01/05ISR Number: 4628234-1Report Type:Expedited (15-DaCompany Report #2005049071
Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other (100 MG, UP TO THREE TIMES A DAY)		Drug Effect Decreased Impaired Healing	Consumer	Neurontin (Gabapentin) Lotrel (Amlodipine, Benazepril Hydrochloride) Atenolol (Atenolol) Celecoxib (Celecoxib)	PS C C C		

Date:04/01/05ISR Number: 4628277-8Report Type:Expedited (15-DaCompany Report #2005049653
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Coma	Consumer	Neurontin			
Other		Nervous System Disorder		(Gabapentin)	PS		
(100 MG)		Pulmonary Embolism					

Date:04/01/05ISR Number: 4628284-5Report Type:Expedited (15-DaCompany Report #2005049411
Age:75 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cerebrovascular Accident	Consumer	Neurontin			
Other		Memory Impairment		(Gabapentin)	PS		ORAL
ORAL		Myocardial Infarction		Warfarin (Warfarin)	C		
				Metformin			
				Hydrochloride			
				(Metformin			
				Hydrochloride)	C		
				Glipizide			
				(Glipizide)	C		
				Quinapril			
				Hydrochloride			
				(Quinapril			
				Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/04/05ISR Number: 4628445-5Report Type:Expedited (15-DaCompany Report #2005036506
 Age:76 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 2400 MG (800 MG, 3 IN 1 D), ORAL 7-8 YEARS AGO		Condition Aggravated Diabetes Mellitus Drug Effect Decreased Gallbladder Operation Insomnia	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG (800 MG, 3 IN 1 D), ORAL		Neuropathy Peripheral Paraesthesia Pharmaceutical Product Complaint		Gabapentin (Tablets) (Gabapentin)	SS		ORAL
				Verapamil Hydrochloride (Verapamil Hydrochloride) Hydrochlorothiazide (Hydrochlorothiazide) Estrogens Conjugated (Estrogens Conjugated) Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotinamide, Lekovit Ca (Calcium Carbonate, Colecalciferol) Omeprazole (Omeprazole)	C C C C C C		

Date:04/04/05ISR Number: 4628488-1Report Type:Expedited (15-DaCompany Report #2005049031
 Age:82 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Grand Mal Convulsion	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Donepezil Hydrochloride (Donepezil Hydrochloride)	PS C		

Date:04/04/05ISR Number: 4629049-0Report Type:Direct
 Age: Gender:Female I/FU:I Company Report #CTU 245098

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Asthenia Pharmaceutical Product Complaint		Neurontin	PS		

Date:04/04/05ISR Number: 4629065-9Report Type:Direct
 Age:46 YR Gender:Male I/FU:I Company Report #CTU 245188

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 400MG TWICE		No Adverse Effect		Neurontin 400mg	PS		

DAILY,1200 MG
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Freedom Of Information (FOI) Report

BEDTIME

Date:04/04/05ISR Number: 4629195-1Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 245103

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Condition Aggravated		Gabapentin	PS		ORAL
1 PO TID	6 WK	Hyperaesthesia Pain Pharmaceutical Product Complaint		Elavil	C		

Date:04/04/05ISR Number: 4629197-5Report Type:Direct
 Age:63 YR Gender:Female I/FU:I

Company Report #CTU 245105

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Insomnia		Gabapentin 300 Mg	PS		ORAL
1 QAM, 1 Q		Pharmaceutical Product Complaint					
NOON 2 QHS		Restlessness		Wellbutrin Sr	C		
ALL PO							

Date:04/04/05ISR Number: 4629251-8Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 245111

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysphagia		Gabapentin	PS		ORAL
600 MG 2 BID		Pharmaceutical Product Complaint					
PO							

Date:04/04/05ISR Number: 4629278-6Report Type:Direct Company Report #CTU 245141
Age:34 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Atrial Fibrillation		Gabapentin 300 (Teva)	PS	Teva	ORAL
300 MG PO				Celebrex Flexeril	C C		

Date:04/04/05ISR Number: 4629281-6Report Type:Direct Company Report #CTU 245386
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated		Gabapentin 300 Mg	PS		
ONE QID		Neuropathy Peripheral Pharmaceutical Product Complaint					

Date:04/05/05ISR Number: 4627160-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0376424A
Age:69 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Abnormal Dreams Cognitive Disorder Confusional State Disorientation Hallucination, Visual

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Hypersomnia Incoherent Metabolic Encephalopathy					
8MG per day	12 DAY	Psychomotor Retardation		Zophren	PS	Glaxosmithkline	ORAL
INTRAVENOUS	2250MG per	Retrograde Amnesia		Zovirax	SS	Glaxosmithkline	
day	6 DAY						
INTRAVENOUS	15MG per day			Polaramin	SS		
	12 DAY			Mopral	SS	Glaxosmithkline	ORAL
20MG Per day	14 DAY			Primperan	SS	Glaxosmithkline	
INTRAVENOUS	60MG per day			Neurontin	SS		ORAL
4 DAY				Fraxodi	C	Glaxosmithkline	
SUBCUTANEOUS				Fungizone	C		
UNKNOWN				Chemotherapy	C		
UNKNOWN							

Date:04/05/05ISR Number: 4627407-1Report Type:Expedited (15-DaCompany Report #2005049905
Age:32 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anticonvulsant Drug Level Increased	Consumer	Neurontin (Gabapentin)	PS		
800 MG (400		Complex Partial Seizures					
MG, 2 IN 1 D)		Disease Recurrence Drug Interaction Hypersensitivity Petit Mal Epilepsy Pharmaceutical Product Complaint		Azelastine Hydrochloride (Azelastine Hydrochloride)	SS		
1000 MG (500				Carbamazepine (Carbamazeine)	SS		
MG, 2 IN 1 D)				Lorazepam(Lorazepam)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/05/05ISR Number: 4627431-9Report Type:Expedited (15-DaCompany Report #2005047915

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		No Adverse Effect	Consumer	Neurontin (Gabapentin)	PS		
	2700 MG (900 MG, 3 IN 1 D)			Bextra (Valdecoxib)	SS		

Date:04/05/05ISR Number: 4627548-9Report Type:Expedited (15-DaCompany Report #2005048449

Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression	Consumer	Neurontin (Gabapentin)	PS		
Other		Completed Suicide Drug Ineffective Self Injurious Behaviour					

Date:04/05/05ISR Number: 4627554-4Report Type:Expedited (15-DaCompany Report #2005048971

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Ineffective Neoplasm Malignant	Consumer	Neurontin (Gabapentin)	PS		ORAL
	300 MG (100 MG, 3 IN 1 D), ORAL	Pain Pharmaceutical Product Complaint		Verapamil (Verapamil) Atenolol (Atenolol) Vitamins (Vitamins) Quinapril Hydrochloride (Quinapril Hydrochloride) Hydrochlorothiazide	C C C C C		

(Hydrochlorothiazide
) C

Date:04/05/05ISR Number: 4627555-6Report Type:Expedited (15-DaCompany Report #2005048898
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cold Sweat Confusional State	Consumer	Neurontin (Gabapentin)	PS		
SEE IMAGE		Drug Ineffective Fear Feeling Abnormal Gallbladder Operation Incision Site Complication Paranoia Reaction To Medical Agent Preservatives		Olopatadine Hydrochloride (Olopatadine Hydrochloride) Lansoprazole (Lansoprazole) Lidocaine (Lidocaine) Carisoprodol (Carisoprodol)			C C C C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/05/05ISR Number: 4627557-XReport Type:Expedited (15-DaCompany Report #2005023528
 Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	5 MG (5 MG, 1 IN 1 D), ORAL	Back Pain	Health Professional	Glucotrol (Glipizide)	PS		ORAL
		Blood Glucose Increased					
		Cough					
	100 MG (100 MG, 3 IN 1 D), ORAL	Muscle Spasms Pain		Neurontin (Gabapentin)	SS		ORAL
		Pharyngitis					
		Treatment Noncompliance					
		Viral Infection		Seretide Mite (Fluticasone Propionate, Salmeterol Xinaifoate)	C		
				Salbutamol (Salbutamol)	C		

Date:04/05/05ISR Number: 4628041-XReport Type:Direct Company Report #CTU 245310
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	300 MG TID 9 DAY	Convulsion	Pharmaceutical Product	Gabapentin 300 Mg (Purepac)	PS	Purepac	
		Complaint		Synthroid	C		
				Methylphenidate	C		
				Fentanyl	C		
				Prevacid	C		
				Klonopin	C		
				Premarin	C		
				Lipitor	C		

Date:04/05/05ISR Number: 4628984-7Report Type:Direct Company Report #CTU 245288
 Age:61 YR Gender:Male I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
100 MG BID		Myositis Pharmaceutical Product		Gabapentin 100 Mg Caps	PS		ORAL
ORAL		Complaint					

Date:04/05/05ISR Number: 4629262-2Report Type:Expedited (15-DaCompany Report #2005049492
 Age:73 YR Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abasia Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL
EVERYDAY, ORAL		Neuropathy Pharmaceutical Product Complaint		Gabapentin (Tablets) (Gabapentin)	SS		ORAL
3600 MG (600 MG, 6 IN 1 D),ORAL				Furosemide (Furosemide)	C		
				Glipizide (Glipizide)	C		
				Fenofibrate (Fenofibrate)	C		
				Lansoprazole			

Freedom Of Information (FOI) Report

(Lansoprazole) C
 Isosorbide
 (Isosorbide) C
 Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C
 Metoprolol Succinate
 (Metoprolol
 Succinate) C
 Trandolapril
 (Trandolapril) C
 Nortriptyline
 (Nortriptyline) C
 Warfarin Sodium
 (Warfarin Sodium) C
 Metformin
 Hydrochloride
 (Metformin
 Hydrochloride) C

Date:04/05/05ISR Number: 4629270-1Report Type:Expedited (15-DaCompany Report #2005049656
 Age:8 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Aggression Self Injurious Behaviour	Consumer	Neurontin (Gabapentin)	PS		
300 MG (2 IN 1 D)		Weight Increased					

Date:04/05/05ISR Number: 4629418-9Report Type:Expedited (15-DaCompany Report #2005052077
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Dependence	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Age:68 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other DF ONCE		Dizziness	Consumer	Diltiazem	PS		
300 MG TID		Feeling Jittery		Gabapentin	SS		
		Heart Rate Increased		Other Unspecified			
DF		Hyperhidrosis		Medications	SS		
		Insomnia		Amitriptyline	C		
		Restlessness		Other Unspecified			
		Vision Blurred		Therapeutic Products	C		

Age:52 YR Gender:Male I/FU:F

Outcome	PT
Other	Abasia
	Back Pain
	Balance Disorder
	Choking

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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
2400 MG (1 D); ORAL		Dyskinesia Dystonia Feeling Abnormal Headache	Consumer	Gabapentin (Gabapentin)	PS		ORAL
ORAL		Mental Impairment Nuclear Magnetic Resonance Imaging Brain Abnormal		Nefazodone Hydrochloride (Nefazodone Hydrochloride)	SS		ORAL
		Opisthotonus Pain Pharyngolaryngeal Pain Road Traffic Accident Speech Disorder Sudden Onset Of Sleep Tremor Vision Blurred		Escitalopram (Escitalopram) Clonazepam All Other Therapeutic Products	SS C C		

Date:04/06/05ISR Number: 4628894-5Report Type:Expedited (15-DaCompany Report #2005043864
Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Abnormal Behaviour Condition Aggravated	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Depression Memory Impairment Mental Disorder Suicidal Ideation Suicide Attempt		All Other Non-Therapeutic Products (All Other Non-Therapeutic Products) Levothyroxine Sodium (Levothyroxine Sodium) Rosuvastatin (Rosuvastatin) Verapamil (Verapamil)	SS C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/06/05ISR Number: 4628905-7Report Type:Expedited (15-DaCompany Report #2004047157
 Age:14 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (200 MG, 3 IN 1 D), ORAL		Emotional Disorder Growth Accelerated Medication Tampering Pharmaceutical Product Complaint Therapeutic Response Decreased	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:04/06/05ISR Number: 4628916-1Report Type:Expedited (15-DaCompany Report #2005044422
 Age:70 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Other		Convulsion Feeling Abnormal Overdose Pain Pharmaceutical Product Complaint	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Irbesartan (Irbesartan)	C		
				Paroxetine Hydrochloride (Paroxetine Hydrochloride)	C		
				Carvedilol (Carvedilol)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Ezetimibe (Ezetimibe)	C		
				Rosiglitazone Maleate (Rosiglitazone Maleate)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Drug Ineffective Hallucination, Auditory	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Head Injury Lower Limb Fracture Nervous System Disorder Skin Laceration Suicidal Ideation		Escitalopram (Escitalopram)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other TAKE 2 CAPSULES BY MOUTH AT BEDTIME	9 DAY	Anxiety Drug Effect Decreased Neuropathic Pain Pharmaceutical Product Complaint		Gabapentin Caps G5027-300mg	PS	Greenstone	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/06/05ISR Number: 4631030-2Report Type:Direct
Age:75 YR Gender:Female I/FU:I

Company Report #CTU 245483

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG BID		Abdominal Pain		Gabapentin- Generic	PS		
		Gamma-Glutamyltransferase Increased		Insulin	C		
		Nausea		Estradiol	C		
		Pharmaceutical Product Complaint		Furosemide	C		
				Plavix	C		
				Atenolol	C		
				Digoxin	C		
				Synthroid	C		
				Lisinopril	C		
				Bromfed	C		

Date:04/06/05ISR Number: 4631274-XReport Type:Expedited (15-DaCompany Report #2005042991
Age:81 YR Gender:Female I/FU:F

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (100 Other MG, 3 IN 1 D), ORAL		Condition Aggravated Fall	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Hypoaesthesia					
		Insomnia					
		Loss Of Consciousness Memory Impairment Pain In Extremity Pharmaceutical Product Complaint Treatment Noncompliance		All Other Therapeutic Products (All Other Therapeutic Products)	C		
				Paracetamol (Paracetamol)	C		

Date:04/07/05ISR Number: 4631235-0Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 245595

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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III QD Pyrexia Neurontin 300mg PS

Urticaria
Vomiting

Date:04/07/05ISR Number: 4631237-4Report Type:Direct Company Report #CTU 245599

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Convulsion		Gabapentin 600mg	PS		ORAL
600MG PO							
Other							

Date:04/07/05ISR Number: 4631396-3Report Type:Expedited (15-DaCompany Report #2005014989

Age:44 YR Gender:Male I/FU:F

Outcome	PT
Disability	Amnesia
	Balance Disorder
	Cervical Spinal Stenosis
	Depression
	Erectile Dysfunction
	Hypertension
	Hypoaesthesia
	Pharmaceutical Product

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Complaint
Spinal Fusion Acquired

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1 D), ORAL		Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Professional				
			Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		ORAL
ORAL			Pantoprazole (Pantoprazole)	C		
			Nabumetone (Nabumetone)	C		
			Potassium Chloride (Potassium Chloride)	C		
			Furosemide (Furosemide)	C		
			Tizanidine Hydrochloride (Tizanidine Hydrochloride)	C		

Date:04/07/05ISR Number: 4631443-9Report Type:Expedited (15-DaCompany Report #2004107063
Age:85 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG (300 MG)		Acute Myocardial Infarction	Health Professional	Neurontin (Gabapentin)	PS		
		Cardiac Failure	Company				
		Congestive Condition Aggravated	Representative	Atorvastatin (Atorvastatin)	C		
				Irbesartan (Irbesartan)	C		
				Glyceryl Trinitrate (Glyceryl Trinitrate)	C		

Furosemide
(Furosemide) C
Insulin (Insulin) C
Glipizide
(Glipizide) C

Date:04/07/05ISR Number: 4631452-XReport Type:Expedited (15-DaCompany Report #2005050566
Age:38 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Drug Hypersensitivity Dystonia Joint Stiffness Swollen Tongue	Consumer	Gabapentin (Gabapentin)	PS		

Date:04/07/05ISR Number: 4631559-7Report Type:Expedited (15-DaCompany Report #2005043001
Age:54 YR Gender:Male I/FU:F

Outcome	PT	Report Source
Hospitalization - Initial or Prolonged	Drug Interaction Vitamin B12 Deficiency	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
600 MG (300 MG, 2 IN 1 D), ORAL			Neurontin (Tablets) (Gabapentin)	PS		ORAL
10 MG, (1 IN 1 D), ORAL			Clonazepam (Clonazepam)	SS		ORAL
			Insulin Glargine (Insulin Glargine)	C		
			Insulin Aspart (Insulin Aspart)	C		
			Ramipril (Ramipril)	C		
			Acetylsalicylate Lysine (Acetylsalicylate Lysine)	C		
			Trimetazidine Hydrochloride (Trimetazidine Hydrochloride)	C		
			Diosmin (Diosmin)	C		

Date:04/07/05ISR Number: 4631640-2Report Type:Expedited (15-DaCompany Report #2005049540

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Hot Flush	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1600 MG (800 MG, 2 IN 1 D), ORAL			Professional				

Date:04/08/05ISR Number: 4632779-8Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 245696

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin (Generic)			
Other		Drug Ineffective		Garbopentin	PS		
400 MG Q 3 H		Pharmaceutical Product					
		Complaint					

Date:04/08/05ISR Number: 4632780-4Report Type:Direct
Age:47 YR Gender:Female I/FU:I

Company Report #CTU 245689

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Gabapentin-Generic			
		Inadequate Analgesia		400 Mg Caps	PS		ORAL
1200 MG PO		Pharmaceutical Product					
		Complaint					
TID							

Date:04/08/05ISR Number: 4632787-7Report Type:Direct
Age:80 YR Gender:Female I/FU:I

Company Report #USP 245691

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin (Generic			
		Drug Ineffective		Form)	PS		
300 MG THREE							
TIMES/DAY							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/08/05ISR Number: 4633284-5Report Type:Expedited (15-DaCompany Report #2005CG00637
 Age:69 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 20 MG QD PO		Abnormal Dreams	Foreign	Mopral	PS		ORAL
Initial or Prolonged INTRAVENOUS	15 MG QD IV	Confusional State	Health	Polaramine	SS		
INTRAVENOUS	60 MG QD IV	Decreased Activity	Professional	Primperan	SS		
INTRAVENOUS	2250 MG QD IV	Disorientation Hallucination, Visual	Other	Zovirax Wellcome	SS	Glaxo Wellcome	
300 MG QD PO		Hypersomnia		Neurontin	SS		ORAL
400 MG QD PO		Incoherent		Neurontin	SS		ORAL
200 MG QD PO		Metabolic Encephalopathy		Neurontin	SS		ORAL
8 MG QD		Retrograde Amnesia		Zophren	SS		
		Somnolence		Fraxodi Fungizone	C C		

Date:04/08/05ISR Number: 4633413-3Report Type:Expedited (15-DaCompany Report #2005051895
 Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600MG 3 TIMES		Condition Aggravated Difficulty In Walking	Consumer	Neurontin (Gabapentin)	PS		ORAL
DAILY AS NECESSARY (600 MG), ORAL		Injury Knee Arthroplasty Pain In Extremity Tendonitis		Temazepam (Temazepam) Ezetimibe (Ezetimibe)	C C		

Date:04/11/05ISR Number: 4633576-XReport Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 245844

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased		Neurontin Generic	PS		ORAL
600MG	ORAL						
		Pharmaceutical Product					
BID		Complaint					

Date:04/11/05ISR Number: 4634946-6Report Type:Expedited (15-DaCompany Report #2005043855
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Amnesia Eye Disorder Feeling Abnormal Hypoaesthesia Pain Visual Disturbance	Consumer	Neurontin (Gabapentin)	PS		

Date:04/11/05ISR Number: 4634951-XReport Type:Expedited (15-DaCompany Report #2005051900
Age:70 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion Vascular Dementia	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG,	ORAL			Aricept (Donepezil)	SS		
10 MG							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/11/05ISR Number: 4634954-5Report Type:Expedited (15-DaCompany Report #2005051911
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blister Coma	Consumer	Neurontin (Gabapentin)	PS		ORAL
200-300MG UP TO THREE TIMES DAILY, ORAL		Drooling Lip Disorder Overdose Speech Disorder		Clonidine (Clonidine) Temazepam (Temazepam)	C C		

Date:04/11/05ISR Number: 4635317-9Report Type:Expedited (15-DaCompany Report #2005049905
 Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Complex Partial Seizures Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		
800 MG (300 MG, 2 IN 1 D)		Potentiation Electroencephalogram Abnormal Petit Mal Epilepsy Pharmaceutical Product Complaint		Azelastine Hydrochloride (Azelastine Hydrochloride) Carbamazepine (Carbamazepine)	SS SS		
1000 MG (500 MG, 2 IN 1 D)				Lorazepam (Lorazepam)	C		

Date:04/11/05ISR Number: 4635384-2Report Type:Expedited (15-DaCompany Report #2005052161
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Pressure Inadequately Controlled	Consumer	Neurontin (Gabepentin)	PS		ORAL
600 MG (300 MG, 2 IN 1D),ORAL		Blood Pressure Increased Condition Aggravated					
5 MG (5 MG, 1 IN 1 D),ORAL		Oedema Peripheral Surgery Therapy Non-Responder		Norvasc (Amlodipine)	SS		ORAL
				Lisinopril (Lisinopril)	C		
				Fursoemide (Furosemide)	C		
				Atenolol (Atenolol)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Amitriptyline (Amitriptyline)	C		
				Esomeprazole (Esomeprazole)	C		
				Simvastatin (Simvastatin)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/11/05ISR Number: 4635394-5Report Type:Expedited (15-DaCompany Report #2005050637

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 800 MG (3 IN Initial or Prolonged 1 D), ORAL Disability Other	Bone Formation Decreased Condition Aggravated Difficulty In Walking Drug Ineffective Fear Foot Operation Migraine Pain Pharmaceutical Product Complaint Road Traffic Accident Sternal Fracture	Consumer	Neurontin (Gabapentin) Quinapril Hydrochloride (Quinapril Hydrochloride)	PS C		ORAL

Date:04/12/05ISR Number: 4633307-3Report Type:Expedited (15-DaCompany Report #2005042430

Age:63 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 400 MG (200 MG, 2 IN 1 D), ORAL	Arthralgia Dizziness Ear Pain Headache Speech Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:04/12/05ISR Number: 4633602-8Report Type:Direct

Company Report #CTU 245952

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG TID PO	Nightmare Pharmaceutical Product Complaint		Neurontin (Generic) 600 Mg	PS		ORAL

Date:04/12/05ISR Number: 4633921-5Report Type:Direct
Age: Gender: I/FU:I

Company Report #CTU 245848

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Muscle Spasms		Gabapentin	PS		ORAL
300 MG PO *							
DAILY		Pharmaceutical Product					
		Complaint		Hctz	C		
				Mysoline	C		
				Atenolol	C		
				K-Dur	C		
				Vit B12	C		

Date:04/12/05ISR Number: 4633979-3Report Type:Direct
Age:42 YR Gender:Male I/FU:I

Company Report #CTU 245901

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Delirium		Gabapentin	PS		
Initial or Prolonged		Depression		Gabitril	SS		
		Mental Status Changes					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/12/05ISR Number: 4634391-3Report Type:Expedited (15-DaCompany Report #2005026789

Age:52 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour Adverse Event	Consumer	Gabapentin (Gabapentin)	PS		ORAL
2400 MG (3 IN		Anxiety					
1 D), ORAL		Back Pain		Nefazodone			
		Balance Disorder		Hydrochloride			
		Choking		(Nefazodone			
		Coordination Abnormal		Hydrochloride)	SS		ORAL
ORAL		Drug Ineffective		Escitalopram			
		Dysphonia		(Escitalopram)	SS		
		Dystonia		Clonazepam			
		Feeling Abnormal		(Clonazepam)	C		
		Gait Disturbance		All Other			
		Headache		Therapeutic Products			
		Memory Impairment		(All Other			
		Neck Pain		Therapeutic			
		Nuclear Magnetic		Products)	C		
		Resonance Imaging Brain					
		Abnormal					
		Opisthotonus					
		Pharyngolaryngeal Pain					
		Road Traffic Accident					
		Somnolence					
		Speech Disorder					
		Thinking Abnormal					
		Tremor					
		Vision Blurred					

Date:04/12/05ISR Number: 4634402-5Report Type:Expedited (15-DaCompany Report #2005040889

Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Disorder	Health	Celebrex (Celecoxib)	PS		ORAL
200 MG (200		Dysstasia	Professional				
MG, 1 IN 1							

D), ORAL	Incontinence			
900 MG (300 MG, 3 IN 1 D), ORAL	Medication Error Muscle Spasms Oedema Peripheral Pharmaceutical Product	Neurontin (Neurontin)	SS	ORAL
300 MG, ORAL	Complaint Treatment Noncompliance Tremor	Gabapentin (Gabapentin)	SS	ORAL
		Esomeprazole (Esomeprazole)	C	
		Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	C	
		Oxycocet (Oxycodone Hydrochloride, Paracetamol)	C	
		Furosemide (Furosemide)	C	
		Clonazepam (Clonazepam)	C	
		Diclofenac Sodium (Diclofenac Sodium)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/12/05ISR Number: 4634418-9Report Type:Expedited (15-DaCompany Report #2005046314

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Abasia	Consumer	Neurontin			
Other		Arthralgia		(Gabapentin)	PS		ORAL
ORAL;		Bedridden					
FOLLOW-UP		Convulsion		Bextra (Valdecoxib)	SS		ORAL
ORAL		Drug Ineffective		Gabapentin			
		Mobility Decreased		(Gabapentin)	SS		
		Weight Increased		Omeprazole			
				(Omeprazole)	C		
				Hyzaar			
				(Hydrochlorothiazide			
				, Losartan			
				Potassium)	C		
				Raloxifene			
				Hydrochloride			
				(Raloxifene			
				Hydrochloride)	C		
				Desipramine			
				(Desipramine)	C		

Date:04/12/05ISR Number: 4634708-XReport Type:Expedited (15-DaCompany Report #2004115149

Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Carotid Artery Stenosis	Foreign	Neurontin			
		Dizziness	Consumer	(Gabapentin)	PS		ORAL
300 MG (300		Dysphagia					
MG, 1 IN 1		Hallucination					
D), ORAL		Headache		Atorvastatin			
		Vertigo		(Atorvastatin)	C		
				Citalopram			
				Hydrobromide			
				(Citalopram			
				Hydrobromide)	C		

Theophylline	
(Theophylline)	C
Diltiazem	
Hydrochloride	
(Diltiazem	
Hydrochloride)	C
Cimicifuga Racemosa	
Root (Cimicifuga	
Racemosa Root)	C
Spirolactone	
(Spirolactone)	C
Fluticasone	
Propionate	
(Fluticasone	
Propionate)	C
Salmeterol Xinafoate	
(Salmeterol	
Xinafoate)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/13/05ISR Number: 4634602-4Report Type:Expedited (15-DaCompany Report #2005040328
 Age:75 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - Initial or Prolonged 1800 MG (600 Other MG, 3 IN 1 D), ORAL	Alpha 2 Globulin Increased Blister Conjunctivitis Drug Interaction	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
(500 MG), ORAL	Erythema Multiforme Prurigo Pruritus Rash Papular		Paracetamol (Paracetamol)	SS		ORAL
500 MCG (2 IN 1 D), ORAL	Stevens-Johnson Syndrome Toxic Skin Eruption		Seretide Mite (Fluticasone Propionate, Salmeterol	C		ORAL

Date:04/13/05ISR Number: 4634673-5Report Type:Direct Company Report #CTU 246065
 Age:47 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300 MG TID	Anxiety Panic Reaction Pharmaceutical Product Complaint		Gabapentin 300 Mg	PS		
Duration 1 MON						

Date:04/13/05ISR Number: 4634680-2Report Type:Expedited (15-DaCompany Report #2005053386
 Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia Dizziness	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL Other		Head Injury Headache	Professional Other	Verapamil (Verapamil)	SS		
ORAL		Loss Of Consciousness Malaise		Paclitaxel (Paclitaxel)	SS		
INTRAVENOUS	INTRAVENOUS	Overdose Vomiting		Cisplatin (Cisplatin)	SS		
INTRAVENOUS	INTRAVENOUS			Aporex (Detropropoxyphene Hydrochloride, Paracetamol)	SS		ORAL
ORAL				Fluindione (Fluindione)	SS		ORAL
ORAL							

Date:04/13/05ISR Number: 4634688-7Report Type:Expedited (15-DaCompany Report #2005052728

Age:53 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Alanine Aminotransferase Abnormal Aspartate Aminotransferase Abnormal Blood Alkaline Phosphatase Abnormal Blood Bilirubin Abnormal

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Freedom Of Information (FOI) Report

Hepatitis Cholestatic

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
1 GRAM (1 D), ORAL		Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Venlafaxine Hydrochloride (Venlafaxine Hydrochloride0 Topiramate (Topiramate) Dihydroergotamine Mesilate (Dihydroergotamine Mesilate) Bromazepam (Bromazepam)	C C C C		

Date:04/13/05ISR Number: 4635504-XReport Type:Expedited (15-DaCompany Report #2005049492
Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other EVERYDAY , ORAL		Abasia Drug Ineffective Nephrectomy	Consumer	Neurontin (Gabapentin)	PS		ORAL
3600 MG (600 MG, 6 IN 1 D), ORAL		Neuropathic Pain Pharmaceutical Product Complaint		Gabapentin (Tablets) (Gabapentin)	SS		ORAL
				Furosemide (Furosemide) Glipizide (Glipizide) Fenofibrate (Fenofibrate)	C C C		

Lansoprazole	
(Lansoprazole)	C
Isosorbide	
(Isosorbide)	C
Acetylsalicylic Acid	
(Acetylsalicylic Acid)	C
Metoprolol Succinate	
(Metoprolol Succinate)	C
Trandolapril	
(Trandolapril)	C
Nortriptyline	
(Nortriptyline)	C
Warfarin Sodium	
(Warfarin Sodium)	C
Metformin	
Hydrochloride	
(Metformin Hydrochloride)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/13/05ISR Number: 4635512-9Report Type:Expedited (15-DaCompany Report #2005055459
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability (300 MG), ORAL		Abasia Asthenia Coordination Abnormal	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
		Diplopia Dysarthria Headache Nystagmus Sensation Of Heaviness		Fentanyl (Fentanyl) Amitriptyline (Amitriptyline) Tramadol (Tramadol) Celecoxib (Celecoxib)	C C C C		

Date:04/13/05ISR Number: 4635557-9Report Type:Expedited (15-DaCompany Report #2004047157
Age:14 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (200 MG , 3 IN 1 D), ORAL		Emotional Disorder Growth Accelerated Pharmaceutical Product Complaint Therapy Non-Responder	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:04/13/05ISR Number: 4635614-7Report Type:Expedited (15-DaCompany Report #KII-2005-0015781
Age:36 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Other ORAL		Acidosis Agitation Bowel Sounds Abnormal Bradycardia Cardiac Arrest	Study Health Professional Other	Morphine Sulfate (Similar To Nd 19-516) (Morphine Sulfate) Unknown Clonazepam(Clonazepa	PS		ORAL

ORAL	Coma	m)	SS	ORAL
ORAL	Disorientation	Baclofen(Baclofen)	SS	ORAL
ORAL	Electrocardiogram Qt Prolonged	Methocarbamol(Methocarbamol)	SS	ORAL
ORAL	Hypertension Intentional Misuse Mydriasis Pupillary Reflex Impaired Somnolence	Gabapentin(Gabapentin)	SS	

Date:04/13/05ISR Number: 4636216-9Report Type:Direct Company Report #CTU 246095
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Chest Pain		Neurontin	PS		ORAL
3000 MGM/D PO		Dyspnoea Myalgia					

Date:04/13/05ISR Number: 4636226-1Report Type:Direct Company Report #CTU 246097
Age:61 YR Gender:Female I/FU:I

Outcome	PT Pain Pharmaceutical Product Complaint
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG BID			Neurontin Generic	PS		

Date:04/13/05ISR Number: 4636228-5Report Type:Direct Company Report #CTU 246099
 Age:80 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other PO 2 Q DAY		Drug Effect Decreased		Neurontin	PS		ORAL
		Mood Altered Pharmaceutical Product Complaint					

Date:04/14/05ISR Number: 4634859-XReport Type:Expedited (15-DaCompany Report #US-KINGPHARMUSA00001-K200500450
 Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Depressed Level Of Consciousness Drug Toxicity Poisoning		Skelaxin	PS	King Pharmaceuticals, Inc.	
				Flurazepam	SS		
				Gabapentin	SS		
				Citalopram	SS		
				Acetaminophen	SS		

Date:04/14/05ISR Number: 4636677-5Report Type:Expedited (15-DaCompany Report #GBR-2005-0001625
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		C-Reactive Protein Increased Inflammation Pruritus	Foreign Health Professional Other	Oxycontin Tablets 10 Mg (Oxycondone Hydrochloride) Cr Tablet	PS		ORAL
20 MG, DAILY,							

ORAL

Movicol (Sodium
Bicarbonate,
Potassium Chloride,
Sodium Chloride,
Macrogol)

SS

ORAL

1 UNIT,

DAILY, ORAL

Keppra
(Levetiracetam)

SS

ORAL

1 GRAM,

DAILY, ORAL

Valproic Acid

SS

ORAL

ORAL

Neurontin
(Gabapentin)

SS

ORAL

ORAL

Praxilene
(Naftidrofuryl
Oxalate)

SS

Mopral (Omeprazole)

SS

ORAL

20 MG, DAILY,

ORAL

Profenid "Rhone
Poulenc"
(Ketoprofen)

SS

ORAL

100 MG,

DAILY, ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/14/05ISR Number: 4636686-6Report Type:Expedited (15-DaCompany Report #8009828

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1000 MG / D Initial or Prolonged PO		C-Reactive Protein Increased	Foreign Health	Keppra	PS		ORAL
1 DF /D PO		Inflammation	Professional	Movicol	SS		ORAL
20 MG /D PO		Rash Morbilliform	Other	Oxycontin	SS		ORAL
PO				Depakine	SS		ORAL
20 MG /D PO				Mopral	SS		ORAL
100 MG /D PO				Profenid	SS		ORAL
PO				Neurontin	SS		ORAL
PO				Praxilene	SS		ORAL

Date:04/14/05ISR Number: 4636711-2Report Type:Expedited (15-DaCompany Report #2005054293

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG (300 MG, 1 D), ORAL		Blood Pressure Diastolic Decreased	Consumer	Gabapentin (Gabapentin)	PS		ORAL
(300 MG), ORAL		Drug Effect Decreased		Gabapentin (Gabapentin)	SS		ORAL
				Valsartan (Valsartan) Amlodipine/Atorvasta tin (Amlodipine,	C		

Atorvastatin) C
 Carvedilol
 (Carvedilol) C
 Acetylsalicylic Acid
 (Acetylsalicylic
 Acid) C

Date:04/14/05ISR Number: 4636771-9Report Type:Expedited (15-DaCompany Report #2005037812
 Age:75 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2400 MG (800 MG, 3 IN 1 D), ORAL	Coordination Abnormal Cystitis Drug Ineffective Feeling Drunk Myoclonus Neuropathy Nystagmus	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Lisinopril (Lisinopril)	C		
			Paroxetine Hydrochloride (Paraxotine Hydrochloride)	C		
			Trazodone (Trazodone)	C		
			Docusate (Docusate)	C		
			Simvastatin (Simvastatin)	C		
			Citalopram Hydrobromide (Citalopram Hydrobromide)	C		
			Tramadol			

Freedom Of Information (FOI) Report

Hydrochloride
(Tramadol
Hydrochloride) C

Date:04/14/05ISR Number: 4637039-7Report Type:Expedited (15-DaCompany Report #2005054277

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
2400 MG (600 MG, 4 IN 1 D)							
ORAL				Clonazepam	C		

Date:04/14/05ISR Number: 4637040-3Report Type:Expedited (15-DaCompany Report #2005054447

Age:61 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other		Myocardial Infarction	Consumer	Neurontin (Gabapentin)	PS		
300 MG (100 MG, 3 IN 1 D)				Metoprolol	C		
				Atorvastatin	C		

Date:04/14/05ISR Number: 4638887-XReport Type:Direct

Company Report #CTU 246174

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion		Neurontin (Generic)	PS		ORAL
600 MG ONCE			Pharmaceutical Product				
PO QHS		Complaint					

Date:04/15/05ISR Number: 4638017-4Report Type:Expedited (15-DaCompany Report #2005055610
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Epilepsy	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
900 MG (1 D),			Professional				
ORAL			Company Representative				

Date:04/15/05ISR Number: 4638121-0Report Type:Expedited (15-DaCompany Report #2005042689
Age:36 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Headache Petit Mal Epilepsy	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
100 MG (100			Professional				
MG, 1 IN 1		Poor Quality Drug					
D), ORAL		Administered					
		Stress		Carbamazepine (Carbamazepine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/15/05ISR Number: 4638125-8Report Type:Expedited (15-DaCompany Report #2005026792

Age:52 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin			
600 MG (300		Exostosis	Health	(Gabapentin)	PS		ORAL
MG, 2 IN 1		Restless Legs Syndrome	Professional				
D), ORAL				Lansoprazole			
				(Lansoprazole)	C		
				Celecoxib			
				(Celecoxib)	C		
				Clonazepam			
				(Clonazepam)	C		
				Ezetimibe			
				(Ezetimibe)	C		

Date:04/15/05ISR Number: 4638315-4Report Type:Expedited (15-DaCompany Report #US014711

Age:56 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Communication Disorder	Foreign	Fentanyl Citrate	PS		
BUCCAL	1600 UG PRN	Disorientation	Study				
BUCCAL		Drug Interaction	Health	Gabapentin	SS		
600 MG DAILY		Overdose	Professional	Oxycontin	SS		
80 MG DAILY			Other	Oxynorm	SS		
30 MG DAILY				Co-Proxamol	SS		
8 TAB/CAP				Paracetamol	C		
DAILY				Dexamethasone-Pix	C		
				Lansoprazole	C		
				Laxoberal			
				"Boehringer			

Ingelheim" C
 Capecitabine C
 Movicol C
 Senna C
 Spironolactone C
 Omeprazole C
 Coproxamol C

Date:04/15/05ISR Number: 4639934-1Report Type:Direct Company Report #CTU 246342
 Age:50 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dry Skin		Gabapentin	PS		
800MG ONCE		Insomnia					
QHS		Pruritus					
		Rash					

Date:04/18/05ISR Number: 4639265-XReport Type:Expedited (15-DaCompany Report #2005056196
 Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased	Consumer	Neurontin			
100 MG		Musculoskeletal Stiffness		(Gabapentin)	PS		ORAL
(100MG, 1 IN1		Self-Medication					
D), ORAL		Spinal Compression					
		Fracture		Valsartan			

Freedom Of Information (FOI) Report

(Valsartan)

C

Date:04/18/05ISR Number: 4639296-XReport Type:Expedited (15-DaCompany Report #USA-2004-0016457

Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Asthenia Blood Glucose Increased Blood Magnesium Decreased Blood Sodium Decreased	Health Professional Other	Oxycontin Tablets (Oxycodone Hydrochloride) Cr Tablet			PS
20 MG, Q12H		Brain Oedema Confusional State Contusion Fall Fatigue		Oxyir Capsules 5 Mg (Oxycodone Hydrochloride, Oxycodone Hydrochloride)			SS
5 MG, Q4H		Injury Mental Status Changes Nausea		Aranesp (Darbepoetin) Kytril (Granisetron)			SS SS
INTRAVENOUS	1 MG,	Pancytopenia					
INTRAVENOUS		Reticulocyte Count		Prograf (Tacrolimus)			SS
SEE IMAGE		Increased Vomiting		Protonix (Pantoprazole)			SS
40 MG, DAILY				Acetylsalicylic Acid (Acetylsalicylic Acid)			SS
81 MG, DAILY				Gabapentin (Gabapentin) Magnesium (Magnesium)			SS SS
400 MG				Donepezil (Donepezil) Inderal (Propranolol Hydrochloride) Dronabinol (Dronabinol) Megace (Megestrol			SS SS SS

Acetate)	SS
Insulin (Insulin)	SS
Aldactone	
(Spironolactone)	SS
Ambien (Zolpidem	
Tartrate)	SS
Senokot (Sennoside	
A+B)	C
Chlorpromazine	
(Chlorpromazine)	C
Glipizide	
(Glipizide)	C
Procardia	C
Pentamidine	
(Pentamidine)	C
Peri-Colace	
(Sennoside A+B,	
Docusate Sodium)	
Tablet	C
Dulcolax	C
Dexamethasone	
(Dexamethasone)	C
Cis-Platinum	
(Cisplatin)	C

Freedom Of Information (FOI) Report

Zoloft (Sertraline
Hydrochloride) C
Leukine
(Sargramostim) C
Adriamycin
(Doxorubicin) C

Date:04/18/05ISR Number: 4639313-7Report Type:Expedited (15-DaCompany Report #2005045701
Age:82 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Dry Mouth	Consumer Health Professional	Detrol La Capsule, Prolonged Release (Tolterodine L-Tartrate)	PS		ORAL
4 MG (4 MG, 1 IN 1 D), ORAL	1	MON					
ORAL	1	MON		Neurontin (Gabapentin)	SS		ORAL
12.5 MG (12.5 MG, 1 IN 1 D), ORAL				Hydrochlorothiazide (Hydrochlorothiazide)	SS		ORAL
ORAL		YR					
ORAL		YR		Ramipril (Ramipril)	SS		ORAL
ORAL		YR		Metoprolol Succinate (Metoprolol Succinate)	SS		ORAL
ORAL				Fenofibrate (Fenofibrate)	SS		ORAL

Date:04/18/05ISR Number: 4639345-9Report Type:Expedited (15-DaCompany Report #2005019428
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Adverse Drug Reaction	Consumer	Lipitor			
Other		Atrial Fibrillation		(Atorvastatin)	PS		ORAL
10 MG (10 MG,		Drug Ineffective					
1 IN 1 D),		Feeling Abnormal					
ORAL		Rash Generalised		Neurontin			
		Therapy Non-Responder		(Gabapentin)	SS		ORAL
600 MG (1 IN		Treatment Noncompliance					
1 D) ORAL		Trigeminal Neuralgia		Atenolol (Atenolol)	SS		ORAL
50 MG (1 IN 1							
D), ORAL				Carbamazepine			
				(Carbamazepine)	SS		ORAL
200 MG (1 IN							
1 D), ORAL				Lisinopril			
				(Lisinopril)	C		
				Hydrochlorothiazide			
				(Hydrochlorothiazide			
)	C		

Date:04/18/05ISR Number: 4639348-4Report Type:Expedited (15-DaCompany Report #2005044442
Age:35 YR Gender:Male I/FU:I

Outcome PT
Other Ageusia
Blood Cholesterol

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D), ORAL

Hallucination

Professional

Headache
Vertigo

Other

Atorvastatin
(Atorvastatin) C
Citalopram
Hydrobromide
(Citalopram
Hydrobromide) C
Theophylline
(Theophylline) C
Diltiazem
Hydrochloride
(Diltiazem
Hydrochloride) C
Cimicifuga Racemosa
Root
(Cimicifuga Racemosa
Root) C
Spironolactone
(Spironolactone) C
Fluticasone
Propionate
(Fluticasone

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Propionate) C
 Salmeterol Xinafoate
 (Salmeterol
 Xinafoate) C

Date:04/18/05ISR Number: 4639899-2Report Type:Expedited (15-DaCompany Report #2005054672
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complex Partial Seizures	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1200 MG (3 IN		Feeling Abnormal	Professional				
1 D), ORAL		Grand Mal Convulsion		Carbamazepine (Carbamazepine)	C		
		Head Injury		Topiramate (Topiramate)	C		
		Loss Of Consciousness					
		Tongue Biting					
		Urinary Incontinence					

Date:04/18/05ISR Number: 4639918-3Report Type:Expedited (15-DaCompany Report #2005056759
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Irritable Bowel Syndrome	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Initial or Prolonged		Serotonin Syndrome	Professional				
(400 MG),			Other				
ORAL							

Date:04/18/05ISR Number: 4640037-0Report Type:Expedited (15-DaCompany Report #2005055287
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anxiety	Consumer	Zoloft (Sertraline)	PS		ORAL
(1 D), ORAL							
Initial or Prolonged		Blood Thyroid Stimulating		Neurontin			

Other (300 MG),	Hormone Decreased	(Gabapentin)	SS	ORAL
ORAL	Carpal Tunnel Syndrome			
	Cartilage Injury	Duloxetine		
	Drug Interaction	Hydrochloride		
	Exercise	(Duloxetine		
30 MG (30 MG,	Electrocardiogram	Hydrochloride)	SS	ORAL
1 IN 1 D),	Abnormal			
ORAL	Gastroesophageal Reflux			
	Disease	Oxygen (Oxygen)	SS	
	Interstitial Lung Disease	All Others		
	Joint Injury	Therapeutic Products		
	Mental Disorder	(All Other		
	Muscle Disorder	Therapeutic		
	Myocardial Infarction	Products)	SS	
	Nightmare	Glyceryl Trinitrate		
	Oxygen Saturation	(Glyceryl		
	Decreased	Trinitrate)	SS	
	Pain			
	Polytraumatism			
	Post Procedural Pain			
	Road Traffic Accident			
	Sleep Apnoea Syndrome			
	Weight Increased			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/19/05ISR Number: 4639127-8Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0552554A
 Age:84 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Decubitus Ulcer		Citrucel Regular			
Other		Diarrhoea		Suspension	PS	Glaxosmithkline	OTHER
9	MON						
		Hyperhidrosis		Gabapentin	SS		OTHER
WK							
		Pneumonia Aspiration		Aricept	C		
		Pyrexia		Premarin	C		
		Vomiting		Acetaminophen	C	Glaxosmithkline	
				Nasocort	C		
				Zaleplon	C		
				Bextra	C		
				Neurontin	C		
YR							

Date:04/19/05ISR Number: 4640409-4Report Type:Expedited (15-DaCompany Report #2005022076
 Age:84 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Back Pain	Health	Lipitor			
Initial or Prolonged		Cellulitis	Professional	(Atorvastatin)	PS		ORAL
20 MG (20							
Disability		Fall					
MG), ORAL							
Other		Feeling Abnormal		Neurontin			
300 MG, ORAL		Foot Fracture		(Gabapentin)	SS		ORAL
		Infarction		Acetylsalicyclic			
		Intervertebral Disc		Acid			
		Protrusion		(Acetylsalicylic			
		Ischaemia		Acid)	C		
		Musculoskeletal Disorder		Alendronate Sodium			
		Neuropathy Peripheral		(Alendronate Sodium)	C		
		Pain In Extremity					
		Spinal Column Stenosis					

Date:04/19/05ISR Number: 4640445-8Report Type:Expedited (15-DaCompany Report #2005043864
 Age:49 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Amnesia Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL Other		Mental Disorder Suicidal Ideation Suicide Attempt		All Other Non-Therapeutic Products (All Other Non-Therapeutic Products)	SS		
UNKNOWN	UNKNOWN			Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Rosuvastatin (Rosuvastatin)	C		
				Verapamil (Verapamil)	C		

Date:04/19/05ISR Number: 4640508-7Report Type:Expedited (15-DaCompany Report #2005055516
Age:68 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged	Blood Pressure Inadequately Controlled
Other	Body Height Decreased

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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
		Burning Sensation Cardiac Failure					
		Coronary Artery Occlusion Depression	Consumer	Glucotrol Xl (Glipizide)	PS		ORAL
ORAL		Diabetes Mellitus		Zoloft (Sertrailne)	SS		ORAL
ORAL		Inadequate Control Disease Recurrence		Gabapentin (Gabapentin)	SS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL		Drug Ineffective Dyspnoea					
		Economic Problem Multiple Fractures Neuropathy Overweight Post Procedural Complication Skin Discolouration Stress Stress Fracture Treatment Noncompliance		Atorvastatin (Atorvastatin) Guaifenesin (Guaifenesin) Dyazide (Hydrochlorothiazide , Triamterene) Salbutamol (Salbutamol) Metoprolol Succinate (Metoprolol Succinate) Valsartan (Valsartan) Human Mixtard (Insulin Human, Insulin Human Injection, Isophane)	C C C C C C C		

Date:04/19/05ISR Number: 4640622-6Report Type:Expedited (15-DaCompany Report #2005058345

Age:63 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1800 MG (1 D) Other 200 MG (1 D)		Abdominal Pain Lower Apathy Asthenia	Foreign Literature Health	Gabapentin (Gabapentin) Tramadol (Tramadol)	PS SS		

Biliary Dilatation	Professional	Enalapril	
Drug Ineffective		(Enalapril)	SS
Drug Interaction		Nebivolol	
Hepatomegaly		(Nebivolol)	SS
Hepatotoxicity		Insulin (Insulin)	SS
Malaise			

Date:04/19/05ISR Number: 4641034-1Report Type:Direct
 Age:51 YR Gender:Male I/FU:I

Company Report #CTU 246569

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Oedema Peripheral		Gabapentin 600 Mg	PS		
600 MG QD							

DIVIDED

Lasix	C
Codeine	C
Glucophage	C
Actos	C
Lescol	C
Lotrin	C
Aggrenox	C
Welchol	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/19/05ISR Number: 4641218-2Report Type:Direct
Age: Gender: I/FU:I

Company Report #CTU 246583

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
PO 800 ONE		Neuropathic Pain		Gabapentin 800mg	PS		ORAL
TID		Pharmaceutical Product					
SWITCHED TO		Complaint					
GENERIC 2		Skin Odour Abnormal					
MONTHS AGO	2 MON	Urine Odour Abnormal					
				Keppra	C		

Date:04/20/05ISR Number: 4641263-7Report Type:Direct
Age: Gender: I/FU:I

Company Report #CTU 246667

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ONE DAILY	2 YR	Drug Ineffective		Gabapentin	PS		

Date:04/20/05ISR Number: 4641265-0Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 246669

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG 3 X		Fatigue Headache		Gabapentin 600 Mg Three Times Daily	PS		ORAL
DAY BY MOUTH		Myalgia					
		Pharmaceutical Product					
		Complaint					

Date:04/20/05ISR Number: 4641266-2Report Type:Direct
Age: Gender: I/FU:I

Company Report #CTU 246671

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Drug Ineffective		Gabapentin	PS		ORAL
600 MG PO QHS	2 MON						

Date:04/20/05ISR Number: 4641969-XReport Type:Expedited (15-DaCompany Report #2005056955

Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Other		Fall Ligament Injury	Consumer	Neurontin (Gabapentin)	PS		

Date:04/20/05ISR Number: 4642453-XReport Type:Direct Company Report #CTU 246652

Age: Gender:Female I/FU:I

Outcome		PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration						
Life-Threatening		Anger		Gabapentin Tablets	PS	Mfd By Greenstone	
1 TAB DAILY		Blister					
AT BEDTIME		Crying		Alprazolam	C		
		Hair Disorder		Hydroxyzine	C		
		Hostility		Lopressor	C		
		Insomnia		Tevetan	C		
		Neuropathic Pain					
		Personality Change					
		Pharmaceutical Product					
		Complaint					
		Rash Pruritic					
		Skin Lesion					
		Suicidal Ideation					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/20/05ISR Number: 4642760-0Report Type:Expedited (15-DaCompany Report #2005052728
 Age:53 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1 GRAM (1 D), ORAL	Hepatitis Cholestatic	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C		
			Topiramate (Topiramate)	C		
			Dihydroergotamine Mesilate (Dihydroergotamine Mesilate)	C		
			Bromazepam (Bromazepam)	C		

Date:04/20/05ISR Number: 4642768-5Report Type:Expedited (15-DaCompany Report #2005057468
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (300 MG, 3 IN 1 D), ORAL	Confusional State Delirium Drug Interaction Feeling Abnormal Paraesthesia	Foreign Health Professional Company Representative	Neurontin (Tablets) (Gabapentin)	PS		ORAL
			Fluoxetine (Fluoxetine)	SS		
			All Other Therapeutic Products(All Other Therapeutic Products)	C		

Date:04/20/05ISR Number: 4643419-6Report Type:Direct Company Report #CTU 246656
Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death 500MG Q 8 Life-Threatening HOURS	Stevens-Johnson Syndrome Toxic Epidermal Necrolysis		Primaxin Neurontin	PS SS		
Hospitalization - 200MG TID Initial or Prolonged Required Intervention to Prevent Permanent Impairment/Damage						

Date:04/21/05ISR Number: 4641760-4Report Type:Expedited (15-DaCompany Report #2005056882
Age:65 YR Gender:Female I/FU:I

Outcome	PT
Other	Blood Pressure Increased Chondropathy Diabetes Mellitus Drug Ineffective Neuropathic Pain Polytraumatism

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Road Traffic Accident

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG (300 MG, 1 IN 1 D), ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL
			Morphine (Morphine) Morphine Sulfate (Morphine Sulfate)	C C		
			Prinzide (Hydrochlorothiazide , Lisinopril)	C		
			Methylphenidate Hydrochloride (Methylphenidate Hydrochloride)	C		
			Metformin Hydrochloride (Metformin Hydrochloride)	C		
			Levothyroxine Sodium(Levothyroxine Sodium)	C		
			Escitalopra (Escitalopram)	C		
			Estrognes C Onjugated (Estrognes Conjugated)	C		

Date:04/21/05ISR Number: 4641979-2Report Type:Expedited (15-DaCompany Report #2004074283

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Balance Disorder Deafness Unilateral Ear Pain Tinnitus	Consumer Health Professional	Neurontin (Gabapentin)	PS		

Date:04/21/05ISR Number: 4642143-3Report Type:Direct
Age:35 YR Gender:Female I/FU:I

Company Report #CTU 246824

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea		Gabapentin	PS		ORAL
2 TAB TID PO		Dizziness Headache Nausea					

Date:04/21/05ISR Number: 4642412-7Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 246771

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Effect Decreased Pharmaceutical Product		Gabapentin Mg	300 PS		ORAL
300 MG ORAL		Complaint Therapeutic Response Decreased					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/21/05ISR Number: 4643044-7Report Type:Expedited (15-DaCompany Report #2005030141

Age:81 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1800 MG (600 MG, 3 IN 1 D), ORAL	Anaphylactic Reaction	Foreign Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
				Carbamazepine (Carbamazepine)	SS		
	0.6 MG (0.2 MG, 3 IN 1 D), ORAL			Buprenorphine (Buprenorphine)	SS		ORAL
				Ibuprofen (Ibuprofen)	C		
				Dyazide (Hydrochlorothiazide , Triamterene)	C		
				Risedronate Sodium (Risedronate Sodium)	C		
				Calcium (Calcium)	C		

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
Other 20 MG, ORAL		Asthma	Consumer	Bextra (Valdecoxib)	PS		ORAL
		Blood Pressure Increased Bronchospasm		Dilantin (Phenytoin Sodium)	SS		ORAL
ORAL		Convulsion Weight Increased		Corticosteroids (Corticosteroids) Bupropion Hydrochloride (Bupropion	SS		

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:04/21/05ISR Number: 4643315-4Report Type:Expedited (15-DaCompany Report #2004074993
Age:55 YR Gender:Male I/FU:F

Outcome	PT
Death	Accidental Overdose
Hospitalization -	Completed Suicide
Initial or Prolonged	Drug Abuser
	Drug Ineffective

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Freedom Of Information (FOI) Report

Dose	Duration	Drug Toxicity Injury Mental Status Changes	Report Source	Product	Role	Manufacturer	Route
		Metabolic Acidosis	Consumer	Neurontin			
		Multiple Drug Overdose		(Gabapentin)	PS		
		Respiratory Distress		Celebrex (Celecoxib)	SS		
		Respiratory Failure		Paroxetine			
		Treatment Noncompliance		Hydrochloride			
				(Paroxetine			
				Hydrochloride)	SS		
				Etrafon-D			
				(Amitriptyline			
				Hydrochloride,			
				Perphenazine)	SS		
				Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	SS		
				Paracetamol			
				(Paracetamol)	SS		
				Esomeprazole			
				(Esomeprazole)	C		
				Baclofen (Baclofen)	C		
				Insulin (Insulin)	C		
				Fentanyl (Fentanyl)	C		
				Metformin			
				Hydrochloride			
				(Metformin			
				Hydrochloride)	C		

Date:04/21/05ISR Number: 4643318-XReport Type:Expedited (15-DaCompany Report #2004058042

Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin			
Other		Completed Suicide		(Gabapentin)	PS		ORAL
300 MG QHS,		Depression					
MAY INCREASE		Drug Ineffective					
TO Q8H OVER		Feeling Abnormal					
ONE), ORAL							

Gun Shot Wound	Valdecoxib	
Hypoaesthesia	(Valdecoxib)	C
Insomnia	Methylprednisolone	
Nervousness	(Methylprednisolone)	C
Pain	Vicodin (Hydrocodone	
Paraesthesia	Bitartrate,	
Polytraumatism	Paracetamol)	C
Stress	Ultracet	
Weight Decreased	(Paracetamol,	
	Tramadol	
	Hydrochloride)	C
	Cyclobenzaprine	
	(Cyclobenzaprine)	C

Date:04/21/05ISR Number: 4643324-5Report Type:Expedited (15-DaCompany Report #2005007423
Age:25 YR Gender:Male I/FU:I

Outcome
Hospitalization -
Initial or Prolonged

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Freedom Of Information (FOI) Report

Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Accident At Work	Consumer	Neurontin			
		Aggression		(Gabapentin)	PS		
		Anxiety		Methadone			
		Arthralgia		Hydrochloride			
		Back Injury		(Methadone			
		Delusion		Hydrochloride)	SS		
		Depression		Clonazepam			
		Drug Abuser		(Clonazepam)	SS		
		Drug Ineffective		Oxycodone			
		Drug Withdrawal Syndrome		Hydrochloride			
		Intentional Misuse		(Oxycodone			
		Joint Sprain		Hydrochloride)	SS		
		Legal Problem					
		Limb Injury					
		Migraine					
		Nephrolithiasis					
		Panic Attack					
		Paranoia					
		Psychotic Disorder					
		Skin Laceration					
		Suicidal Ideation					
		Suicide Attempt					
		Testicular Pain					
		Varicocele					

Date:04/21/05ISR Number: 4643369-5Report Type:Expedited (15-DaCompany Report #2004059901

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acute Respiratory Failure	Consumer	Neurontin			
Hospitalization -		Agitation		(Gabapentin)	PS		
Initial or Prolonged		Anxiety		Xanax Tablet			
Other		Aortic Injury		(Alprazolam)	SS		ORAL
4 MG (1 MG, 4		Completed Suicide					
IN 1 D), ORAL		Haemothorax		Lithium (Lithium)	SS		ORAL
600 MG (300		Lung Injury					
MG, 2 IN 1							

D), ORAL	Overdose			
	Pain	Fluoxetine		
	Pubic Rami Fracture	Hydrochloride		
	Rib Fracture	(Fluoxetine		
	Suicidal Ideation	Hydrochloride)	SS	ORAL
20 MG (20 MG,				
1 IN 1 D),	Suicide Attempt			
ORAL				
		Lorazepam		
		(Lorazepam)	SS	
		Zolpidem Tartrate		
		(Zolpidem Tartrate)	SS	ORAL
10 MG (10 MG,				
1 IN 1 D),				
ORAL				
		Temazepam		
		(Temazepam)	C	
		Diazepam (Diazepam)	C	
		Olanzapine		
		(Olanzapine)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/21/05ISR Number: 4643370-1Report Type:Expedited (15-DaCompany Report #2004058034
 Age:50 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG Disability (TOTAL) Other 2400 MG (600 MG, 4 IN 1 D), ORAL	Affect Lability Anxiety Chest Pain Drug Ineffective Grandiosity Hallucination, Auditory Hyperhidrosis Insomnia Limb Injury Nephritis Interstitial Off Label Use Opiates Positive Overdose Pain Pressure Of Speech Restlessness Speech Disorder Stress Suicidal Ideation Suicide Attempt Tearfulness	Consumer	Neurontin (Gabapentin) Ibuprofen (Ibuprofen) Olanzapine (Olanzapine)	PS SS C		ORAL

Date:04/22/05ISR Number: 4642948-9Report Type:Expedited (15-DaCompany Report #FR-ABBOTT-05P-056-0297333-00
 Age:78 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 2 DAY 8 DAY	C-Reactive Protein Increased Rash Morbilliform Systemic Inflammatory Response Syndrome		Micropakine Granule Profenid Nulytely Levetiracetam Oxycodone	PS SS SS SS		ORAL ORAL ORAL ORAL

9	DAY		Hydrochloride	SS	ORAL
4	DAY		Omeprazole	SS	ORAL
			Gabapentin	SS	ORAL
			Naftidrofuryl Oxalate	SS	ORAL

Date:04/22/05ISR Number: 4643032-0Report Type:Expedited (15-DaCompany Report #2004079234
 Age:83 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Balance Disorder	Consumer	Neurontin			
Other		Delirium		(Gabapentin)	PS		ORAL
1800 MG (600		Dementia					
MG, 3 IN 1		Dizziness					
D), ORAL		Fall					
		Somnolence					

Date:04/22/05ISR Number: 4643068-XReport Type:Expedited (15-DaCompany Report #2005003387
 Age:59 YR Gender:Male I/FU:I

Outcome	PT
Life-Threatening	Corneal Ulcer
Other	Drug Ineffective
	Gun Shot Wound

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Freedom Of Information (FOI) Report

Dose	Duration	Self Injurious Behaviour Spinal Fracture Suicide Attempt	Report Source Consumer	Product Neurontin (Gabapentin)	Role PS	Manufacturer	Route
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Date:04/22/05ISR Number: 4643182-9Report Type:Expedited (15-DaCompany Report #2004071970
Age:39 YR Gender:Female I/FU:F

Outcome Dose Death Other 1800 MG (600 MG, 3 IN 1 D) ORAL	Duration	PT Anxiety Brain Death Brain Herniation Brain Oedema Cerebral Atherosclerosis Cerebral Haemorrhage Completed Suicide Drug Ineffective Facial Bones Fracture Gun Shot Wound Head Injury Injury Intentional Self-Injury Intraventricular Haemorrhage Pain Pneumocephalus Polytraumatism Sinus Bradycardia Skull Fractured Base Subarachnoid Haemorrhage Subdural Haemorrhage Suicidal Ideation	Report Source Consumer	Product Neurontin (Gabapentin) Nortriptyline Esomeprazole Fexofenadine Hydrochloride Amitriptyline Multivitamins W/Minerals (Minerals Nos, Vitamins Nos) Ascorbic Acid	Role PS C C C C C C C C C	Manufacturer	Route ORAL
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Date:04/22/05ISR Number: 4643183-0Report Type:Expedited (15-DaCompany Report #2004070397
Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1200 MG (300 Disability MG, 4 IN 1 D)	Aggression Agitation	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other ORAL		Alcohol Use Deformity					
		Depression Drug Tolerance Overdose		Clonazepam (Clonazepam) (Clonazepam)	SS		ORAL
2 MG (0.5 MG, 4 IN 1 D)		Pain					
ORAL		Polytraumatism					
		Stress At Work Suicide Attempt		Fluoxetine Lithium	C C		

Date:04/22/05ISR Number: 4643192-1Report Type:Expedited (15-DaCompany Report #2004052118
Age:20 YR Gender:Male I/FU:F

Outcome	PT
Death	Acute Respiratory
Other	Distress Syndrome Aggression Anoxic Encephalopathy

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Anxiety					
		Brain Oedema					
		Cardiac Arrest					
		Completed Suicide	Consumer	Neurontin			
		Drug Prescribing Error		(Gabapentin)	PS		
		Glasgow Coma Scale		Venlafaxine			
		Abnormal		Hydrochloride	C		
		Injury Asphyxiation		Risperidone	C		
		Loss Of Consciousness					
		Neck Injury					
		Pain					
		Pneumomediastinum					
		Pneumonia Aspiration					
		Pneumonitis					
		Respiratory Arrest					
		Resuscitation					
		Vomiting					

Date:04/22/05ISR Number: 4643387-7Report Type:Expedited (15-DaCompany Report #2004083654
 Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin			
Other		Completed Suicide		(Gabapentin)	PS		ORAL
ORAL		Depression					
		Injury					
		Pain					
		Suicidal Ideation					

Date:04/22/05ISR Number: 4643389-0Report Type:Expedited (15-DaCompany Report #2004065552
 Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Abdominal Abscess	Consumer	Neurontin			
Hospitalization -		Abdominal Pain		(Gabapentin)	PS		
300 MG (100							
Initial or Prolonged		Candidiasis					
MG, 3 IN 1 D)							
Disability		Coagulopathy		Clonazepam			

Other	Colon Injury	(Clonazepam)	SS
0.5 MG			
	Drug Ineffective	Oxycocet (Oxycodone	
	Emotional Disorder	Hydrochloride,	
	Gun Shot Wound	Paracetamol)	SS
	Haemothorax	Vicodin (Hydrocodone	
	Hypotension	Bitartrate,	
	Major Depression	Paracetamol)	SS
60 MG, (10			
MG, 1 IN 4	Mediastinal Haemorrhage		
HR),	Overdose		
	Pericardial Excision	Pantoprazole	
	Post Procedural Pain	(Pantoprazole)	C
	Postoperative Fever	Latanoprost	
	Somnolence	(Lantanoprost)	C
	Splenic Injury	Esomeprazole	
	Staphylococcal Abscess	(Esomeprazole)	C
	Suicide Attempt	Gemfibrozil	
		(Gemfibrozil)	C
		Pravastatin Sodium	
		(Pravastatin Sodium)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/22/05ISR Number: 4643390-7Report Type:Expedited (15-DaCompany Report #2004077855

Age:14 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Affective Disorder	Consumer	Neurontin			
ORAL		Alcoholism		(Gabapentin)	PS		ORAL
		Anger		Levothyroxine Sodium			
		Depression		(Levothyroxine			
		Drug Abuser		Sodium)	C		
		Hypothyroidism		Clindamycin			
		Learning Disorder		(Clindamycin)	C		
		Psychotic Disorder		Quetiapine Fumarate			
		Schizoaffective Disorder		(Quetiapine			
		Suicide Attempt		Fumarate)	C		
				Lithium (Lithium)	C		
				Multivitamins			
				(Ascorbic Acid,			
				Ergocalciferol,			
				Folic Acid,			
				Nicotinamide,	C		
				Paroxetine			
				Hydrochloride			
				(Paroxetine			
				Hydrochloride)	C		
				Clozapine			
				(Clozapine)	C		
				Lorazepam			
				(Lorazepam)	C		
				Docusate Sodium			
				(Docusate Sodium)	C		

Date:04/22/05ISR Number: 4643391-9Report Type:Expedited (15-DaCompany Report #2004106658

Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Consumer	Neurontin			
		Depression		(Gabapentin)	PS		
		Drug Ineffective					
		Panic Attack					

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin			
Hospitalization - 600 MG (300 Initial or Prolonged MG, 2 IN 1 Other D), ORAL		Cognitive Disorder		(Gabapentin)	PS		ORAL
		Completed Suicide					
		Depressed Level Of Consciousness Depression		Midazolam (Midazolam)	SS		
INTRAVENOUS	INTRAVENOUS	Fatigue		Tramadol			
		Intentional Misuse		Hydrochloride			
		Libido Decreased		(Tramadol			
ORAL		Major Depression		Hydrochloride)	SS		ORAL
		Respiratory Distress		Citalopram			
		Sleep Disorder		(Citalopram)	SS		
		Sluggishness		Pancuronium			
		Somnolence		(Pancuronium)	SS		
INTRAVENOUS	INTRAVENOUS	Suicidal Ideation		All Other Therapeutic Products			

Freedom Of Information (FOI) Report

(All Other
Therapeutic
Products) SS
Zolpidem Tartrate
(Zolpidem Tartrate) C
Ibuprofen
(Ibuprofen) C
Clonazepam
(Clonazepam) C
Mirtazapine
(Mirtazapine) C

Date:04/22/05ISR Number: 4644318-6Report Type:Expedited (15-DaCompany Report #2005059333
Age:85 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Cerebral Disorder	Foreign	Neurontin			
Other		Drug Withdrawal	Health	(Gabapentin)	PS		ORAL
		Convulsions	Professional	Oxazepam (Oxazepam)	SS		ORAL
75 MG (1 IN 1		Leukoaraiosis		Sulpiride			
				(Sulpiride)	C		
				Bromazepam			
				(Bromazepam)	C		
				Amlodipine Besilate			
				(Amlodipine			
				Besilate)	C		
				Buflomedil			
				(Buflomedil)	C		
				Betahistine			
				(Betahistine)	C		
				Alendronate Sodium			
				(Alendronate Sodium)	C		
				Omeprazole			
				(Omeprazole)	C		
				Mianserin			
				Hyrdochloride			
				(Mianserin			
				Hydrochloride)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged (600 MG, Other UNKNOWN), ORAL	Anger Incorrect Dose Administered Influenza Irritable Bowel Syndrome Jaw Disorder Mood Altered Muscle Spasms Nausea Neuralgia Pain Pain In Extremity Pharmaceutical Product Complaint Spinal Operation Trigeminal Neuralgia	Consumer Health Professional	Neurontin (Tablets) (Gabapentin) Oxycodone Hydrochloride (Oxycodone Hydrochloride) Lansoprazole (Lansoprazole) Librax (Chlordiazepoxide Hydrochloride, Clidinium Bromide) Provella-14 (Estrogens	PS C C C		ORAL

Freedom Of Information (FOI) Report

Conjugated,
 Medroxyprogesterone
 Acetate) C
 Zolpidem Tartrate
 (Zolpidem Tartrate) C
 Prochlorperazine
 Edisylate
 (Prochlorperazine
 Edisylate) C
 Ondansetron
 Hydrochloride
 (Ondansetron
 Hydrochloride) C
 Oxcarbazepine
 (Oxcarbazepine) C
 Vicodine
 (Hydrocodone
 Bitartrate,
 Paracetamol) C

Date:04/22/05ISR Number: 4645147-XReport Type:Expedited (15-DaCompany Report #2005058414
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	2.4 GRAM (1.2 GRAM, 2 IN 1	Drug Ineffective Insomnia Stent Placement	Foreign Consumer	Neurontin (Gabapentin)	PS		

D),

Date:04/22/05ISR Number: 4645920-8Report Type:Direct Company Report #CTU 246885
 Age:56 YR Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 800 MG QID		Dry Mouth Dysphonia Pharyngolaryngeal Pain Therapeutic Response Unexpected With Drug		Neurontin (Generic)	PS		

Substitution

Date:04/22/05ISR Number: 4645926-9Report Type:Direct Company Report #CTU 246890
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Therapeutic Response		Neurontin	PS		
100 MG		Unexpected With Drug Substitution Unevaluable Event					

Date:04/25/05ISR Number: 4644531-8Report Type:Direct Company Report #CTU 247026
 Age:9 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Erythema		Neurontin	PS		ORAL
Other		Pruritus					
100 MG 2 CAPS							
PO				Motrin	SS		ORAL
400 MG PO							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/25/05ISR Number: 4644688-9Report Type:Direct
Age:46 YR Gender:Female I/FU:I

Company Report #CTU 247114

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG	TID	Apathy		Neurontin 600 Mg	PS		
		Cognitive Disorder Delusion Drug Ineffective Drug Prescribing Error Hallucination Impaired Work Ability Intelligence Test Abnormal Personality Change Psychotic Disorder Suicidal Ideation					

Date:04/25/05ISR Number: 4645369-8Report Type:Expedited (15-DaCompany Report #2005058410
Age:53 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (900 MG)		Neutropenic Sepsis White Blood Cell Count Decreased	Foreign Health Professional	Gabapentin (Gabapentin)	PS		
				Morphine (Morphine) Lansoprazole (Lansoprazole) Phenoxyethylpenicil lin (Phenoxyethylpenici llin)	C C C		

Date:04/25/05ISR Number: 4645371-6Report Type:Expedited (15-DaCompany Report #2005057945
Age:73 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Confusional State	Foreign	Gabapentin			

900 MG (300	Dizziness	Health	(Gabapentin)	PS	ORAL
MG, 3 IN 1	Dry Mouth	Professional			
D), ORAL	Scotoma				
300 MG (150			Tramadol (Tramadol)	SS	ORAL
MG, 2 IN 1					
D), ORAL					
			Alendronate Sodium		
			(Alendronate Sodium)	C	
			Calcium		
			Carbonate/Colecalciferol (Calcium		
			Carbonate,		
			Colecalciferol)	C	
			Drug Used In		
			Diabetes (Drug Used		
			In Diabetes)	C	
			Tramadol (Tramadol)	C	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/25/05ISR Number: 4645392-3Report Type:Expedited (15-DaCompany Report #2005059888
Age:59 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening Hospitalization - (3 IN 1 D), Initial or Prolonged ORAL Disability	Circulatory Collapse Dizziness Hypoaesthesia Loss Of Consciousness Orthostatic Hypotension	Foreign Health Professional	Gabapentin (Gabapentin) Amitriptyline (Amitriptyline)	PS C		ORAL

Date:04/26/05ISR Number: 4644795-0Report Type:Expedited (15-DaCompany Report #US-BOEHRINGER INGELHEIM PHARMACEUTICALS, INC.-2005-
Age: 10BP Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged TPV/r 1000/400 mg	Atypical Mycobacterial Infection Ecthyma Hallucination Lymphadenopathy Mental Status Changes Oedema Peripheral Somnolence Tremor		Tipranavir / Ritonavir Capsules Neurontin Ms Contin Morphine Ir Vicodin T-20 Truvada Diflucan Asocal Imodium Dto Dapsone Florinef Celexa Klonopin Valcyte Procrit Tylenol	PS SS SS SS C C C C C C C C C C C C C C C C	B.I. Pharmaceuticals, Inc. /Ridgefield Roxane Laboratories, Inc.	ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL ORAL

SUBCUTANEOUS

1 tab

2 tabs	Lomotil	C	ORAL
	Rantidine	C	
	Flagyl	C	ORAL
	Trazadone	C	ORAL
	Calcium	C	ORAL
600/200			
	Lovenox	C	
SUBCUTANEOUS			
	Colace	C	ORAL
	Neupogen	C	
SUBCUTANEOUS			
	Fluronazole	C	ORAL
	Lactaid	C	ORAL
	Valganciclovir	C	ORAL
	Folate	C	ORAL
	Azithromycin	C	ORAL
	Nexium	C	ORAL

Date:04/26/05ISR Number: 4647213-1Report Type:Expedited (15-DaCompany Report #2005054672
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Complex Partial Seizures Grand Mal Convulsion	Foreign Study	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1200 MG (3 IN							
1 D), ORAL		Head Injury	Health				
		Malaise	Professional	Carbamazepine			

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(Carbamazepine) C
 Topiramate
 (Topiramate) C

Date:04/26/05ISR Number: 4647219-2Report Type:Expedited (15-DaCompany Report #2005059757
 Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Foreign	Neurontin			
		Drug Exposure During	Study	(Gabapentin)	PS		ORAL
ORAL		Pregnancy	Health	Folic Acid (Folic			
		Haemorrhagic Disorder	Professional	Acid)	C		
		Pregnancy On Oral					
		Contraceptive					

Date:04/26/05ISR Number: 4647250-7Report Type:Expedited (15-DaCompany Report #2005058799
 Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebral Disorder	Consumer	Neurontin			
		Cold Sweat		(Gabapentin)	PS		ORAL
200 MG (100		Crying					
MG 2 IN 1 D)		Depression					
ORAL		Dizziness		Venlafaxine			
		Feeling Abnormal		Hydrochloride			
		Homicidal Ideation		(Venlafaxine			
		Hostility		Hydrochloride)	C		
		Syncope		Atenolol (Atenolol)	C		
		Treatment Noncompliance		Folic Acid (Folic			
				Acid)	C		
				Alprazolam			
				(Alprazolam)	C		
				Potassium			
				(Potassium)	C		
				Amlodipine Besilate			
				(Amlodipine			
				Besilate)	C		
				Percogesic			

(Paracetamol, Phenyltoloxamine Citrate)	C
Oxycocet (Oxycodone Hydrochloride, Paracetamol)	C
Hydrocodone (Hydrocodone)	C
Potassium (Potassium)	C
Lisinopril (Lisinopril)	C
Propacet (Dextropropoxyphene Napsilate, Paracetamol)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/26/05ISR Number: 4647255-6Report Type:Expedited (15-DaCompany Report #2005058875
 Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Cholesterol Increased	Consumer	Gabapentin (Gabapentin)	PS		ORAL
4500 MG (1500 MG 3 IN 1 D) ORAL		Blood Thyroid Stimulating Hormone Increased					
		Blood Triglycerides Increased		Carbamazepine (Carbamazepine)	C		
		Drug Ineffective		Atorvastatin (Atorvastatin)	C		
		Hysterectomy		Levothyroxine (Levothyroxine)	C		
		Low Density Lipoprotein Increased		Sertraline Hydrochloride (Sertraline Hydrochloride)	C		
		Therapeutic Response Unexpected With Drug Substitution		Insulin (Insulin) Simvastatin (Simvastatin)	C		

Date:04/26/05ISR Number: 4647290-8Report Type:Expedited (15-DaCompany Report #2005059041
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Ineffective Pain In Extremity	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (100 MG, 3 IN 1 D), ORAL		Thrombosis					
				All Other Therapeutic Products (All Other Therapeutic Products)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated	Consumer	Neurontin			
		Drug Ineffective		(Gabapentin)	PS		ORAL
ORAL		Pharmaceutical Product		Gabapentin (Tablets)			
		Complaint		(Gabapentin)	SS		ORAL
3600 MG (600		Renal Disorder					
MG, 6 IN 1							
D), ORAL							
				Furosemide			
				(Furosemide)	C		
				Glipizide			
				(Glipizide)	C		
				Fenofibrate			
				(Fenofibrate)	C		
				Lansoprazole			
				(Lansoprazole)	C		
				Isosorbide			
				(Isosorbide)	C		
				Acetylsalicylic Acid			
				(Acetylsalicylic			
				Acid)	C		
				Metoprolol Succinate			

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Metoprolol Succinate)	C
Trandolapril (Trandolapril)	C
Nortriptyline (Nortriptyline)	C
Warfarin Sodium (Warfarin Sodium)	C
Metformin Hydrochloride (Metformin Hydrochloride)	C

Date:04/26/05ISR Number: 4647452-XReport Type:Expedited (15-DaCompany Report #2005059645
 Age:58 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin			
Other		Completed Suicide		(Gabapentin)	PS		
		Drug Ineffective					
		Suicidal Ideation					

Date:04/26/05ISR Number: 4647806-1Report Type:Expedited (15-DaCompany Report #05P-056-0297333-00
 Age:78 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Rash Morbilliform Systemic Inflammatory Response Syndrome	Foreign Health Professional	Micropakine Granule (Sodium Valproate)	PS		ORAL
				Profenid (Ketoprofen)			
				(Ketoprofen)			
				(Ketoprofen)	SS		ORAL
100 MG, 1 IN							
1 D ORAL							
				Nulytely	SS		ORAL
13.125 GM, 1							
IN 1 D ORAL							

1 GM, 1 IN 1	Levetiracetam	SS	ORAL
D ORAL			
20 MG, 1 IN 1	Oxycodone Hydrochloride	SS	ORAL
D, ORAL			
20 MG, 1 IN 1	Omeprazole	SS	ORAL
D, ORAL			
ORAL	Gabapentin	SS	ORAL
	Naftidrofuryl Oxalate	SS	ORAL
ORAL			

Date:04/26/05ISR Number: 4647856-5Report Type:Expedited (15-DaCompany Report #2005AP02339
Age:66 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Atrial Fibrillation	Health	Atenolol	PS		ORAL
50 MG DAILY		Drug Ineffective	Professional				
PO		Malaise		Atenolol	SS		ORAL
50 MG DAILY		Rash Generalised					
PO		Trigeminal Neuralgia		Atenolol	SS		ORAL
50 MG DAILY							
PO				Lipitor	SS		ORAL
10 MG DAILY							
PO							

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20 MG DAILY	Lipitor	SS	ORAL
PO			
600 MG DAILY	Neurontin	SS	ORAL
PO			
1800 MG DAILY	Neurontin	SS	ORAL
PO			
200 MG DAILY	Tegretol	SS	ORAL
PO			

Date:04/26/05ISR Number: 4648286-2Report Type:Expedited (15-DaCompany Report #2005037725

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Body Temperature Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL; 300 MG,		Brain Operation					
ORAL		Condition Aggravated Convulsion Epileptic Aura Grand Mal Convulsion		Valproate Semisodium (Valproate Semisodium)	C		
				Risperidone (Risperidone)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Guanfacine Hydrochloride (Guanfacine Hydrochloride)	C		
				Benztropeine (Benztropeine)	C		
				Quetiapine Fumarate (Quetiapine Fumarate)	C		
				Diazepam (Diazepam)	C		

Date:04/26/05ISR Number: 4648328-4Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 247123

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Therapeutic Response Unexpected With Drug Substitution		Gabapentin	PS		

Date:04/27/05ISR Number: 4647717-1Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 247223

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Convulsion Drug Effect Decreased Therapeutic Response Unexpected With Drug Substitution		Gabapentin	PS		
Other							

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/27/05ISR Number: 4648473-3Report Type:Expedited (15-DaCompany Report #2005063015

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dysarthria	Foreign	Neurontin			
ORAL		Tension	Health	(Gabapentin)	PS		ORAL
			Professional				
			Company				
			Representative				

Date:04/27/05ISR Number: 4648536-2Report Type:Expedited (15-DaCompany Report #2005060987

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Alopecia	Foreign	Neurontin			
Initial or Prolonged		Dysphemia	Health	(Gabpentin)	PS		ORAL
ORAL							
		Somatoform Disorder	Professional	Valproate Sodium			
		Tremor	Company	(Valproate Sodium)	SS		
			Representative	Sertraline			
				(Sertraline)	C		
				Panadeine Co (Codine			
				Phosphate,			
				Paracetamol)	C		
				Tramadol (Tramadol)	C		

Date:04/27/05ISR Number: 4648645-8Report Type:Expedited (15-DaCompany Report #PAR_0005_2005

Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Asthenia	Foreign	Isoptine	PS		
Initial or Prolonged		Head Injury	Health	Paclitaxel	SS		
		Loss Of Consciousness	Professional	Cisplatine	SS		
		Malaise	Other	Fluidione	SS		
		Metastases To Central		Di-Gesic	SS		
		Nervous System		Gabapentin	SS		
		Open Wound					
		Overdose					
		Vomiting					

Date:04/27/05ISR Number: 4648938-4Report Type:Expedited (15-DaCompany Report #2005038428

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Neurontin			
		Haemorrhage		(Gabapentin)	PS		
				Norvasc (Amlodipine)	SS		

Date:04/27/05ISR Number: 4649475-3Report Type:Expedited (15-DaCompany Report #2004106809

Age:24 YR Gender:Male I/FU:F

Outcome	PT
Death	Apnoea
	Completed Suicide
	Euphoric Mood
	Foaming At Mouth
	Loss Of Consciousness
	Mental Disorder
	Mouth Haemorrhage

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Freedom Of Information (FOI) Report

Multiple Drug Overdose
Polytraumatism

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
(3 IN 1 D)		Consumer	Neurontin (Gabapentin)	PS		
			Morphine (Morphine)	SS		
			Fentanyl (Fentanyl)	SS		
TRANSDERMAL	175 MG (1 IN					
1 D),						
TRANSDERMAL						

Date:04/27/05ISR Number: 4649562-XReport Type:Expedited (15-DaCompany Report #2005042730
Age:42 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Agitation	Consumer	Neurontin (Gabapentin)	PS		
Hospitalization -		Anorexia					
900 MG (300							
Initial or Prolonged		Atelectasis					
MG, 3 IN 1 D)							
Disability		Blindness		Paroxetine Hydrochloride			
Other		Cerebral Haemorrhage		(Paroxetine Hydrochloride)	C		
		Cognitive Disorder		Chlorpromazine Hydrochloride			
		Contusion		(Chlorpromazine Hydrochloride)	C		
		Delirium		Zolpidem Tartrate			
		Diabetes Insipidus		(Zolpidem Tartrate)	C		
		Dyspepsia					
		Dysphagia					
		Eyeball Rupture					
		Facial Bones Fracture					
		Fracture					
		Gun Shot Wound					
		Hypocalcaemia					
		Hypomagnesaemia					
		Intentional Self-Injury					
		Loss Of Consciousness					
		Pleural Effusion					
		Pneumocephalus					
		Pneumonia Bacterial					
		Pseudomonas Infection					

Respiratory Failure
Sinus Disorder
Soft Tissue Injury
Suicide Attempt

Date:04/28/05ISR Number: 4649372-3Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 247315

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Required 300MG PO		Dyspepsia		Gabapentin	PS		ORAL
Intervention to TID		Gastrooesophageal Reflux Disease					
Prevent Permanent Impairment/Damage		Pharmaceutical Product Complaint					

Date:04/28/05ISR Number: 4649443-1Report Type:Direct
Age:64 YR Gender:Female I/FU:I

Company Report #CTU 247356

Outcome	PT
Other	Drug Effect Decreased 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
600MG (3 TIMES DAILY)			Neurontin 600mg	PS		

Date:04/28/05ISR Number: 4650245-0Report Type:Expedited (15-DaCompany Report #2005049653
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 100 MG (100 Initial or Prolonged MG, 1 IN 1 Other D), UNKNOWN		Pulmonary Embolism	Consumer Health Professional	Neurontin (Gabapentin)	PS		
				All Other Therapeutic Products (All Other Therapeutic Products)	C		

Date:04/28/05ISR Number: 4650271-1Report Type:Expedited (15-DaCompany Report #KII-2005-0016077
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other DAILY, ORAL		Agitation Anxiety Blood Ph Decreased Blood Ph Increased	Study Health Professional Other	Oxycodone Hydrochloride (Oxycodone Hydrochloride)	PS		ORAL
QID, QID, ORAL		Blood Potassium Increased Coma Drooling Electrocardiogram St		Morphine Sulfate (Morphine Sulfate)	SS		
				Carisoprodol (Carisoprodol)	SS		ORAL

TID, ORAL	Segment Elevation Hyperhidrosis	Gabapentin (Gabapentin)	SS	ORAL
ORAL	Oxygen Saturation	Antidepressants ()	SS	ORAL
ORAL	Decreased Tremor	Oral Antidiabetics ()	SS	ORAL
		Triamterene (Triamterene)	SS	
		Proton Pump Inhibitor ()	SS	
		Thyroid Preparations ()	SS	
		Serumlipidreducing Agents ()	SS	
		Estradiol (Estradiol)	SS	
		Amfetamine (Amfetamine)	SS	
		Barbiturates ()	SS	

Date:04/28/05ISR Number: 4650314-5Report Type:Expedited (15-DaCompany Report #2005051911
Age:52 YR Gender:Male I/FU:F

Outcome PT
Other Aphasia
Blister
Depressed Level Of

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Consciousness Drooling Overdose	Report Source	Product	Role	Manufacturer	Route
200-300MG UP TO THREE TIMES DAILY, ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Clonidine (Clonidine)	C		
				Temazepam (Temazepam)	C		

Date:04/28/05ISR Number: 4650653-8Report Type:Direct
Age: Gender:Female I/FU:I Company Report #CTU 247383

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Ineffective Therapeutic Response Unexpected With Drug Substitution		Gabapentin 600mg	PS		

Date:04/29/05ISR Number: 4651011-2Report Type:Expedited (15-DaCompany Report #2005040445
Age:20 YR Gender:Male I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
4800 MG (800 MG, 6 IN 1 D), ORAL		Disease Recurrence Petit Mal Epilepsy	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
		Poor Quality Drug Administered	Professional	Insulin (Insulin) Lamotrigine (Lamotrigine)	C C		

Paxil (Paroxetine
Hydrochloride) C
Clonazepam
(Clonazepam) C

Date:04/29/05ISR Number: 4651022-7Report Type:Expedited (15-DaCompany Report #2005051895
Age:79 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Lower Limb Fracture	Consumer	Neurontin (Gabapentin)	PS		ORAL
600MG 3 TIMES		Tendonitis					
DAILY AS		Treatment Noncompliance					
NECESSARY							
(600 MG),							
ORAL				Gabapentin (Gabapentin)	SS		
				Temazepam (Temazepam)	C		
				Ezetimbe (Ezetimbe)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:04/29/05ISR Number: 4651041-0Report Type:Expedited (15-DaCompany Report #2005061466

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	10 MG (10 MG,	Arthralgia	Consumer	Bextra (Valdecoxib)	PS		ORAL
1 IN 1 D),		Cough					
ORAL		Gingival Atrophy					
		Gingival Discolouration		Neurontin			
900 MG (300		Gingival Oedema		(Gabapentin)	SS		ORAL
MG, 3 IN 1		Gingival Pain					
D), ORAL		Gingival Recession					
		Gingivitis		Atenolol (Atenolol)	C		
		Impaired Healing		Lisinopril			
		Impaired Work Ability		(Lisinopril)	C		
		Nausea		Potassium			
		Oral Mucosal Exfoliation		(Potassium)	C		
		Post Procedural		Insulin Glargine			
		Complication		(Insulin Glargine)	C		
		Skin Disorder		Insulin Lispro			
		Therapy Non-Responder		(Insulin Lispro)	C		
				Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	C		
				Vicodin (Hydrocodone			
				Bitartrate,			
				Paracetamol)	C		
				Clonazepam			
				(Clonazepam)	C		
				Temazepam			
				(Temazepam)	C		
				Clopidogrel Sulfate			
				(Clopidogrel			
				Sulfate)	C		
				Lansoprazole			
				(Lansoprazole)	C		
				Pancrelipase			
				(Pancrelipase)	C		
				Paroxetine			
				Hydrochloride			

(Paroxetine
Hydrochloride) C
All Other
Therapeutic Products
(All Other
Therapeutic
Products) C
Carbidopa
(Carbidopa) C

Date:04/29/05ISR Number: 4651075-6Report Type:Expedited (15-DaCompany Report #2005063311

Age: Gender:Female I/FU:I

Outcome PT
Other Asthenia
Condition Aggravated
Drug Effect Decreased
Drug Ineffective
Fatigue
Feeling Abnormal
Pharmaceutical Product

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Complaint Sleep Apnoea Syndrome Somnolence Weight Decreased	Report Source	Product	Role	Manufacturer	Route
900 MG (900 MG, 1 IN 1 D), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
900 MG (900 MG, 1 IN 1 D), ORAL				Gabapentin (Gabapentin)	SS		ORAL
				Glipizide (Glipizide)	C		
				Metformin Hydrochloride (Metformin Hydrochloride)	C		
				Levothyroxine Sodium (Levothyroxine Sodium)	C		
				Medroxyprogesterone Acetate (Medroxyprogesterone Acetate)	C		
				Estradiol (Estradiol)	C		
				Folic Acid (Folic Acid)	C		
				Methotrexate (Methotrexate)	C		
				Multivitamins (Ascorbic Acid, Ergocalciferol, Folic Acid, Nicotinamide,	C		
				Latanoprost (Latanoprost)	C		
				Alprazolam (Alprazolam)	C		
				Lomoitil (Atropine			

Sulfate, Diphenoxylate Hydrochloride)	C
Galenic /Paracetamol/Codeine / (Codeine, Paracetamol)	C
Aporex (Dextropropoxyphene Hydrochloride, Paracetamol)	C
Respaire-Sr-120 (Guaifenesin, Pseudosphedrine Hydrochloride)	C
Vicodin (Hydrochloride Bitartrate, Paracematol)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/02/05ISR Number: 4651682-0Report Type:Expedited (15-DaCompany Report #2005061466

Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cough	Consumer	Bextra (Valdecoxib)	PS		ORAL
10 MG (10 MG,		Gingival Disorder					
1 IN 1 D),		Gingival Pain					
ORAL		Gingival Swelling		Neurontin			
		Gingivitis		(Gabapentin)	SS		ORAL
900 MG (300		Impaired Healing					
MG, 3 IN 1		Nausea					
D), ORAL		Oral Mucosal Exfoliation		Atenolol (Atenolol)	C		
				Lisinopril			
				(Lisinopril)	C		
				Potassium			
				(Potassium)	C		
				Insulin Glargine			
				(Insulin Glargine)	C		
				Insulin Lispro			
				(Insulin Lispro)	C		
				Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	C		
				Vicodin (Hydrocodone			
				Bitartrate,			
				Paracetamol)	C		
				Clonazepam			
				(Clonazepam)	C		
				Temazepam			
				(Temazepam)	C		
				Clopidogrel Sulfate			
				(Clopidogrel			
				Sulfate)	C		
				Lansoprazole			
				(Lansoprazole)	C		
				Pancrelipase			
				(Pancrelipase)	C		
				Paroxetine			
				Hydrochloride			

(Paroxetine
 Hydrochloride) C
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C
 Carbidopa
 (Carbidopa) C

Date:05/02/05ISR Number: 4652246-5Report Type:Expedited (15-DaCompany Report #2005063453
 Age:80 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Thrombophlebitis	Health Professional	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 MG, 3 IN 1 D), ORAL				Prednisone Tablet (Prednisone) Valaciclovir	SS		

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Freedom Of Information (FOI) Report

Hydrochloride
(Valaciclovir
Hydrochloride) C

Date:05/02/05ISR Number: 4652681-5Report Type:Expedited (15-DaCompany Report #2005063347

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Feeling Abnormal Movement Disorder	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL			Professional Company Representative	Sertraline Hydrochloride (Sertraline Hydrochloride) All Other Therapeutic Products (All Other Therapeutic Products)	C C		

Date:05/02/05ISR Number: 4652729-8Report Type:Expedited (15-DaCompany Report #2005-BP-01716PF

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 2 IN 1 D		Anxiety	Consumer	Pramipexole	PS		
Initial or Prolonged Required		Asthma Balance Disorder	Health Professional	Neurontin (Gabapentin)	SS		ORAL
Intervention to Prevent Permanent 2 IN 1 D		Chronic Obstructive Pulmonary Disease		Zyrtec-D 12 Hour (Cirrus)	SS		
Impairment/Damage 2 IN 1 D		Condition Aggravated Dental Operation		Clonazepam (Clonazepam)	SS		
		Disturbance In Attention Dizziness		Mirtazapine (Mirtazapine)	SS		
		Memory Impairment Multiple Allergies		Amitriptyline (Amitriptyline)	SS		
		Sinus Disorder		Montelukast Sodium (Montelukast Sodium)	C		

Combivent
(Combivent) C
Mirtazapine
(Mirtazapine) C
Esomeprazole
(Esomeprazole) C
Salbutamol
(Salbutamol) C
Calamine/Camphor/Dip
henhydramine C
Bupropion
(Bupropion) C

Date:05/03/05ISR Number: 4652969-8Report Type:Expedited (15-DaCompany Report #2005055516
Age:68 YR Gender:Female I/FU:F

Outcome PT
Hospitalization - Blood Pressure
Initial or Prolonged Inadequately Controlled
Other Blue Toe Syndrome
Burning Sensation
Cardiac Failure

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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
		Coronary Artery Occlusion Depression					
		Diabetes Mellitus Inadequate Control	Consumer	Glucotrol Xl (Glipizide)	PS		ORAL
ORAL							
		Difficulty In Walking		Zoloft (Sertraline)	SS		ORAL
ORAL							
		Drug Effect Decreased Drug Ineffective		Gabepentin (Gabapentin)	SS		ORAL
900 MG (300							
MG, 3 IN 1		Dyspnoea					
		Economic Problem					
D), ORAL							
		Neuropathy		Atorvastatin (Atorvastatin)	C		
		Overweight					
		Pharmaceutical Product		Guaifenesin (Guaifenesin)	C		
		Complaint					
		Stress		Dyazide (Hydrochlorothiazide			
		Stress Fracture		, Triamterene)	C		
		Treatment Noncompliance		Salbutamol (Salbutamol)	C		
				Metoprolol Succinate (Metoprolol			
				Succinate)	C		
				Valsartan (Valsartan)	C		
				Human Mixtard (Insulin Human,			
				Insulin Human			
				Injection, Isophane)	C		

Date:05/03/05ISR Number: 4653182-0Report Type:Expedited (15-DaCompany Report #2005063329

Age:53 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Crying Depression	Consumer	Neurontin (Gabapentin)	PS		
350 MG (1 D)							
		Dizziness Pharmaceutical Product		Gabapentin (Gabapentin)	SS		

Complaint
Thyroidectomy

Levothyroxine Sodium
(Levothyroxine
Sodium) SS

Date:05/03/05ISR Number: 4653328-4Report Type:Direct
Age:81 YR Gender:Female I/FU:I

Company Report #CTU 247661

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 20 MG TWICE	Cerebral Haemorrhage Cerebrovascular Accident		Oxycontin 20 Mg Twice Daily	PS		
Initial or Prolonged DAILY Disability 300 MGS 3 DAILY	Contusion Eye Haemorrhage Fall Hip Fracture Upper Limb Fracture		Neurontin 300mgs 3 Daily	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/03/05ISR Number: 4653348-XReport Type:Expedited (15-DaCompany Report #2005006212

Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Amnesia	Foreign	Neurontin			
Other		Blood Thyroid Stimulating	Health	(Gabapentin)	PS		ORAL
1800 MG (600		Hormone Increased	Professional				
MG, 3 IN 1		Fatigue					
D), ORAL		High Density Lipoprotein		Estradiol			
		Increased		(Estradiol)	C		
		Hypercholesterolaemia		Dydrogesterone			
		Hypertriglyceridaemia		(Dydrogesterone)	C		
		Somnolence		Clonazepam			
		Weight Increased		(Clonazepam)	C		
				Tramadol			
				Hydrochloride			
				(Tramadol			
				Hydrochloride)	C		

Date:05/03/05ISR Number: 4653436-8Report Type:Expedited (15-DaCompany Report #2005052077

Age:81 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Foreign	Neurontin			
Other		Asthenia	Health	(Gabapentin)	PS		ORAL
1800 MG (600		Depression	Professional				
MG, 3 IN 1		Drug Dependence	Company				
D), ORAL		Drug Ineffective	Representative	Losartan Potassium			
		Psychomotor Retardation		(Losartan Potassium)	C		

Date:05/04/05ISR Number: 4652588-3Report Type:Expedited (15-DaCompany Report #GB-JNJFOC-20041007637

Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other
 OROPHARINGEAL 1.5MG 21 DAY
 OROPHARINGEAL 21 DAY
 OROPHARINGEAL
 OROPHARINGEAL
 OROPHARINGEAL
 OROPHARINGEAL
 OROPHARINGEAL

Risperdal PS
 Risperdal SS
 Clozapine SS
 Ranitidine SS
 Gabapentin SS
 Lamotrigine SS
 Lamotrigine SS

Date:05/04/05ISR Number: 4653362-4Report Type:Direct
 Age:56 YR Gender:Female I/FU:I

Company Report #CTU 247738

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	600 MG PO TID	Performance Status		Neurontin	PS		ORAL
		Decreased Sedation Therapeutic Response Unexpected With Drug Substitution		Synthroid	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/04/05ISR Number: 4653385-5Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 247713

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
600 MG @ HS		Drug Ineffective		Neurontin	PS		
		Therapeutic Response Unexpected With Drug Substitution					

Date:05/04/05ISR Number: 4653806-8Report Type:Expedited (15-DaCompany Report #KII-2005-0016305
 Age:12 MON Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Accidental Exposure	Study	Oxycontin			
		Agitation	Health	Tablets(Oxycodone			
		Aspiration	Professional	Hydrochloride) Cr			
ORAL		Cough	Other	Tablet	PS		ORAL
		Lethargy		Gabapentin			
ORAL		Miosis		(Gabapentin)	SS		ORAL
		Oxygen Saturation Decreased					
		Rhonchi					
		Vomiting					

Date:05/04/05ISR Number: 4653959-1Report Type:Expedited (15-DaCompany Report #2005063842
 Age:45 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cerebrovascular Accident	Foreign	Neurontin			
		Condition Aggravated	Health	(Gabapentin)	PS		ORAL
ORAL		Dysphagia	Professional				

Date:05/04/05ISR Number: 4654125-6Report Type:Expedited (15-DaCompany Report #2005042029
 Age:29 YR Gender:Male I/FU:F

Outcome Dose Other	Duration	PT Neutropenia	Report Source Foreign Health Professional	Product Gabapentin (Gabapentin)	Role PS	Manufacturer	Route ORAL
100 MG (100 MG, 1 IN 1 D), ORAL				Clozapine (Clozapine)	SS		ORAL
150 MG (150 MG, 1 IN 1 D)				Ranitidine Hydrochloride (Ranitidine Hydrochloride)	SS		
100 MG (100 MG, 1 IN 1 D), ORAL				Risperidone (Risperidone)	SS		ORAL
50 MG (50 MG, 1 IN 1 D), ORAL				Lamotrigine (Lamotrigine)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/04/05ISR Number: 4654190-6Report Type:Expedited (15-DaCompany Report #2005042541
Age:90 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abasia	Consumer	Neurontin			
		Asthenia		(Gabapentin)	PS		ORAL
		Condition Aggravated					
		Dysstasia		Terazosin	C		
		Essential Tremor		Lisinopril	C		
		Fall		Simvastatin	C		
		Feeling Abnormal		All Other			
		Movement Disorder		Therapeutic Products	C		

Date:05/04/05ISR Number: 4654206-7Report Type:Expedited (15-DaCompany Report #2005040445
Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disease Recurrence	Consumer	Neurontin (Tablets)			
		Petit Mal Epilepsy	Health	(Gabapentin)	PS		ORAL
		Pharmaceutical Product					
			Professional				
		Complaint					
				Insulin	C		
				Lamotrigine	C		
				Paxil (Paroxetine			
				Hydrochloride)	C		
				Clonazepam	C		

Date:05/04/05ISR Number: 4654207-9Report Type:Expedited (15-DaCompany Report #2005045468
Age:72 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pain In Extremity	Consumer	Neurontin			
		Peripheral Coldness	Health	(Gabapentin)	PS		ORAL
		300 MG (100					

MG, 3 IN 1 D) Prostate Cancer Professional
 ORAL
 Glipizide C
 Metformin
 Hydrochloride C

Date:05/04/05ISR Number: 4654740-XReport Type:Expedited (15-DaCompany Report #2005063472
 Age:78 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Circulatory Collapse Drug Dose Omission	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
1500 MG (1 D), ORAL		Grand Mal Convulsion	Professional				
		Treatment Noncompliance	Company Representative	Antihypertensives (Antihypertensives) Hmg Coa Reductase Inhibitors (Hmg Coa Reductase Inhibitors)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/04/05ISR Number: 4655044-1Report Type:Expedited (15-DaCompany Report #2005065732

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1200 MG (600	Anaemia Infection	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other MG, 2 IN 1 D)		Renal Failure	Professional				
ORAL				All Other Therapeutic Products (All Other Therapeutic Products)	SS		

Date:05/04/05ISR Number: 4655045-3Report Type:Expedited (15-DaCompany Report #2004075580

Age:23 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (300		Blood Pressure Increased Chest Pain	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
MG, 3 IN 1		Confusional State	Professional				
D), ORAL		Gingival Pain					
		Gingival Swelling Headache		Dilantin (Phenytoin Sodium)	SS		ORAL
300 MG (100		Laboratory Test Abnormal					
MG, 3 IN 1		Memory Impairment					
D),ORAL		Paraesthesia Tongue Biting		Topiramate (Topiramate)	SS		ORAL
300 MG (200MG		Tremor					
AM, 100MG		Vomiting					
PM), ORAL		Weight Increased		Citalopram			

Hydrobromide
(Citalopram
Hydrobromide) C

Date:05/05/05ISR Number: 4655431-1Report Type:Expedited (15-DaCompany Report #US015151
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 4 MG BID ORAL	Depressed Level Of Consciousness	Health Professional	Gabitril Neurontin	PS SS		ORAL ORAL
Initial or Prolonged 300 MG, ORAL	Drug Interaction Dyskinesia Tremor					

Date:05/05/05ISR Number: 4655532-8Report Type:Expedited (15-DaCompany Report #2005042704
Age:72 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (100 MG, 6 IN 1 D), ORAL	Anaemia Cataract Operation Dyspnoea Fatigue Feeling Abnormal Hypersensitivity Post Procedural Complication Restlessness Visual Disturbance	Consumer Health Professional	Neurontin (Gabapentin) Furosemide (Furosemide) Asasantin (Acetylsalicylic Acid, Dipyridamole) Sertraline	PS C C		ORAL

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Freedom Of Information (FOI) Report

Hydrochloride (Sertraline Hydrochloride)	C
Amlodipine Besilate (Amlodipine Besilate)	C
Zonisamide (Zonisamide)	C
Levothyroxine Sodium (Levothyroxine Sodium)	C
Losartan Potassium (Losartan Potassium)	C
Centrum Silver (Ascorbic Acid, Calcium, Minerals Nos, Retinol, Tocopheryl Acetate, Multivitamins (Ascorbic Acid/Ergocalciferol/ Folic Acid/Nicotinamide/Pa Calcium Carbonate (Calcium Carbonate)	C
Dicycloverine (Dicycloverine)	C

Date:05/05/05ISR Number: 4655551-1Report Type:Expedited (15-DaCompany Report #2004048907
Age:51 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abdominal Distension
Initial or Prolonged	Abdominal Pain
Other	Abnormal Behaviour Agitation Anger Anxiety Arterial Injury Atrial Fibrillation Biliary Tract Disorder Blood Calcium Decreased Body Dysmorphic Disorder Chest Pain Cognitive Disorder Condition Aggravated

Confusional State
Cough
Crying
Dependence On Respirator
Depressed Mood
Depression
Diaphragmatic Injury
Drug Screen Positive
Dysphagia
Faecaloma
Feeling Abnormal
Feeling Guilty
Gun Shot Wound
Haematocrit Decreased
Haemoglobin Decreased

Hydrochloride (Fluoxetine Hydrochloride)	C
Raloxifene Hydrochloride (Raloxifene Hydrochloride)	C
Lithium Carbonate (Lithium Carbonate)	C
Tramadol Hydrochloride (Tramadol Hydrochloride)	C
Atenolol (Atenolol)	C
Trazodone (Trazodone)	C
Propacet	

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(Dextropropoxyphene Napsilate, Paracetamol)	C
Estratest Hs (Estrogens Esterified, Methyltestosterone)	C
Estradiol (Estradiol)	C
Venlafaxine Hydrochloride (Venlafaxine Hydrochloride)	C
Nefazodone Hydrochloride (Nefazodone Hydrochloride)	C
Zolmitriptan (Zolmitriptan)	C
Sodium Fluoride (Sodium Fluoride)	C
Amoxicillin (Amoxicillin)	C
Meperidine W/Promethazine (Pethidine, Promethazine)	C
Azithromycin (Azithromycin)	C
Aquatab C (Dextromethorphan, Guaifenesin, Phenylpropanolamine)	C
Prednisone (Prednisone)	C
Pravastatin Sodium (Pravastatin Sodium)	C
Clindamycin (Clindamycin)	C
Chlorhexidine (Chlorhexidine)	C
Vicodin (Hydrochloride Bitartrate, Paracetamol)	C
Docusate Sodium (Docusate Sodium)	C
Risperidone	

(Risperidone)	C
Mirtazapine	
(Mirtazapine)	C
Bupropion	
Hydrochloride	
(Bupropion	
Hydrochloride)	C
Propranolol	
Hydrochloride	
(Propranolol	
Hydrochloride)	C
Sumatriptan	
Succinate	
(Sumatriptan	

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Succinate)	C
Methocarbamol	
(Methocarbamol)	C
Naproxen (Naproxen)	C
Narine Repetabe	
(Loratadine,	
Pseudoephedrine	
Sulfate)	C
Temazepam	
(Temazepam)	C
Quetiapine Fumarate	
(Quetiapine	
Fumarate)	C
Robitussin A-C /Old	
Form/ (Codeine	
Phosphate,	
Guaifenesin,	
Pheniramine Maleate)	C
Flurazepam	
Hydrochloride	
(Flurazepam	
Hydrochloride)	C
Mometasone Furoate	
(Mometasone Furoate)	C
Adapalene	
(Adapalene)	C
Ronatic	
(Chlorphenamine	
Tannate, Mepyramine	
Tannate,	
Phenylephrine	C
Modafinil	
(Modafinil)	C
Tizanidine	
Hydrochloride	
(Tizanidine	
Hydrochloride)	C
Estrogens Conjugated	
(Estrogens	
Conjugated)	C
Estradiol	
(Estradiol)	C
Tiagabine	
Hydrochloride	
(Tiagabine	
Hydrochloride)	C
Sildenafil Citrate	
(Sildenafil Citrate)	C

Butorphanol Tartrate
(Butorphanol
Tartrate) C
Trimethobenzamide
Hydrochloride
(Trimethobenzamide
Hydrochloride) C
Lithium Carbonate
(Lithium Carbonate) C
Valproate Semisodium
(Valproate
Semisodium) C
Beclometasone
Dipropionate

Freedom Of Information (FOI) Report

(Beclometasone
Dipropionate) C
Clavulin
(Amoxicillin
Trihydrate,
Clavulanate
Potassium) C
Midrid
(Dichloralphenazone,
Isometheptene,
Paracetamol) C

Date:05/05/05ISR Number: 4655553-5Report Type:Expedited (15-DaCompany Report #2004106809
Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Apnoea	Consumer	Neurontin			
		Atherosclerosis		(Gabapentin)	PS		
		Brain Oedema		Morphine (Morphine)	SS		
3 IN 1 D		Cerebral Disorder		Fentanyl (Fentanyl)	SS		
TRANSDERMAL	175 MG	(1 IN					
1 D),		Completed Suicide					
TRANSDERMAL		Euphoric Mood					
		Foaming At Mouth		Diamorphine			
		Laryngeal Disorder		(Diamorphine)	SS		
		Loss Of Consciousness		Citalopram			
		Mouth Haemorrhage		(Citalopram)	SS		
		Multiple Drug Overdose		Dextropropoxyphene			
		Muscle Rigidity		(Dextropropoxyphene)	SS		
		Polytraumatism					
		Pulmonary Congestion					
		Pulmonary Oedema					
		Pulmonary Vascular					
		Disorder					
		Rhinorrhoea					
		Ventricular Hypertrophy					
		Vomiting					

Date:05/05/05ISR Number: 4655583-3Report Type:Expedited (15-DaCompany Report #2005042698
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Anaemia Asthenia	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
100 MG (100 Other MG, 1 IN 1 D), ORAL		Body Height Decreased Coronary Artery Occlusion	Professional				
		Gastrointestinal Angiodysplasia Haemorrhagic Nausea Pharmaceutical Product Complaint		All Other Therapeutic Products (All Other Therapeutic Products) Acetylsalicylic Acid (Acetylsalicylic Acid)	SS SS		ORAL
ORAL				Lansoprazole (Lansoprazole) Diltiazem Hydrochloride (Diltiazem Hydrochloride) Alprazolam	C C		

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(Alprazolam)	C
Amitriptyline	
Hydrochloride	
(Amitriptyline	
Hydrochloride)	C
Tolterodine	
L-Tartrate	
(Tolterodine	
L-Tartrate)	C
Furosemide	
(Furosemide)	C
Atorvastatin	
(Atorvastatin)	C
Thiamine	
Hydrochloride	
(Thiamine	
Hydrochloride)	C
Paracetamol	
(Paracetamol)	C
Calcium (Calcium)	C

Date:05/05/05ISR Number: 4655852-7Report Type:Expedited (15-DaCompany Report #2005065410
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Foreign	Neurontin			
		Aphasia	Consumer	(Gabapentin)	PS		
		Brain Scan Abnormal		Morphine (Morphine)	C		
		Convulsion		Amitriptyline			
		Dysgeusia		Hydrochloride			
		Dysgraphia		(Amitriptyline			
		Oral Intake Reduced		Hydrochloride)	C		
		Personality Change Due To					
		A General Medical					
		Condition					
		Reading Disorder					
		Weight Decreased					

Date:05/06/05ISR Number: 4655842-4Report Type:Direct Company Report #CTU 247939
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Abnormal Behaviour Neurontin PS
1200MG TID

Aggression
Pharmaceutical Product
Complaint

Date:05/06/05ISR Number: 4656627-5Report Type:Expedited (15-DaCompany Report #8009828
Age:78 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 1000 MG /D PO		Inflammation	Foreign	Keppra	PS		ORAL
Initial or Prolonged 1 DF /D PO		Rash Morbilliform	Health	Movicol	SS		ORAL
			Professional	Oxycontin	SS		ORAL
20 MG /D PO			Other	Depakine	SS		ORAL
PO				Mopral	SS		ORAL
20 MG / D PO				Profenid	SS		ORAL
100 MG / D PO				Neurontin	SS		ORAL
PO				Praxilene	SS		ORAL
PO							

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Date:05/06/05ISR Number: 4657121-8Report Type:Expedited (15-DaCompany Report #2005042638

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 400 MG (100 Other MG, 4 IN 1 D), ORAL		Drug Ineffective Eczema Insomnia Mental Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
(0.25 MG), ORAL		Post-Traumatic Stress Disorder Weight Fluctuation		Xanax Tablet (Alprazolam)	SS		ORAL
ORAL				Hydroxyzine Pamoate (Caps) (Hydroxyzine Pamoate)	SS		ORAL
				Quetiapine Fumarate (Quetiapine Fumarate)	SS		
				Propylthiouracil (Propylthiouracil)	C		
				Lisinopril (Lisinopril)	C		
				Lekovit Ca (Calcium Carbonate, Colecalciferol)	C		

Date:05/06/05ISR Number: 4657124-3Report Type:Expedited (15-DaCompany Report #2005024066

Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 1200 MG (600 MG, 2 IN 1 D), ORAL		Completed Suicide Injury Asphyxiation	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

ORAL

Zoloft (Sertraline) SS

ORAL

Liothyronine Sodium
(Liothyronine Sodium) C
Levothyroxine Sodium
(Levothyroxine Sodium) C

Date:05/06/05ISR Number: 4657283-2Report Type:Expedited (15-DaCompany Report #2005046003

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer	Phenytoin			
		Diarrhoea		Suspension(Phenytoin			
		Drug Ineffective		Sodium)	PS		
300 MG							
		Nausea		Gabapentin			
		Transient Ischaemic		(Gabapentin)	SS		
		Attack		Lipitor			
				(Atorvastatin)	C		

Date:05/09/05ISR Number: 4658544-3Report Type:Expedited (15-DaCompany Report #2005068784

Age: Gender:Male I/FU:I

Outcome
Hospitalization -
Initial or Prolonged

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Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (300 MG, 1 IN 1 D), ORAL		Convulsion Drug Interaction Lethargy	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Lidocaine (Lidocaine)	SS		
				Insulin (Insulin) Cardiovascular System Drugs	C C		

Date:05/09/05ISR Number: 4658668-0Report Type:Expedited (15-DaCompany Report #GBR-2005-0001625
Age:78 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		C-Reactive Protein Increased Inflammation Rash Morbilliform	Foreign Health Professional Other	Oxycontin Tablets 10 Mg (Oxycodone Hydrochloride) Cr Tablet	PS		ORAL
20 MG, DAILY, ORAL				Movicol(Sodium Bicarbonate, Potassium Chloride, Sodium Chloride, Macrogol)	SS		ORAL
1 UNIT, DAILY, ORAL				Keppra (Levetiracetam)	SS		ORAL
1 GRAM DAILY, ORAL				Valproic Acid (Valproic Acid)	SS		ORAL
ORAL							

20 MG, DAILY,		Mopral (Omeprazole)	SS		ORAL
ORAL					
		Profenid "Rhone-Poulenc" (Ketoprofen)	SS	Rhone-Poulenc	ORAL
100 MG,					
DAILY, ORAL					
ORAL		Neurontin (Gabapentin)	SS		ORAL
ORAL		Praxilene (Naftidrofuryl Oxalate)	SS		ORAL

Date:05/09/05ISR Number: 4658774-0Report Type:Expedited (15-DaCompany Report #2005026776
Age:64 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Blindness	Foreign	Neurontin	PS		ORAL
2400 MG (800 Other MG, 3 IN 1 D), ORAL		Hypermetropia	Health				
		Macular Degeneration	Professional				
		Macular Oedema Retinopathy	Company Representative	Topiramate (Topiramate)	SS		ORAL
200 MG (1 D), ORAL			Other				
				Fentanyl (Fentanyl) Sertraline Hydrochloride (Sertraline)	C		

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Hydrochloride) C
 Clorazepate
 Dipotassium
 (Clorazepate
 Dipotassium) C
 Diazepam (Diazepam) C
 Clonazepam
 (Clonazepam) C

Date:05/09/05ISR Number: 4658791-0Report Type:Expedited (15-DaCompany Report #2005065422
 Age: Gender:Unknown I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG (300 MG, 1 IN 1 D), ORAL	Chest Pain Hot Flush Ocular Hyperaemia Pyrexia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:05/09/05ISR Number: 4658825-3Report Type:Expedited (15-DaCompany Report #2005067852
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (900 MG, 1 IN 1 D), ORAL	Blood Creatinine Increased Myoclonus	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL

Date:05/10/05ISR Number: 4658497-8Report Type:Direct Company Report #CTU 248082
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening	Back Pain		Gabapentin Capsules			

Other	Drug Ineffective	300 Mg, Mfd By Teva	PS	Teva	ORAL
300 MG ORALLY					
	Headache				
4X/ DAY	2 WK				
	Neck Pain	Bextra	C		
	Pain In Extremity	Hydrocodone	C		
	Shoulder Pain	Robaxin	C		
	Suicidal Ideation	Chlorzoxazone	C		
	Therapeutic Response	Protonix	C		
	Unexpected With Drug	Ambien	C		
	Substitution	Migrin-A	C		
	Tremor				

Date:05/10/05ISR Number: 4659058-7Report Type:Direct Company Report #CTU 248166
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Level Changed		Generic Gabapentin	PS		ORAL
600 MG TID,		Pharmaceutical Product					
P.O.		Complaint		Dilantin	C		
				Phenobarbital	C		

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Freedom Of Information (FOI) Report

Date:05/10/05ISR Number: 4660141-0Report Type:Expedited (15-DaCompany Report #2005067797

Age:41 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Aseptic Necrosis Bone	Consumer	Neurontin			
Other		Hypertension		(Gabapentin)	PS		ORAL
1200 MG (300		Malaise					
MG, 4 IN 1							
D), ORAL							
				All Other			
				Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	SS		
				Sertaline			
				Hydrochloride			
				(Sertraline			
				Hydrochloride)	C		
				Zolpidem Tartrate			
				(Zolpidem Tartrate)	C		

Date:05/11/05ISR Number: 4658627-8Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0379673A

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Abortion Induced		Lamictal	PS	Glaxosmithkline	ORAL
Other		Drug Exposure During		Neurontin	SS		ORAL
		Pregnancy					

Date:05/11/05ISR Number: 4658628-XReport Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0379673B

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Drug Exposure During		Lamictal	PS	Glaxosmithkline	
Congenital Anomaly		Pregnancy		Neurontin	SS		
		Holoprosencephaly					

Freedom Of Information (FOI) Report

Potassium
(Potassium) C
Warfarin (Warfarin) C
Terazosin
(Terazosin) C
Nifedipine
(Nifedipine) C
All Other
Therapeutic Products
(All Other
Therapeutic
Products) C
Simvastatin
(Simvastatin) C

Date:05/11/05ISR Number: 4660944-2Report Type:Expedited (15-DaCompany Report #2005070451
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Economic Problem Hypertonic Bladder	Consumer	Neurontin (Gabapentin)	PS		ORAL
1800 MG (600 MG, 1 IN 3D), ORAL		Hypoaesthesia Oral Hysterectomy					
2.5 MG (2.5 MG, 1 IN 1 D), ORAL		Post Procedural Complication Treatment Noncompliance		Chlorpropamide (Chlorpropamide)	SS		ORAL
4 MG (4 MG, 1 IN 1 D), ORAL				Tolterodine (Tolterodine L-Tartrate) (Tolterodine)	SS		ORAL
				Amitriptyline Hydrochloride (Amitriptyline			

100 MG 9100

Hydrochloride) SS

ORAL

MG , 1 IN 1

D), ORAL

Karvea Hct
 (Hydrochlorothiazide
 , Irbesartan) SS
 Metformin
 Hydrochloride
 (Metformin
 Hydrochloride) C
 Rosuvastatin
 (Rosuvastatin) C
 Atorvastatin
 (Atorvastatin) C

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

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Blood Alkaline Phosphatase Increased Blood Creatine	Study	Hydromorphone Hcl	PS		ORAL
ORAL		Phosphokinase Increased	Health Professional	Bupropion	SS		ORAL
ORAL		Blood Creatinine Increased Blood Ph Decreased	Other	(Amfebutamone)	SS		ORAL
ORAL		Blood Pressure Fluctuation		Citalorпам (Citalorпам)	SS		ORAL
ORAL		Blood Urea Increased Bradycardia		Trazodone (Trazodone)	SS		ORAL
ORAL		Cardio-Respiratory Arrest Confusional State Electromechanical		Norvasc (Amlodipine Besilate)	SS		ORAL
ORAL		Dissociation Haemodynamic Instability		Neurontin (Gabapentin)	SS		ORAL
ORAL		Hypoxia		Propranolol	SS		ORAL
ORAL		Loss Of Consciousness Myoclonus		Methocarbamol (Methocarbamol)	SS		ORAL
ORAL		Pneumonia		Herbal Preparation	SS		ORAL
		Sinus Tachycardia Vomiting		Benzodiazepine Derivatives	SS		

Date:05/11/05ISR Number: 4660975-2Report Type:Expedited (15-DaCompany Report #8010302

Age:2 DY Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - TRANSPLACENTAL	TRP	Convulsion Neonatal	Health	Levetiracetam "Ucb"	PS	Ucb	
Initial or Prolonged TRANSPLACENTAL	TRP	Drug Exposure During	Professional	Tegretol	SS		
TRANSPLACENTAL	TRP	Pregnancy		Klonopin	SS		

TRANSPLACENTAL	TRP	Clozaril	SS
TRANSPLACENTAL	TRP	Diazepam	SS
TRANSPLACENTAL	TRP	Neurontin	SS
TRANSPLACENTAL	TRP	Lamictal	SS
TRANSPLACENTAL	TRP	Trileptal	SS
TRANSPLACENTAL	TRP	Phenobarbital	SS
TRANSPLACENTAL	TRP	Dilantin	SS
TRANSPLACENTAL	TRP	Topamax	SS
TRANSPLACENTAL	TRP	Tridione	SS
TRANSPLACENTAL	TRP	Depakote	SS

Date:05/11/05ISR Number: 4660990-9Report Type:Expedited (15-DaCompany Report #2005045777
Age:72 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Ankle Fracture
Initial or Prolonged	Back Pain
Congenital Anomaly	Cataract
Other	Eye Laser Surgery
	Feeling Abnormal
	Hand Deformity
	Herpes Zoster
	Intestinal Obstruction
	Joint Range Of Motion
	Decreased
	Ligament Injury
	Pain In Extremity
	Pharmaceutical Product
	Complaint

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Post Laminectomy Syndrome Scoliosis Spinal Compression Fracture Vertebroplasty	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, 3 IN 1 D)			Consumer Health Professional	Neurontin (Gabapentin)	PS		
				Amlodipine Besilate (Amlodipine Besilate)	C		
				Valsartan (Valsartan)	C		
				Carbamazepine (Carbamazepine)	C		
				Levothyroxine (Levothyroxine)	C		
				Lansoprazole (Lansoprazole)	C		
				Montelukast Sodium (Montelukast Sodium)	C		
				Clindamycin (Clindamycin)	C		
				Hydrocodone (Hydrocodone)	C		
				Seretide Mite (Fluticasone Propionate, Salmeterol Xinafoate)	C		
				Combivent (Ipratropium Bromide, Salbutamol Sulfate)	C		
				Oxycodone Hydrochloride (Oxycodone Hydrochloride)	C		
				Cyclobenzaprine (Cyclobenzaprine)	C		
				Risedronate Sodium (Risedronate Sodium)	C		
				Tolterodine L-Tartrate (Tolterodine L-Tartrate)	C		

Vicodin (Hydrocodone
Bitartrate,
Paracetamol) C
Selenium (Selenium) C
Ascorbic Acid
(Ascorbic Acid) C
Multivitamins
(Ascorbic
Acid/Ergocalciferol/
Folic
Acid/Nicotinamide/Pa C
Calcium Citrate
(Calcium Citrate) C
Becosym Forte
(Nicotinamide,
Pyridoxine
Hydrochloride,

Freedom Of Information (FOI) Report

Riboflavin, Thiamine C
 Cetirizine
 Hydrochloride
 (Cetirizine
 Hydrochloride) C
 Mometasone Furoate
 (Mometasone Furoate) C
 Tolterodine-L
 Tartrate
 (Tolterodine-L
 Tartrate) C
 Proctofoam Hc
 (Hydrocortisone
 Acetate, Pramocaine
 Hydrochloride) C

Date:05/11/05ISR Number: 4661053-9Report Type:Expedited (15-DaCompany Report #2005067549
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1.5 MG (0.5 Other MG, 3 IN 1 D), ORAL	Anger Anxiety Disorder	Consumer	Xanax Tablet (Alprazolam)	PS		ORAL
		Aphagia Drug Dependence Drug Ineffective Drug Withdrawal Syndrome		Neurontin (Gabapentin)	SS		ORAL
		Emotional Distress Intentional Misuse		Geodon (Ziprasidone)	SS		ORAL
		Overdose Phobia Unevaluable Event		Obetrol (Amfetamine Aspartate, Amfetamine Sulfate, Dexamfetamine Saccharate, Fluoxetine Hydrochloride (Fluoxetine Hydrochloride)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Venlafaxine
Hydrochloride) C
Clonazepam
(Clonazepam) C
Zolpidem Tartrate
(Zolpidem Tartrate) C

Date:05/11/05ISR Number: 4661056-4Report Type:Expedited (15-DaCompany Report #2005045395
Age:77 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	500 MG (100 Other MG, 5 IN 1 D), ORAL	Grand Mal Convulsion Pancreatitis	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Pharmaceutical Product Complaint	Professional				

Ranitidine
Hydrochloride
(Ranitidine
Hydrochloride) C
Methylphenobarbital
(Methylphenobarbital
) C
Nabumetone
(Nabumetone) C
Atenolol (Atenolol) C
Paroxetine
Hydrochloride
(Paroxetine
Hydrochloride) C

Date:05/11/05ISR Number: 4661082-5Report Type:Expedited (15-DaCompany Report #PTWYE359621JAN05
Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Dizziness Drug Withdrawal Syndrome Feeling Cold Feeling Hot	Foreign Health Professional Other	Efexor Xr (Venlafaxine Hydrochloride, Capsule, Extended			

150 MG 1X PER	Hyperhidrosis	Release, 0)	PS	ORAL
1 DAY ORAL	Malaise			
600 MG 1X PER		Gabapentin (Gabapentin, , 0)	SS	ORAL
1 DAY ORAL				
80 MG 1X PER		Propranolol Hydrochloride (Propranolol Hydrochloride, , 0)	SS	ORAL
1 DAY ORAL				

Date:05/11/05ISR Number: 4661104-1Report Type:Expedited (15-DaCompany Report #2005069871
Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Convulsion	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 2 IN 1 D), ORAL			Professional				
				Senna (Senna) Glycerol (Glycerol)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Thiamine (Thiamine) C
 Dihydrocodeine
 (Dihydrocodeine) C
 Meloxicam
 (Meloxicam) C

Date:05/11/05ISR Number: 4661771-2Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 248280

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - 300MG PO BID 5 DAY	Dizziness		Gabapentin 300mg	PS		ORAL
Initial or Prolonged	Fall		Meclize	C		
Other	Hallucination		Bortezomib	C		

Date:05/12/05ISR Number: 4660662-0Report Type:Expedited (15-DaCompany Report #FR-ABBOTT-05P-056-0297333-00
 Age:78 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged 2 DAY	C-Reactive Protein Increased		Micropakine Granule Profenid	PS SS		ORAL ORAL
8 DAY	Rash Morbilliform		Nulytely	SS		ORAL
9 DAY	Systemic Inflammatory Response Syndrome		Levetiracetam Oxycodone Hydrochloride	SS SS		ORAL ORAL
4 DAY			Omeprazole	SS		ORAL
			Gabapentin Naftidrofuryl Oxalate	SS SS		ORAL ORAL

Date:05/12/05ISR Number: 4661781-5Report Type:Direct
 Age:22 YR Gender:Male I/FU:I

Company Report #CTU 248359

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						

400 MG TID	Abnormal Behaviour	Neurontin	PS
800 MG TID	Convulsion	Neurontin	SS

Date:05/12/05ISR Number: 4661983-8Report Type:Expedited (15-DaCompany Report #2005056882
 Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Back Injury	Consumer	Neurontin			
Other		Blood Glucose Increased	Health	(Gabapentin)	PS		ORAL
300 MG (300		Blood Pressure Increased	Professional				
MG, 1 IN 1		Drug Ineffective					
D), ORAL		Joint Injury		Morphine (Morphine)	C		
		Neck Injury		Morphine Sulfate			
		Paraesthesia		(Morphine Sulfate)	C		
		Road Traffic Accident		Prinzide			
				(Hydrochlorothiazide			
				, Lisinopril)	C		
				Methylphenidate			
				Hydrochloride			
				(Methylphenidate			
				Hydrochloride)	C		
				Metformin			
				Hydrochloride			
				(Metformin			

Furosemide
(Furosemide) C
Tizanidine
(Tizanidine) C
Prednisone
(Prednisone) C

Date:05/12/05ISR Number: 4662111-5Report Type:Expedited (15-DaCompany Report #2005070434

Age:51 YR Gender:Female I/FU:I

Outcome PT
Hospitalization - Agoraphobia
Initial or Prolonged Blood Cholesterol
Disability Increased
Other Body Height Decreased
Bone Density Decreased
Cerebrovascular Accident
Communication Disorder
Condition Aggravated
Convulsion

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, 3 IN 1 D)		Disturbance In Attention Economic Problem Insomnia Intervertebral Disc Degeneration Mental Disorder Nervousness Osteoarthritis Panic Attack Treatment Noncompliance	Consumer	Neurontin (Gabapentin)	PS		
				Estrogens Conjugated (Estrogens Conjugated) Fluoxetine Hydrochloride (Fluoxetine Hydrochloride) Quetiapine Fumarate (Quetiapine Fumarate)	C C C		

Date:05/12/05ISR Number: 4662124-3Report Type:Expedited (15-DaCompany Report #2005045792
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 200 MG (100 MG, 2 IN 1 D), ORAL		Bacterial Infection Fungal Infection Pneumonia	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Fluconazole (Fluconazole) Allegra-D (Fexofenadine, Pseudoephedrine Hydrochloride) Rosiglitazone Maleate (Rosiglitazone Maleate) Metoprolol (Metoprolol) Enalapril Maleate	C C C C		

(Enalapril Maleate)	C
Warfarin Sodium	
(Warfarin Sodium)	C
Famotidine	
(Famotidine)	C
Granisetron	
(Granisetron)	C
Lansoprazole	
(Lansoprazole)	C
Ondansetron	
Hydrochloride	
(Ondansetron	
Hydrochloride)	C
Pyridoxine	
Hydrochloride	
(Pyridoxine	
Hydrochloride)	C
Vicodin (Hydrocodone	
Bitartrate,	
Paracetamol)	C
Paracetamol	
(Paracetamol)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Temazepam
 (Temazepam) C
 Promethazine
 Hydrochloride
 (Promethazine
 Hydrochloride) C
 Chemotherapy Nos
 (Chemotherapy Nos) C

Date:05/12/05ISR Number: 4662228-5Report Type:Expedited (15-DaCompany Report #2005070314
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Health Professional	Neurontin (Gabapentin)	PS		

Date:05/12/05ISR Number: 4662534-4Report Type:Expedited (15-DaCompany Report #2005068428
 Age: Gender:I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abortion Induced	Foreign Health Professional	Neurontin (Gabapentin)	PS		
Congenital Anomaly		Drug Exposure During Pregnancy		Lamotrigine (Lamotrigine)	SS		
		Holoprosencephaly					
		Mental Impairment					

Date:05/13/05ISR Number: 4661329-5Report Type:Expedited (15-DaCompany Report #PHNU2005DE01944
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Circulatory Collapse		Exelon	PS	Novartis Sector: Pharma	ORAL
3 mg/day							
Unknown				Neurontin	SS		ORAL
Unknown				Nephral	C		

Unknown

Detrusitol C

Unknown

Madopar C

Date:05/13/05ISR Number: 4663200-1Report Type:Expedited (15-DaCompany Report #2005071190

Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1600 MG (800 Other MG, 2 IN 1 D)		Cardiac Operation Chest Pain Dyspnoea	Health Professional	Neurontin (Gabapentin)	PS		
				Atenolol (Atenolol)	C		
				Metformin Hydrochloride/Rosigl itazone (Metformin Hydrochloride, Rosiglitazone)	C		
				Rosuvastatin (Rosuvastatin)	C		
				Insulin Injection, Isophane (Insulin Injection, Isophane)	C		
				Insulin (Insulin)	C		
				Ultracet			

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Freedom Of Information (FOI) Report

(Paracetamol,
Tramadol
Hydrochloride) C

Date:05/13/05ISR Number: 4663228-1Report Type:Expedited (15-DaCompany Report #2005043174
Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 MG, 2 IN 1 D), ORAL		Condition Aggravated Electric Shock Gastric Disorder Neuropathy	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Pharmaceutical Product Complaint Spinal Fusion Surgery		Allopurinol (Allopurinol) Levothyroxine Sodium (Levothyroxine Sodium) Omeprazole (Omeprazole)	C C C		

Date:05/13/05ISR Number: 4663251-7Report Type:Expedited (15-DaCompany Report #2004065552
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 300MG (100 MG, 3 IN 1 Disability D), Other 1 MG (0.5 MG, 2 IN 1 D), UNKNOWN		Abdominal Pain Abdominal Tenderness Candidiasis Coagulopathy Disability Drug Ineffective Gastrointestinal Injury Gun Shot Wound Haemorrhage	Consumer	Neurontin (Gabapentin) Clonazepam (Clonazepam)	PS SS		
				Oxycocet (Oxycodone)			

3 TABLETS (3	Haemothorax	Hydrochloride,	
	Hypotension	Paracetamol)	SS
IN 1 D),	Major Depression		
UNKNOWN	Mental Disorder		
	Overdose	Vicodin (Hydrocodone	
	Pain	Bitartrate,	
	Polytraumatism	Paracetamol)	SS
60 MG (10 MG,	Post Procedural		
1 IN 4 HR),	Complication		
UNKNOWN	Postoperative Abscess	Pantoprazole	
	Splenic Injury	(Pantoprazole)	C
	Staphylococcal Infection	Latanoprost	
	Suicide Attempt	(Latanoprost)	C
		Esomeprazole	
		(Esomeprazole)	C
		Gemfibrozil	
		(Gemfibrozil)	C
		Pravastatin Sodium	
		(Pravastatin Sodium)	C
		Fluticasone	
		Propionate	
		(Fluticasone	
		Propionate)	C
		Gaviscon (Sodium	
		Alginate, Sodium	
		Bicarbonate)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/13/05ISR Number: 4663343-2Report Type:Expedited (15-DaCompany Report #2005059757

Age:28 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Foreign	Neurontin			
ORAL		Drug Exposure During	Study	(Gabapentin)	PS		ORAL
		Pregnancy	Health	Folic Acid (Folic			
		Metrorrhagia	Professional	Acid)	C		
		Pregnancy					

Date:05/16/05ISR Number: 4663723-5Report Type:Expedited (15-DaCompany Report #2004117393

Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Alcohol Use	Consumer	Neurontin			
Initial or Prolonged		Anxiety		(Gabapentin)	PS		
900 MG (300		Condition Aggravated					
Disability		Depression		Trazodone			
MG, 3 IN 1 D)		Drug Ineffective		(Trazodone)	C		
Other		Dysthymic Disorder		Mirtazapine			
		Facial Bones Fracture		(Mirtazapine)	C		
		Gun Shot Wound		Naltrexone			
		Insomnia		(Naltrexone)	C		
		Irritability					
		Overdose					
		Poisoning					
		Soft Tissue Injury					
		Stress					
		Suicide Attempt					

Date:05/16/05ISR Number: 4663724-7Report Type:Expedited (15-DaCompany Report #2003115340

Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Abnormal Behaviour
Other	Agitation
	Anorexia

Anxiety
Arthritis
Balance Disorder
Dehydration
Depression
Diabetes Mellitus
Non-Insulin-Dependent
Diarrhoea
Economic Problem
Emotional Disorder
Hostility
Impulsive Behaviour
Injury
Insomnia
Mania
Monoplegia
Mood Swings
Muscle Spasms
Obesity
Pneumonia
Social Problem
Suicidal Ideation

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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
			Suicide Attempt Tremor	Consumer	Neurontin (Gabapentin)	PS		

Date:05/16/05ISR Number: 4663795-8Report Type:Expedited (15-DaCompany Report #2005072171
Age:50 YR Gender:Male I/FU:I

Dose	Duration	Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Other 1D, ORAL			Angioneurotic Oedema	Consumer	Celebrex (Celecoxib)	PS		ORAL
10 MG (10 MG, 1 IN 1 D), ORAL			Diarrhoea Disease Progression Drug Effect Decreased		Neurontin (Gabapentin)	SS		ORAL
2400 MG (800 MG, 3 IN 1 D)			Fibromyalgia Gastrointestinal Pain Inflammation Musculoskeletal Disorder Osteoarthritis		Bextra (Valdecoxib) Ibuprofen	SS		
					Oxycodone Hydrochloride Oxycocet (Oxycodone Hydrochloride, Paracetamol) Sertraline Hydrochloride	C C C		

Date:05/16/05ISR Number: 4663799-5Report Type:Expedited (15-DaCompany Report #2004043537
Age:60 YR Gender:Male I/FU:F

Dose	Duration	Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 600 MG (300			Anxiety Cognitive Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL

Initial or Prolonged	Completed Suicide			
MG, 2 IN 2				
Other	Depressed Level Of			
D), ORAL				
	Consciousness	Midazolam	SS	
INTRAVENOUS	INTRAVENOUS			
	Depression	Citalopram	SS	
	Fatigue	Tramadol		
ORAL	Injury	Hydrochloride	SS	ORAL
	Libido Decreased	Pancuronium	SS	
INTRAVENOUS	INTRAVENOUS			
	Major Depression	All Other		
	Neuropathy Peripheral	Therapeutic Products	SS	
	Pain	Zolpidem Tartrate	C	
	Respiratory Distress	Ibuprofen	C	
	Self-Medication	Mirtazapine	C	
	Sleep Disorder	Clonazepam	C	
	Sluggishness			
	Somnolence			

Date:05/16/05ISR Number: 4664067-8Report Type:Direct Company Report #CTU 248687
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Burning Sensation		Gabapentin 400 Mg	PS		ORAL
1 PO TID		Condition Aggravated					
		Therapeutic Response					
		Unexpected With Drug					
		Substitution					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/16/05ISR Number: 4664105-2Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 248646

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Insomnia		Gabapentin 300 Mg Bid	PS		

Date:05/16/05ISR Number: 4664218-5Report Type:Expedited (15-DaCompany Report #2005071473
 Age:67 YR Gender:Unknown I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG (1200 MG, 2 IN 1 D)		Circadian Rhythm Sleep Disorder Dyspnoea Sedation	Foreign Health Professional	Gabapentin (Gabapentin) Ramipril (Ramipril) Bisoprolol (Bisoprolol) Amiodarone (Amiodarone) Ipratropium Bromide (Ipratropium Bromide) Hydrocortisone (Hydrocortisone) Ciprofloxacin (Ciprofloxacin)	PS C C C C C C		

Date:05/16/05ISR Number: 4664274-4Report Type:Direct
 Age:45 YR Gender:Male I/FU:I

Company Report #CTU 248627

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 300 MG ONE AM , TWO N , TWO		Pharmaceutical Product Complaint		Neurontin - Generic	PS		

HS

Date:05/16/05ISR Number: 4664655-9Report Type:Expedited (15-DaCompany Report #2004242949TH
Age:49 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 20 MG (20 MG, Initial or Prolonged 1 IN 1 D), Other ORAL	Blood Cholesterol Increased Blood Glucose Increased Blood Triglycerides Increased Drug Eruption	Foreign Health Professional	Bextra (Valdecoxib) Neurontin (Gabapentin) (Gabapentin)	PS SS		ORAL ORAL
300 MG (300 MG, 1 IN 1 D), ORAL	Drug Hypersensitivity Gouty Arthritis Low Density Lipoprotein Increased Skin Lesion Stevens-Johnson Syndrome		Colchicine (Colchicine)	C		

Date:05/17/05ISR Number: 4663529-7Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0376424A
Age:69 YR Gender:Female I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged	Abnormal Dreams Cognitive 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
8MG per day	12 DAY	Confusional State Disorientation Hallucination, Visual		Zophren	PS	Glaxosmithkline	ORAL
INTRAVENOUS	2250MG per	Hypersomnia		Zovirax	SS	Glaxosmithkline	
day	6 DAY	Incoherent Metabolic Encephalopathy		Polaramin	SS		
INTRAVENOUS	15MG per day 12 DAY	Psychomotor Retardation		Mopral	SS	Glaxosmithkline	ORAL
20MG Per day	14 DAY	Retrograde Amnesia		Primperan	SS	Glaxosmithkline	
INTRAVENOUS	60MG per day			Neurontin	SS		ORAL
4 DAY				Fraxodi	C	Glaxosmithkline	
SUBCUTANEOUS				Fungizone	C		
UNKNOWN				Chemotherapy	C		
UNKNOWN							

Date:05/17/05ISR Number: 4665156-4Report Type:Expedited (15-DaCompany Report #2005073479
Age:45 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Arrhythmia	Foreign	Neurontin			
Other		Blood Pressure	Consumer	(Gabapentin)	PS		ORAL
900 MG (300		Fluctuation					
MG, 3 IN 1		Cardiac Failure					
D), ORAL		Gastric Perforation Spinal Laminectomy		Ixel (Milnacipran)	C		

Date:05/17/05ISR Number: 4665298-3Report Type:Expedited (15-DaCompany Report #2004106748
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	600 MG (300 Other MG, 2 IN 1 D)	Abnormal Behaviour Agitation	Consumer	Neurontin (Tablets) (Gabapentin)	PS		
		Condition Aggravated					
		Delusion		Haldol (Haloperidol)	SS		
		Depression		Risperidone			
		Drug Ineffective		(Risperidone)	SS		
6 MG (3 MG, 2 IN 1 D)		Homicidal Ideation					
		Intentional Misuse		Metformin			
		Multiple Drug Overdose		(Metformin)	C		
		Psychotic Disorder					
		Suicide Attempt					
		Treatment Noncompliance					

Date:05/17/05ISR Number: 4665301-0Report Type:Expedited (15-DaCompany Report #2005007423
Age:25 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Accident At Work
Initial or Prolonged	Anxiety
Other	Arthralgia
	Back Injury
	Delusion
	Depression
	Drug Abuser
	Drug Ineffective
	Drug Withdrawal Syndrome
	Intentional Misuse
	Joint Sprain
	Migraine

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Morbid Thoughts Multiple Drug Overdose Nephrolithiasis	Consumer	Neurontin (Gabapentin)	PS		
		Panic Attack Paranoia Psychotic Disorder Skin Laceration Spousal Abuse Suicidal Ideation Suicide Attempt Tearfulness Varicocele	Consumer	Methadone Hydrochloride (Methadone Hydrochloride) Klonopin (Clonazepam) Oxycontin (Oxycodone Hydrochloride)	SS SS		

Date:05/17/05ISR Number: 4665304-6Report Type:Expedited (15-DaCompany Report #2005003387
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Other		Cervical Vertebral Fracture Conjunctivitis Corneal Ulcer Difficult To Wean From Ventilator Drug Ineffective Glasgow Coma Scale Abnormal Gun Shot Wound Head Injury Nosocomial Infection Pneumonia Haemophilus Self Injurious Behaviour Subdural Haematoma Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:05/17/05ISR Number: 4665305-8Report Type:Expedited (15-DaCompany Report #2004106719
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour	Consumer	Neurontin			

Freedom Of Information (FOI) Report

4 MG, 1-2
 EVERY 4-6
 HOURS AS
 NECESSARY,
 ORAL

Hydrochloride) SS

Prednisone
 (Prednisone) C
 Fentanyl (Fentanyl) C
 Felodipine
 (Felodipine) C
 Clonidine
 (Clonidine) C
 Ramipril (Ramipril) C
 Potassium
 (Potassium) C
 Lansoprazole
 (Lansoprazole) C
 Clonazepam
 (Clonazepam) C
 Ursodeoxycholic Acid
 (Ursodeoxycholic
 Acid) C
 Nateglinide
 (Nateglinide) C
 Lorazepam (Lorazepam) C

Date:05/17/05ISR Number: 4665320-4Report Type:Expedited (15-DaCompany Report #2004077855
 Age:14 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Affective Disorder Aggression	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG ORAL		Alcoholism Anger Depression Drug Abuser Dysarthria Feeling Drunk Gait Disturbance		Levothyroxine Sodium (Levothyroxine Sodium) Clindamycin (Clindamycin) Quetiapine Fumarate (Quetiapine	C C		

Hypothyroidism	Fumarate)	C
Injury	Lithium (Lithium)	C
Learning Disorder	Multivitamins	
Loss Of Consciousness	(Ascorbic Acid,	
Pain	Ergocalciferol,	
Psychotic Disorder	Folic Acid,	
Schizoaffective Disorder	Nicotinamide,	C
Suicide Attempt	Paroxetine	
	Hydrochloride	
	(Paroxetine	
	Hydrochloride)	C
	Clozapine	
	(Clozapine)	C
	Lorazepam	
	(Lorazepam)	C
	Docusate Sodium	
	(Docusate Sodium)	C
	Modafinil	
	(Modafinil)	C
	Topiramate	
	(Topiramate)	C
	Naltrexone	

Freedom Of Information (FOI) Report

(Naltrexone) C
 Medroxyprogesterone
 Acetate
 (Medroxyprogesterone
 Acetate) C
 Salbutamol
 (Salbutamol) C

Date:05/17/05ISR Number: 4665321-6Report Type:Expedited (15-DaCompany Report #2004058042
 Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin			
Other		Completed Suicide	Health	(Gabapentin)	PS		ORAL
300 MG QHS, MAY INCREASE TO Q8H OVER ONE, ORAL		Depression	Professional				
		Drug Ineffective					
		Gun Shot Wound					
		Hypoaesthesia		Bextra	C		
		Injury		Medrol			
		Insomnia		(Methylprednisolone)	C		
		Nervousness		Hydrocodone			
		Pain		(Hydrocodone)	C		
		Paraesthesia		Ultracet			
		Stress		(Paracetamol,			
		Suicidal Ideation		Tramadol			
		Weight Decreased		Hydrochloride)	C		
				Cyclobenzaprine			
				(Cyclobenzaprine)	C		

Date:05/17/05ISR Number: 4665322-8Report Type:Expedited (15-DaCompany Report #2004070397
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1200 MG (300 Disability MG, 4 IN 1		Abnormal Behaviour Agitation	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Anger					

Other D), ORAL	Anxiety			
2 MG (0.5 MG, 4 IN 1 D), ORAL	Depression Drug Ineffective Drug Tolerance Intentional Misuse Orgasm Abnormal Overdose Sexual Dysfunction Somnolence Stress At Work Suicidal Ideation Suicide Attempt	Clonazepam (Clonazepam)	SS	ORAL
		Prozac (Fluoxetine Hydrochloride) Eskalith (Lithium Carbonate)	C C	

Date:05/17/05ISR Number: 4665517-3Report Type:Expedited (15-DaCompany Report #2004058034
Age:50 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Anger
Initial or Prolonged	Anxiety
Disability	Chest Pain
Other	Delusion
	Drug Ineffective
	Drug Screen Positive

Date:05/17/05ISR Number: 4665554-9Report Type:Expedited (15-DaCompany Report #2004012957
Age:49 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 600 MG (300 Disability MG, 2 IN 1 D) Other	Accident Deformity Face Injury Gun Shot Wound Mental Disorder Pain Polytraumatism Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:05/17/05ISR Number: 4665556-2Report Type:Expedited (15-DaCompany Report #2004083654
Age:61 YR Gender:Female I/FU:F

Outcome	PT
Death Other	Anxiety Completed Suicide Depression

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Dose	Duration	Injury Pain	Report Source	Product	Role	Manufacturer	Route
ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:05/17/05ISR Number: 4666361-3Report Type:Expedited (15-DaCompany Report #2004052118
Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Acute Respiratory	Consumer	Neurontin (Gabapentin)	PS		
Other		Distress Syndrome		Venlafaxine			
		Aggression		Hydrochloride			
		Anger		(Venlafaxine			
		Anoxic Encephalopathy		Hydrochloride)	C		
		Anxiety		Risperidone			
		Brain Oedema		(Risperidone)	C		
		Cardio-Respiratory Arrest					
		Completed Suicide					
		Injury					
		Injury Asphyxiation					
		Loss Of Consciousness					
		Pain					
		Pneumomediastinum					
		Pneumonia Aspiration					
		Pneumonitis					
		Prescribed Overdose					
		Respiratory Failure					
		Vomiting					

Date:05/18/05ISR Number: 4665005-4Report Type:Direct Company Report #CTU 249041
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aggression		Gabapentin 600 Mg 2			
PO 2 TID		Pharmaceutical Product		Tid	PS		ORAL
		Complaint					
		Therapeutic Response					

Unexpected With Drug
Substitution

Date:05/18/05ISR Number: 4665389-7Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 249023

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Drug Effect Decreased		Gabapentin			
		Drug Intolerance		(Neurontin)	PS		
INTRAVENOUS	600 MG TWO 9	Pharmaceutical Product					
AM IV Q HS		Complaint					

Date:05/18/05ISR Number: 4665654-3Report Type:Direct
Age:51 YR Gender:Female I/FU:I

Company Report #CTU 248975

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Therapeutic Response		Gabapentin 300 Mg			
Other		Unexpected With Drug		Ivax	PS	Ivax	ORAL
300 MG 3		Substitution					
TIMES DAILY							

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Freedom Of Information (FOI) Report

ORAL

Date:05/18/05ISR Number: 4666215-2Report Type:Expedited (15-DaCompany Report #2005016316

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anorexia	Foreign	Neurontin			
SEE IMAGE		Asthma	Consumer	(Gabapentin)	PS		ORAL
150 MG (75		Blood Glucose Increased	Health	Topamax (Topiramate)	SS		ORAL
MG, 2 IN 1		Blood Pressure Increased	Professional				
D), ORAL		Chronic Fatigue Syndrome					
		Complex Regional Pain Syndrome		Prednisone (Prednisone)	C		
		Condition Aggravated		Elavil			
		Drug Ineffective		(Amitriptyline Hydrochloride)	C		
		Fatigue		Flexeril			
		Fibromyalgia		(Cyclobenzaprine Hydrochloride)	C		
		Headache		Rivotril			
		Hepatic Enzyme Increased		(Clonazepam)	C		
		Incision Site Complication					
		Incisional Hernia Repair					
		Muscle Atrophy					
		Nausea					
		Pharmaceutical Product Complaint					
		Post Procedural Complication					
		Weight Increased					

Date:05/18/05ISR Number: 4666216-4Report Type:Expedited (15-DaCompany Report #2005015729

Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Dose Omission	Foreign	Gabapentin (Tablets)			
600 MG (600		Drug Withdrawal Syndrome	Health	(Gabapentin)	PS		ORAL

MG, 1 IN 1	Feeling Hot And Cold	Professional			
D), ORAL	Malaise				
150 MG (150	Syncope		Venlafaxine (Venlafaxine)	SS	ORAL
MG, 1 IN 1					
D), ORAL			Propranolol (Propranolol)	SS	ORAL
80 MG					
(DAILY), ORAL					

Date:05/18/05ISR Number: 4666224-3Report Type:Expedited (15-DaCompany Report #2005071513
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Life-Threatening (300 MG), ORAL	Anaphylactic Reaction Blood Pressure Diastolic Decreased Pharyngeal Oedema Swollen Tongue	Foreign Health Professional	Gabapentin (Gabapentin)	PS		ORAL
			Panadeine Co (Codeine Phosphate, Paracetamol)	C		

FDA - Adverse Event Reporting System (AERS)

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Date:05/18/05ISR Number: 4666530-2Report Type:Expedited (15-DaCompany Report #2005071771

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG (300 Other MG, 2 IN 1 D), ORAL	Condition Aggravated Grand Mal Convulsion Wrong Technique In Drug Usage Process	Health Professional	Neurontin (Solution) (Gabapentin)	PS		ORAL
			Depakote (Valproate Semisodium)	C		
			Tegretol (Carbamazepine)	C		
			Diazepam (Diazepam)	C		
			Synthroid (Levothyroxine Sodium)	C		
			Oscal 500-D (Calcium, Colecalciferol)	C		
			Pancrelipase (Pancrelipase)	C		
			Periostat (Doxycycline Hyclate)	C		
			Ferrous Sulfate (Ferrous Sulfate)	C		
			Nasonex (Mometasone Furoate)	C		
			Azelaic Acid (Azelaic Acid)	C		
			Atrovent (Ipratropium Bromide)	C		
			Motrin (Ibuprofen)	C		
			Isosurce (Carbohydrates Nos, Fats Nos, Minerals Nos, Proteins Nos, Vitamin Nos)	C		
			Promod (Lecithinum, Whey Proteins)	C		
			Neutrogena T/Gel Therapeutic Shampoo			

Date:05/18/05ISR Number: 4666533-8Report Type:Expedited (15-DaCompany Report #2005071384
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hip Arthroplasty	Consumer	Neurontin			
		Therapy Non-Responder		(Gabapentin)	PS		
300 MG (100							
MG, 3 IN 1 D)							

Date:05/19/05ISR Number: 4668590-1Report Type:Expedited (15-DaCompany Report #2005071187
Age:84 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Grand Mal Convulsion
Initial or Prolonged	Pharmaceutical Product
Other	Complaint

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Freedom Of Information (FOI) Report

Dose	Duration	Status Epilepticus Tremor Underdose	Report Source	Product	Role	Manufacturer	Route
100 MG (100 MG, 1 IN 1 D), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
				Zocor (Simvastatin)	C		
				Dilantin (Phenytoin Sodium)	C		
				Apirin (Acetylsalicylic Acid)	C		
				Potassium Chloride (Potassium Chloride)	C		

Date:05/19/05ISR Number: 4668633-5Report Type:Expedited (15-DaCompany Report #2005045397
Age:50 YR Gender:Female I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1 D), ORAL		Burning Sensation Constipation	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Feeling Abnormal	Professional				
		Muscle Spasms					
		Paraesthesia Pharmaceutical Product Complaint Poor Quality Drug Administered		Estradiol (Estradiol)	C		
				Verapamil (Verapamil)	C		
				Motrin (Ibuprofen)	C		
				Lorazepam (Lorazepam)	C		
				Flexeril (Cyclobenzaprine Hydrochloride)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 400 MG (1D), Other ORAL	Bipolar Disorder Convulsion Difficulty In Walking Dizziness Drug Ineffective Gait Disturbance Increased Appetite Insomnia Metabolic Disorder Migraine Suicidal Ideation Weight Increased	Consumer	Neurontin (Gabapentin) Lamictal (Lamotrigine) Zyprexa (Olanzapine) Seroquel (Quetiapine Fumarate) Depakote (Valproate Sodium) Depakene (Valproate Sodium) Claritin (Loratadine)	PS SS SS SS SS C C		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/19/05ISR Number: 4668738-9Report Type:Expedited (15-DaCompany Report #2005073892
Age:93 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Eye Haemorrhage Iris Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Mydriasis Pain Visual Acuity Reduced Vitreous Opacities		Antihypertensives (Antihypertensives) Cholesterol- And Triglyceride Reducers (Cholesterol- And Triglyceride	C C		

Date:05/19/05ISR Number: 4668748-1Report Type:Expedited (15-DaCompany Report #2005073888
Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Haemorrhage Intracranial	Health Professional	Neurontin (Gabapentin)	PS		ORAL
300 MG (300 MG, 1 IN 1 D), ORAL				Warfarin (Warfarin)	SS		ORAL
2 MG (2 MG, 1 IN 1 D), ORAL				Digoxin (Digoxin) Atenolol (Atenolol)	C C		

Date:05/19/05ISR Number: 4668749-3Report Type:Expedited (15-DaCompany Report #2005072525
Age:67 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Drug Ineffective Feeling Abnormal	Consumer	Neurontin (Gabapentin)	PS		

Other
 200 MG (200
 MG, 1 IN 1 D)

Pulmonary Thrombosis
 Seroma
 Thrombosis
 Weight Increased

Celebrex (Celecoxib) SS
 Estrogens
 (Estrogens) SS
 Synthroid
 Levothyroxine
 (Synthroid
 Levothyroxine) C

Date:05/20/05ISR Number: 4667382-7Report Type:Expedited (15-DaCompany Report #GB-ABBOTT-05P-167-0300147-00
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Gingival Hyperplasia		Epilim Tablets	PS		ORAL
UNKNOWN				Gabapentin	SS		
				Metformin	C		
				Gliclazide	C		

Date:05/20/05ISR Number: 4669970-0Report Type:Expedited (15-DaCompany Report #2005075005
 Age: Gender:Female I/FU:I

Outcome	PT
Disability	Back Pain
Other	Blood Alkaline Phosphatase Increased

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1.5 MG (0.5 MG, 3 IN 1 D), ORAL		Blood Cholesterol Increased Confusional State Dependence Fluid Retention Foot Fracture High Density Lipoprotein Decreased Intervertebral Disc Disorder Memory Impairment	Consumer	Xanax Tablet (Alprazolam)	PS		ORAL
500 MG (100 MG, 3 IN 1 D), ORAL		Oedema Peripheral Vitreous Detachment Weight Increased Weight Loss Poor		Neurontin (Gabapentin)	SS		ORAL
				Axotal (Old Form) (Butalbital, Caffeine, Paracetamol)	SS		
				Irbesartan (Irbesartan)	C		
				Lansoprazole (Lansoprazole)	C		

Date:05/20/05ISR Number: 4670128-XReport Type:Expedited (15-DaCompany Report #2005074294

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Aphasia Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
2400 MG (1 D), ORAL		Hyperhidrosis					
		Paralysis Skin Discolouration		Xanax Tablet (Alprazolam)	SS		
				Remeron (Mirtazapine)	C		
				Periactin (Cyproheptadine Hydrochloride)	C		

Date:05/20/05ISR Number: 4673223-4Report Type:Direct Company Report #CTU 249216
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Neurontin	PS		
Other		Chills Headache Nausea Vomiting					

Date:05/23/05ISR Number: 4672810-7Report Type:Expedited (15-DaCompany Report #2005073373
 Age:57 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (900 Other MG, 1 IN 1 D), ORAL		Drug Level Increased Dysarthria Dyskinesia Lower Respiratory Tract Infection Renal Failure Acute	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
20 MG (20 MG, 1 IN 1 D), ORAL				Piroxicam (Piroxicam)	SS		ORAL
				Antibiotics (Antibiotics)	SS		

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Salbutamol	
(Salbutamol)	C
Estradiol	
(Estradiol)	C
Dihydrocodeine	
(Dihydrocodeine)	C
Omeprazole	
(Omeprazole)	C
Methocarbamol	
(Methocarbamol)	C
Beclomethasone	
(Beclometasone)	C
Amitriptyline	
(Amitriptyline)	C

Date:05/23/05ISR Number: 4672885-5Report Type:Expedited (15-DaCompany Report #2005055287
 Age:54 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 1 D ORAL	Anxiety	Consumer	Zoloft (Sertraline)	PS		ORAL
Initial or Prolonged Other 300 MG ORAL	Carpal Tunnel Syndrome Cartilage Injury		Neurontin (Gabapentin)	SS		ORAL
30 MG (30 MG, 1 IN 1 D)	Drug Interaction Gastrooesophageal Reflux Disease Interstitial Lung Disease		Duloxetine Hydrochloride (Duloxetine Hydrochloride)	SS		ORAL
ORAL	Loose Body In Joint Muscle Disorder					
	Myocardial Infarction Nightmare Polytraumatism Road Traffic Accident Sleep Apnoea Syndrome Weight Increased		Oxygen (Oxygen) All Other Therapeutic Products (All Other Therapeutic Products) Glyceryl Trinitrate (Glyceryl Trinitrate)	SS SS SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Body Height Decreased	Consumer	Neurontin			
Hospitalization -		Cardiac Failure		(Gabapentin)	PS		ORAL
600 MG (200							
Initial or Prolonged		Congestive					
MG, 3 IN 1							
		Chronic Obstructive					
D), ORAL							
		Pulmonary Disease		Seretide Mite			
		Pharmaceutical Product		(Fluticasone			
		Complaint		Propionate,			
		Pulmonary Oedema		Salmeterol			
				Xinafoate)	C		
				All Other			
				Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	C		
				Oxygen (Oxygen)	C		

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Date:05/23/05ISR Number: 4672980-0Report Type:Expedited (15-DaCompany Report #2005049492

Age:73 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Abasia Crying	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
3600 MG (600 MG, 6 IN 1 D), ORAL		Disease Recurrence Drug Ineffective Middle Insomnia Nephrectomy Neuropathic Pain Nocturia Pharmaceutical Product Complaint	Professional	Gabapentin (Tablets) (Gabapentin)	SS		ORAL
				Furosemide (Furosemide)	C		
				Glipizide (Glipizide)	C		
				Fenofibrate (Fenofibrate)	C		
				Lansoprazole (Lansoprazole)	C		
				Isosorbide (Isosorbide)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Metoprolol Succinate (Metoprolol Succinate0	C		
				Trandolapril (Trandolapril)	C		
				Nortriptyline (Nortriptyline)	C		
				Warfarin Sodium (Warfarin Sodium)	C		
				Metformin Hydrochloride (Metformin Hydrochloride)	C		

Date:05/23/05ISR Number: 4673723-7Report Type:Expedited (15-DaCompany Report #2005072861

Age:56 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Condition Aggravated	Health	Neurontin			
Other		Drug Ineffective	Professional	(Gabapentin)	PS		ORAL
300 MG (100		Nerve Compression					
MG, 3 IN 1 D)		Nerve Injury					
ORAL		Pharmaceutical Product		Gabapentin			
100 MG (100		Complaint		(Gabapentin)	SS		
MG, 3 IN 1 D)		Therapeutic Response					
		Unexpected With Drug		Motrin (Ibuprofen)	C		
		Substitution		Extra Strength			
				Tylenol			
				(Paracetamol)	C		

Date:05/23/05ISR Number: 4673739-0Report Type:Expedited (15-DaCompany Report #2005074293
Age:45 YR Gender:Female I/FU:I

Outcome	PT
Other	Brain Damage
	Condition Aggravated
	Drug Effect Decreased

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Drug Interaction Feeling Abnormal Insomnia	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL (800 MG,), ORAL		Intervertebral Disc Protrusion Nerve Compression Neuropathy Pain Pain In Extremity		Motrin (Ibuprofen)	SS		ORAL
		Shoulder Pain Treatment Noncompliance		Hydrocodone (Hydrocodone) Mobic (Meloxicam)	SS SS		ORAL
				Ultram (Tramadol Hydrochloride)	C		

Date:05/23/05ISR Number: 4673742-0Report Type:Expedited (15-DaCompany Report #2005072689
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Condition Aggravated Diarrhoea Gastric Disorder	Consumer	Bextra (Valdecoxib) Neurontin (Gabapentin)	PS SS		ORAL
900 MG (300 MG, 3 IN 1 D); ORAL		Headache Hypertension					
UNKNOWN (200 MG, UNKNOWN); ORAL		Ligament Injury Osteoarthritis Spinal Fracture		Celebrex (Celecoxib)	SS		ORAL
				All Other Therapeutic Products	C		
				All Other Therapeutic Products	C		

Vicodin (Hydrocodone
Bitartrate,
Paracetamol) C
Propacet
(Dextropropoxyphene
Napsilate,
Paracetamol) C

Date:05/24/05ISR Number: 4674078-4Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 249363

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
P.O 800 MG		Neuropathic Pain		Gabapentin 800mg.	PS		
ONCE TID		Skin Odour Abnormal					
(SWITCHED TO		Therapeutic Response					
GENERIC 2		Unexpected With Drug					
MONTHS AGO)		Substitution					
		Urine Odour Abnormal		Keppra	C		

Date:05/24/05ISR Number: 4674130-3Report Type:Direct
Age:57 YR Gender:Female I/FU:I

Company Report #CTU 249372

Outcome	PT
Hospitalization - Initial or Prolonged	Dyskinesia Hallucination Mental Status Changes Muscle Contractions

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FDA - Adverse Event Reporting System (AERS)

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Involuntary
Muscle Twitching

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
600 MG PO	24 HR		Gabapentin	PS		ORAL
			Methadone	C		
			Reglan	C		
			Novolin Ins	C		
			Actos	C		
			Demerol	C		
			Allopurinol	C		
			Humulin R	C		
			Verapamil Sr	C		
			Bis. Na. Bicarb	C		
			Synthroid	C		
			Clonidine	C		
			Amitriptyline	C		

Date:05/24/05ISR Number: 4674153-4Report Type:Expedited (15-DaCompany Report #KII-2005-0016647
Age:30 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Agitation Bowel Sounds Abnormal Coma Disorientation Hypotension	Study Health Professional Other	Hydromorphone Hcl (Similar To Nda 21-044) (Hydromorphone Hydrochloride)	PS		ORAL
ORAL		Mydriasis Tachycardia		Methadone (Methadone)	SS		ORAL
ORAL				Gabapentin (Gabapentin)	SS		ORAL
ORAL				Amitriptyline (Amitriptyline)	SS		ORAL
ORAL				Baclofen (Baclofen)	SS		ORAL
ORAL				Benzodiazepine Derivatives ()	SS		ORAL

ORAL				Antiepileptics()	SS		ORAL
				Warfarin (Warfarin)	SS		ORAL
ORAL							

Date:05/24/05ISR Number: 4674225-4Report Type:Expedited (15-DaCompany Report #2005074695
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Diverticulitis Hypoaesthesia Medication Error Pharmaceutical Product Complaint Visual Disturbance	Consumer	Neurontin (Gabapentin)	PS		

Date:05/24/05ISR Number: 4674274-6Report Type:Expedited (15-DaCompany Report #2005063453
 Age:80 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (300 MG, 3 IN 1 D), ORAL		Condition Aggravated Thrombophlebitis Superficial Varicose Vein	Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Prednisone			

Freedom Of Information (FOI) Report

60 MG 91 IN 1

(Prednisone) SS

D)

Valtrex 9valciclovir
Hydrochloride) SS

3 GRAM 91

GRAM, 3 IN 1

D)

Date:05/25/05ISR Number: 4673773-0Report Type:Expedited (15-DaCompany Report #FR-ABBOTT-05P-056-0297333-00
Age:78 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2 DAY	C-Reactive Protein Increased		Micropakine Granule Profenid	PS SS		ORAL ORAL
	8 DAY	Rash Morbilliform		Nulytely	SS		ORAL
	9 DAY	Systemic Inflammatory Response Syndrome		Levetiracetam Oxycodone Hydrochloride	SS SS		ORAL ORAL
	4 DAY			Omeprazole	SS		ORAL
				Gabapentin Naftidrofuryl Oxalate	SS SS		ORAL ORAL

Date:05/25/05ISR Number: 4674020-6Report Type:Expedited (15-DaCompany Report #2005056028
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2100 MG (900 MG, 3 IN 1	Drug Withdrawal Syndrome Post Procedural Complication	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Thinking Abnormal	Company				

D), ORAL

Representative

All Other
Therapeutic Products
(All Other
Therapeutic
Products) C

Date:05/25/05ISR Number: 4674046-2Report Type:Expedited (15-DaCompany Report #CA-2005-007813
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Dyspnoea Mobility Decreased Nausea Oedema Peripheral	Foreign Consumer Other	Betaseron(Intreferon Beta - 1b) Injection, 250ug	PS		
SUBCUTANEOUS	8 MIU,	EVERY					
2D,		Pyrexia					
SUBCUTANEOUS				Gabapentin (Gabapentin)	SS		
				Baclofen	C		
				Zanaflex (Tizanidine Hydrochloride)	C		
				Amitriptyline (Amitriptyline)	C		
				Diovan Novartis (Valsartan)	C		
				Nexium	C		
				Tylenol Extra-Strength	C		

Freedom Of Information (FOI) Report

Date:05/25/05ISR Number: 4674066-8Report Type:Expedited (15-DaCompany Report #KII-2005-0016638
 Age:56 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Agitation	Study	Oxycontin Tablets			
Initial or Prolonged	Alanine Aminotransferase	Health	(Oxycodone			
Other	Increased	Professional	Hydrocholride) Cr			
	Anion Gap Increased	Other	Tablet	PS		
	Aspartate		Gabapentin			
	Aminotransferase		(Gabapentin)	SS		
	Increased		Metaxalone			
	Blood Amylase Increased		(Metaxalone)	SS		
	Blood Bicarbonate		Orubal	SS		
	Decreased		Zolpidem	SS		
	Blood Creatine		Clonazepam(Clonazepa			
	Phosphokinase Increased		m)	SS		
	Blood Creatine		Modafinil(Modafinil)	SS		
	Phosphokinase Mb		Oral Antidiabetics	SS		
	Increased		Biguanides	SS		
	Blood Creatinine		Niacin (Nicotinic			
	Increased		Acid)	SS		
	Blood Glucose Increased		Ace Inhibitor	SS		
	Blood Lactate		Lidocaine			
	Dehydrogenase Increased		(Lidocaine)	SS		
	Blood Ph Decreased		Thiazides	SS		
	Blood Potassium Increased		Ssri	SS		
	Blood Urea Increased		Antidepressants	SS		
	Coma		Antihistamines For			
	Drug Screen Positive		Systemic Use	SS		
	Haemodialysis		Antibiotics	SS		
	Haemoglobin Decreased		Monistat(Miconazole			
	Lipase Increased		Nitrate)	SS		
	Prothrombin Time		Norethindrone			
	Prolonged		(Norethisterone)	SS		
	Renal Function Test		Albuterol			
	Abnormal		Sulfate(Salbutamol			
			Sulfate)	SS		
			Terbutaline And			
			Other Beta-2 Agonist	SS		
			Warfarin (Warfarin)	SS		
			Fentanyl (Fentanyl)	SS		
			Rhinocort Aqua			
			(Budesonide)	SS		
			Ethanol(Ethanol)	SS		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abnormal Behaviour	Consumer	Neurontin			
Disability		Anhedonia		(Gabapentin)	PS		
Other		Anxiety					
		Emotional Disorder					
		Mental Disorder					
		Pain					
		Pharmaceutical Product					
		Complaint					
		Polytraumatism					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/25/05ISR Number: 4674962-1Report Type:Expedited (15-DaCompany Report #2005071384

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Effect Decreased Hip Arthroplasty	Consumer	Neurontin (Gabapentin)	PS		
300 MG (100 MG, 3 IN 1 D)		Neuropathy		Procardia (Nifedipine)	C		
				Lipitor (Atorvastatin)	C		

Date:05/25/05ISR Number: 4675300-0Report Type:Expedited (15-DaCompany Report #2005075182

Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Other		Drug Interaction	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
1800 MG (300 MG, 6 IN 1 D), ORAL			Professional	Methadone Hydrochloride (Methadone Hydrochloride)	SS		ORAL
			Other	Debrisoquine (Debrisoquine)	SS		
				Tramadol (Tramadol)	C		
				Morphine (Morphine)	C		
				Co-Proxamol	C		
				Pethidine (Pethidine)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Appendix Disorder	Foreign	Neurontin			
Other		Drug Withdrawal Syndrome	Health	(Gabapentin)	PS		ORAL
900 MG (300		Hyperhidrosis	Professional				
MG, 3 IN 3		Surgery					
D), ORAL		Tremor		Hydroxocobalamin (Hydroxocobalamin)	C		
				Lorazepam (Lorazepam)	C		
				Meloxicam (Meloxicam)	C		
				Omeprazole (Omeprazole)	C		
				Paroxetine (Paroxetine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/25/05ISR Number: 4676467-0Report Type:Expedited (15-DaCompany Report #2005071955
 Age:41 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Hypertension	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
400 MG (200		Retinal Disorder					
MG, 1 IN 2		Vision Blurred	Professional				
D), ORAL		Visual Disturbance					
				Naxen (Naproxen)	C		
				Muscle Relaxants	C		

Date:05/25/05ISR Number: 4676472-4Report Type:Expedited (15-DaCompany Report #2005074861
 Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Ventricular Tachycardia	Foreign Health	Gabapentin (Gabapentin)	PS		
3600 MG (900							
MG, 4 IN 1			Professional				
D), UNKNOWN							

Date:05/25/05ISR Number: 4676499-2Report Type:Expedited (15-DaCompany Report #2005075577
 Age:67 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Disability		Arthralgia	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
600 MG (200		Condition Aggravated					
MG, 3 IN 3			Professional				
D), ORAL							
				Tolterodine (Tolterodine)	C		

Date:05/25/05ISR Number: 4676507-9Report Type:Expedited (15-DaCompany Report #2005076936
Age:57 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 2.4 GRAM (1 IN 1 D), ORAL	Toxic Skin Eruption	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:05/25/05ISR Number: 4676525-0Report Type:Expedited (15-DaCompany Report #2005065422
Age:37 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 300 MG (300 MG, 1 IN 1 D); ORAL	Chest Pain Eyelid Oedema Hot Flush Ocular Hyperaemia Pyrexia	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Date:05/25/05ISR Number: 4676590-0Report Type:Expedited (15-DaCompany Report #2005063472
Age:68 YR Gender:Male I/FU:F

Outcome	PT	Report Source
Other	Drug Dose Omission Fall Grand Mal Convulsion	Foreign Health Professional Company

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Representative
Other

Dose	Duration	Product	Role	Manufacturer	Route
1200 MG (400 MG, 3 IN 1 D), ORAL		Neurontin (Gabapentin)	PS		ORAL
		Antihypertensives (Antihypertensives) Statins (Hmg Coa Reductase Inhibitors) Panadeine Forte	C		
		(Codeine Phosphate, Paracetamol)	C		

Date:05/25/05ISR Number: 4676596-1Report Type:Expedited (15-DaCompany Report #2005049031
Age:82 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Facial Palsy Meningitis Metastases To Meninges Tonic Clonic Movements	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Aricept (Donepezil)	PS C		

Date:05/26/05ISR Number: 4675153-0Report Type:Expedited (15-DaCompany Report #2005031290
Age:76 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 10 MG (10 MG,		Drug Ineffective Feeling Drunk	Consumer Health	Lipitor (Atorvastatin)	PS		ORAL

1 IN 1 D),	Feeling Hot And Cold	Professional		
	Insomnia			
ORAL	Neuropathy Peripheral		Neurontin	
	Osteoarthritis		(Gabapentin)	SS ORAL
ORAL	Spinal Fracture		Prozac (Fluoxetine	
	Vision Blurred		Hydrochloride)	SS ORAL
ORAL			Avapro	C
			Digitex (Digoxin)	C
			Coumadine (Warfarin	
			Sodium)	C
			Lexapro	
			(Escitalopram)	C
			Trileptal	
			Ciba-Geigy	
			(Oxcarbazepine)	C Ciba-Geigy
			Potassium Chloride	
			(Potassium Chloride)	C
			Dyazide	
			(Hydrochlorothiazide	
			, Triamterene)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/26/05ISR Number: 4675169-4Report Type:Expedited (15-DaCompany Report #2005051608
 Age:34 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 3600 MG (1200 Other MG, 3 IN 1 D), ORAL	Depressed Level Of Consciousness Respiratory Arrest Sleep Disorder Suicide Attempt	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Strattera (Atomoxetine Hydrochloride)	C		
			Heroin (Diamorphine)	C		
			Cocaine (Cocaine)	C		
			Cannabis (Cannabis)	C		
			Valium (Diazepam)	C		
			Vicodin (Hydrocodone Bitartrate, Paracetamol)	C		
			Percocet (Oxycodone Hydrochloride, Paracetamol)	C		
			Nexium (Esomeprazole)	C		

Date:05/26/05ISR Number: 4675178-5Report Type:Expedited (15-DaCompany Report #2005008387
 Age:92 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG (300 MG, 2 IN 1 D), ORAL	Confusional State Dysarthria Hyperventilation Mental Status Changes Mobility Decreased Speech Disorder Visual Acuity Reduced	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Insulin (Insulin)	SS		
			Calcium (Calcium)	C		
			Coumadin "Boots" (Warfarin Sodium) Amelorad	C		

(Eicosapentaenoic
Acid, Gamolenic
Acid) C
Protonix C
(Pantoprazole)
Miacalcin C
(Citcitonin, Salmon)
Vitamin C (Vitamin
C) C
Multivitamins C
(Multivitamins)

Date:05/26/05ISR Number: 4676657-7Report Type:Expedited (15-DaCompany Report #2004059901

Age:50 YR Gender:Female I/FU:F

Outcome	PT
Death	Acute Respiratory Failure
Hospitalization -	Agitation
Initial or Prolonged	Anxiety
Other	Aortic Injury
	Completed Suicide
	Haemothorax
	Lung Injury

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Disease Recurrence		Gabapentin 400mg			
		Drug Effect Decreased		(Generic)	PS		
400MG ONE BID		Therapeutic Response					
TWO QHS		Unexpected With Drug Substitution					

Date:05/26/05ISR Number: 4687351-0Report Type:Expedited (15-DaCompany Report #2004043537
Age:55 YR Gender:Male I/FU:F

Outcome	PT
Death	Anxiety
Hospitalization -	Cognitive Deterioration
Initial or Prolonged	Completed Suicide
Other	Depressed Level Of
	Consciousness
	Fatigue
	Feeling Jittery
	Head Discomfort
	Intentional Misuse
	Libido Decreased
	Major Depression
	Mental Disorder
	Pain
	Pain In Extremity
	Paraesthesia
	Respiratory Distress

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Freedom Of Information (FOI) Report

Dose	Duration	Restlessness Sleep Disorder Sluggishness Somnolence	Report Source	Product	Role	Manufacturer	Route
600 MG (300 MG, 2 IN 1 D), ORAL			Consumer	Neurontin (Gabapentin)	PS		ORAL
	INTRAVENOUS			Midazolam	SS		
	INTRAVENOUS			Citalopram Pancuronium	SS SS		
	INTRAVENOUS			Ultram (Tramadol Hydrochloride)	SS		ORAL
				All Other Therapeutic Products Ambien (Zolpidem Tartrate) Ibuprofen Klonopin Remeron (Mirtazapine)	SS C C C C		

Date:05/27/05ISR Number: 4676738-8Report Type:Expedited (15-DaCompany Report #2005076985
Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Confusional State Inadequate Analgesia Memory Impairment Neuropathic Pain Sciatica	Consumer	Neurontin (Gabapentin)	PS		

Date:05/27/05ISR Number: 4676863-1Report Type:Expedited (15-DaCompany Report #2004106719
Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Asthenia	Consumer	Neurontin			
Life-Threatening		Completed Suicide		(Gabapentin)	PS		ORAL
1500 MG (300							
Hospitalization -		Depression					
MG, 5 IN 1							
Initial or Prolonged		Dizziness					
D), ORAL							
Other		Drug Ineffective		Diphenhydramine			
		Drug Toxicity		(Diphenhydramine)	SS		ORAL
ORAL							
		Electroencephalogram		Metoprolol			
20 MG (1 IN 1		Abnormal		(Metoprolol)	SS		
D)							
		Fatigue					
		Self Injurious Behaviour		Carisoprodol			
(350 MG, 1 OR		Suicidal Ideation		(Carisoprodol)	SS		ORAL
2 EVERY							
4-6HOURS AS							
NECESSARY),							
ORAL							
				Meprobamate			
				(Meprobamate)	SS		
				Hydromorphone			
				Hydrochloride			
(4 MG, 1-2				(Hydromorphone			
EVERY				Hydrochloride)	SS		ORAL
4-6HOURS AS							
NECESSARY),							
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ORAL

Prednisone	
(Prednisone)	C
Duragesic (Fentanyl)	C
Plendil (Felodipine)	C
Catapres	
(Clonidipine)	C
Altace (Ramipril)	C
Potassium	
(Potassium)	C
Prevacid	
(Lansoprazole)	C
Clonazepam	
(Clonazepam)	C
Ursodiol	
(Ursodeoxycholic	
Acid)	C
Starlix	
(Nateglinide)	C
Lorazepam	
(Lorazepam)	C
Albuterol	
(Salbutamol)	C
Lasix (Furosemide)	C

Date:05/27/05ISR Number: 4677759-1Report Type:Direct
 Age:21 YR Gender:Female I/FU:I

Company Report #CTU 249883

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion		Neurontin	PS		
300 MG 2 TID		Therapeutic Response Unexpected With Drug Substitution					

Date:05/27/05ISR Number: 4677766-9Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 249863

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required Intervention to		Blindness Confusional State		Neurontin (Gabapentine)			

Prevent Permanent
300 MG PO QID
Impairment/Damage

Dizziness
Therapeutic Response
Unexpected With Drug
Substitution

(Generic) PS
Digoxin C
Warfarin C
Diazepam C
Hydrocodone C
Kcl C
Paxil C

ORAL

Date:05/27/05ISR Number: 4677923-1Report Type:Direct
Age:42 YR Gender:Female I/FU:I

Company Report #CTU 249868

Outcome
Life-Threatening
Hospitalization -
Initial or Prolonged
Disability
Required
Intervention to
Prevent Permanent

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Impairment/Damage

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900MG 4X DAY		Drug Dose Omission Suicide Attempt		Neurontin 300 Mg 3600 Mg Day	PS		

Date:05/27/05ISR Number: 4678159-0Report Type:Expedited (15-DaCompany Report #2005078236
Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Gingival Hyperplasia	Foreign Health	Gabapentin (Gabapentin)	PS		
1200 MG (1 IN 1 D)			Professional				
800 MG (1 IN 1 D), ORAL				Epilim (Valproate Sodium)	SS		ORAL
				Metformin (Metformin)	C		
				Gliclazide (Gliclazide)	C		

Date:05/27/05ISR Number: 4678242-XReport Type:Expedited (15-DaCompany Report #2004058042
Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other (300 MG QHS, MAY INCREASE TO Q8H OVER ONE), ORAL		Anxiety Completed Suicide	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Depression	Professional				
		Drug Ineffective					
		Injury					
		Insomnia		Valdecoxib			

Nervousness	(Valdecoxib)	C
Stress	Medrol	
Suicidal Ideation	(Methylprednisolone)	C
Weight Decreased	Hydrocodone	
	(Hydrocodone)	C
	Ultracet	
	(Paracetamol,	
	Tramadol	
	Hydrochloride)	C
	Cyclobenzaprine	
	(Cyclobenzaprine)	C

Date:05/27/05ISR Number: 4678279-0Report Type:Expedited (15-DaCompany Report #2005057468

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1200 MG (400 MG, 3 IN 1 D), ORAL	Anxiety Confusional State Delirium Drug Interaction Paraesthesia Sensory Disturbance	Health Professional Company Representative	Neurontin (Tablets) (Gabapentin) Fluoxetine (Fluoxetine) All Other Therapeutic Products (All Other Therapeutic Products)	 PS SS C		ORAL

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Freedom Of Information (FOI) Report

Voltaren (Diclofenac Sodium) C
 Baclofen (Baclofen) C
 Nitrofurantoin (Nitrofurantoin) C
 Alprazolam (Alprazolam) C
 Omeprazol (Omeprazole) C
 Enoxaparin (Heparin) C
 Lactulose (Lactulose) C
 Bisacodyl (Bisacodyl) C
 Effortil (Etilefrine Hydrochloride) C
 Multivitamins (Multivitamins) C

Date:05/27/05ISR Number: 4678282-0Report Type:Expedited (15-DaCompany Report #2005077350
 Age:77 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG, 3 IN 1 D), ORAL		Cardiac Failure Condition Aggravated	Foreign Consumer	Gabapentin (Gabapentin)	PS		ORAL

Dextropropoxifen (Dextropropoxyphen) C
 Acenocoumarol (Acenocoumarol) C

Date:05/27/05ISR Number: 4678300-XReport Type:Expedited (15-DaCompany Report #2005077209
 Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Amnesia	Consumer	Gabapentin			

Initial or Prolonged	Incorrect Dose	(Gabapentin)	PS	ORAL
3600 MG (1200				
Other	Administered			
MG, 3 IN 1	Neuropathic Pain			
D), ORAL				
		Percocet (Oxycodone Hydrochloride, Paracetamol)	SS	ORAL
ORAL				

Date:05/27/05ISR Number: 4678417-XReport Type:Expedited (15-DaCompany Report #2005041846
Age:69 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Adverse Drug Reaction	Health	Neurontin			
		Meningitis Chemical	Professional	(Gabapentin)	PS		ORAL
1000 MG (2 IN		Rash					
1 D), ORAL		Spinal Disorder		Metoprolol			
				(Metoprolol)	C		
				Tegretol			
				(Carbamazepine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4678633-7Report Type:Expedited (15-DaCompany Report #2005076431

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (600 MG, 1/2 TABLET 3 TIMES DAILY), ORAL		Deep Vein Thrombosis Oedema Peripheral	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL
				Domepridone (Domperidone) Mist Carminative (Camphor, Cardamom Oil, Rhizoma Zingiberæ, Tinctura Capscii) Diclofenac (Diclofenac Sodium)	C C C		

Date:05/27/05ISR Number: 4678672-6Report Type:Expedited (15-DaCompany Report #2005077020

Age:71 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other OPHTHALMIC MG, 1 IN 3 D), ORAL	900 MG (300	Oedema Peripheral	Consumer	Neurontin (Gabapentin)	PS		
				Valium (Diazepam) Paxil (Paroxetine Hydrochloride) Protonix (Pantoprazole) Fosamax (Alendronate Sodium)	C C C C		

Zetia (Ezetimibe) C
 Requip (Ropinirole Hydrochloride) C
 Vitamins (Vitamins) C
 Mobic (Meloxicam) C
 Calcium (Calcium) C
 Detrol La (Tolterodine L-Tartrate) C

Date:05/27/05ISR Number: 4678677-5Report Type:Expedited (15-DaCompany Report #2005040437

Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Complex Regional Pain Syndrome	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
100 MG (100 MG, 1 IN 1 D), ORAL		Condition Aggravated Drug Ineffective	Professional				
		Gallbladder Operation Incorrect Dose Administered Pharmaceutical Product Complaint		Vitamin B (Vitamin B)	C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4678681-7Report Type:Expedited (15-DaCompany Report #2005004215

Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin			
Other		Completed Suicide		(Gabapentin)	PS		
		Drug Ineffective					

Date:05/27/05ISR Number: 4678682-9Report Type:Expedited (15-DaCompany Report #2005003323

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Adjustment Disorder	Consumer	Neurontin			
Other		Back Pain		(Gabapentin)	PS		ORAL
1800 MG		Convulsion					
(DAILY), ORAL		Drug Ineffective					
		Dyspnoea					
		Fear					
		Headache					
		Hyperhidrosis					
		Marital Problem					
		Pain In Extremity					
		Palpitations					
		Panic Disorder					
		Suicide Attempt					

Date:05/27/05ISR Number: 4679041-5Report Type:Expedited (15-DaCompany Report #2004106658

Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agitation	Consumer	Neurontin			
		Completed Suicide		(Gabapentin)	PS		
		Depression					
		Drug Ineffective					
		Gun Shot Wound					
		Panic Attack					

Outcome	PT
Death	Anxiety
Other	Atherosclerosis
	Brain Death
	Brain Herniation
	Brain Oedema
	Completed Suicide
	Drug Ineffective
	Facial Bones Fracture
	Fear
	Gun Shot Wound
	Injury
	Intraventricular
	Haemorrhage
	Pain
	Pneumocephalus
	Sinus Bradycardia
	Sinus Disorder
	Skull Fractured Base

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Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, 3 IN 1 D), ORAL		Subarachnoid Haemorrhage Subdural Haemorrhage Suicidal Ideation Traumatic Brain Injury	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Nortriptyline (Nortriptyline)	C		
				Nexium (Esomeprazole)	C		
				Allegra (Fexofenadine Hydrochloride)	C		
				Amitriptyline (Amitriptyline)	C		
				Multivitamins W/Minerals (Minerals Nos, Vitamins Nos)	C		
				Vitamin C (Vitamin C)	C		

Date:05/27/05ISR Number: 4679043-9Report Type:Expedited (15-DaCompany Report #2004065552
Age:49 YR Gender:Male I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG, 3 IN 1 D), Other 1 MG (0.5 MG, 2 IN 1 D),		Life-Threatening Hospitalization - Abscess Abscess Drainage Candidiasis Chest Injury Coagulopathy Colon Injury Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		
				Klonopin (Clonazepam)	SS		
				Percocet (Oxycodone Hydrochloride,			

3 TABLETS (3	Gun Shot Wound	Paracetamol)	SS
IN 1 D),	Haemorrhage		
	Haemothorax	Lorcet (Hydrocodone	
	Hypotension	Bitartrate,	
60 MG (10 MG,	Injury	Paracetamol)	SS
1 IN 4 HR),	Intentional Misuse		
	Major Depression	Protonix	
	Overdose	(Pantoprazole)	C
	Pain	Xalatan	
	Pyrexia	(Latanoprost)	C
	Somnolence	Nexium	
	Splenectomy	(Esomeprazole)	C
	Splenic Injury	Gemfibrozil	
	Staphylococcal Infection	(Gemfibrozil)	C
	Suicide Attempt	Pravachol	
		(Pravastatin Sodium)	C
		Flonase (Fluticasone	
		Propionate)	C
		Gaviscon (Sodium	
		Alginate, Sodium	
		Bicarbonate)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4679044-0Report Type:Expedited (15-DaCompany Report #2004058040
Age:41 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	900 MG (300 MG, 3 IN 1 D), ORAL	Areflexia Brain Scan Abnormal Completed Suicide Computerised Tomogram Abnormal Corneal Reflex Decreased Drug Ineffective Ear Haemorrhage Epistaxis Fall Gun Shot Wound Head Injury Heart Rate Increased Hypotension Injury Mouth Haemorrhage Movement Disorder Nervous System Disorder Oxygen Saturation Decreased Pupil Fixed Respiration Abnormal Skin Discolouration Traumatic Brain Injury	Consumer	Neurontin (Gabapentin) Remeron (Mirtazapine)	PS C		

Date:05/27/05ISR Number: 4679045-2Report Type:Expedited (15-DaCompany Report #2004117310
Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	SEE IMAGE	Completed Suicide Drug Ineffective Loss Of Consciousness Multiple Drug Overdose Accidental	Consumer	Neurontin (Gabapentin) Ibuprofen (Ibuprofen) Amitriptyline Hydrochloride	PS SS		

(Amitriptyline Hydrochloride)	SS
Nortriptyline Hydrochloride (Nortriptyline Hydrochloride)	SS
Venlafaxine (Venlafaxine)	SS
Oxazepam (Oxazepam)	C
Benzodiazepine Derivatives (Benzodiazepine Derivatives)	C
Opioids (Opioids)	C

Date:05/27/05ISR Number: 4679048-8Report Type:Expedited (15-DaCompany Report #2005077748
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Asthenia	Consumer	Neurontin (Gabapentin)	PS		

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Freedom Of Information (FOI) Report

Date:05/27/05ISR Number: 4679286-4Report Type:Expedited (15-DaCompany Report #2004074993

Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Overdose	Consumer	Neurontin			
Hospitalization -		Completed Suicide		(Gabapentin)	PS		
Initial or Prolonged		Condition Aggravated		Celebrex (Celecoxib)	SS		
Other		Depression		Paxil (Paroxetine			
		Drug Abuser		Hydrochloride)	SS		
		Drug Ineffective		Amitriptyline With			
		Injury		Perphenazine			
		Intentional Misuse		(Amitriptyline			
		Mental Status Changes		Hydrochloride,			
		Metabolic Acidosis		Perphenazine)	SS		
		Overdose		Oxycontin (Oxycodone			
		Respiratory Distress		Hydrochloride)	SS		
		Respiratory Failure		Paracetamol			
		Suicide Attempt		(Paracetamol)	SS		
				Nexium			
				(Esomeprazole)	C		
				Baclofen (Baclofen)	C		
				Insulin (Insulin)	C		
				Duragesic (Fentanyl)	C		
				Glucophage			
				(Metformin			
				Hydrochloride)	C		

Date:05/31/05ISR Number: 4677955-3Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20050507147

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Ibuprofen	PS		
		Multiple Drug Overdose		Nortriptyline			
		Accidental		Hydrochloride	SS		
				Amitriptyline			
				Hydrochloride	SS		
				Venlafaxine	SS		
				Neurontin	SS		
				Neurontin	SS		
				Neurontin	SS		

800mg to 1800

mg daily

Oxazepam C
Benzodiazepine C
Derivatives C
Opioids C

Date:05/31/05ISR Number: 4678802-6Report Type:Expedited (15-DaCompany Report #2004043537
Age:55 YR Gender:Male I/FU:F

Outcome PT
Death Anxiety
Hospitalization - Cognitive Disorder
Initial or Prolonged Completed Suicide
Other Depressed Level Of
Consciousness
Fatigue
Feeling Jittery
Head Discomfort
Inadequate Analgesia
Injury

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Insomnia Intentional Misuse Libido Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 2 IN 1 D), ORAL		Major Depression Neck Pain Pain In Extremity Paraesthesia					
INTRAVENOUS	INTRAVENOUS	Polysubstance Abuse Respiratory Distress		Midazolam (Midazolam)	SS		
ORAL		Restlessness Sleep Disorder Sluggishness Somnolence		Tramadol Hydrochloride (Tramadol Hydrochloride)	SS		ORAL
INTRAVENOUS	INTRAVENOUS	Suicidal Ideation		Citalopram (Citalopram) Pancuronium (Pancuronium)	SS SS		
				All Other Therapeutic Products (All Other Therapeutic Products)	SS		
				Zolpidem Tartrate (Zolpidem Tartrate)	C		
				Ibuprofen (Ibuprofen)	C		
				Clonazepam (Clonazepam)	C		
				Mirtazapine (Mirtazapine)	C		

Date:05/31/05ISR Number: 4678820-8Report Type:Expedited (15-DaCompany Report #2005067549

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Anger	Consumer	Xanax Tablet			

Initial or Prolonged 4 MG (0.5 MG, Other 3 IN 1 D),	Anxiety Disorder	Health	(Alprazolam)	PS	ORAL
ORAL	Diet Refusal	Professional			
	Disturbance In Social Behaviour		Neurontin		
(3 IN 1 D),	Drug Dependence		(Gabapentin)	SS	ORAL
ORAL	Drug Withdrawal Syndrome				
ORAL	Intentional Misuse		Geodon (Ziprasidone)	SS	
	Overdose		Nyquil		
	Phobia		(Dextromethorphan Hydrobromide, Doxylamine Succinate, Ephedrine	SS	
	Psychotic Disorder		All Other Non-Therapeutic Products (All Other Non-Therapeutic Products)	SS	
	Self-Medication		Adderall (Amfetamine Aspartate, Amfetamine Sulfate, Dexamfetamine Saccharate,	C	
	Treatment Noncompliance		Prozac (Fluoxetine Hydrochloride)	C	

Freedom Of Information (FOI) Report

Ambien (Zolpidem
Tartrate) C
Soma (Carisoprodol) C

Date:05/31/05ISR Number: 4678840-3Report Type:Expedited (15-DaCompany Report #2004077855
Age:14 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour	Consumer	Neurontin			
300 MG, ORAL		Affective Disorder		(Gabapentin)	PS		ORAL
		Aggression		Levothyroid			
		Alcoholism		(Levothyroxine			
		Anger		Sodium)	C		
		Coma		Clindamycin			
		Concussion		(Clindamycin)	C		
		Depression		Seroquel (Quetiapine			
		Developmental Delay		Fumarate)	C		
		Drug Abuser		Lithium (Lithium)	C		
		Dysarthria		Multivitamins			
		Gait Disturbance		(Multivitamins)	C		
		Hypothyroidism		Paxil (Paroxetine			
		Injury		Hydrochloride)	C		
		Learning Disorder		Clozaril (Clozapine)	C		
		Loss Of Consciousness		Ativan (Lorazepam)	C		
		Pain		Colace (Docusate			
		Personality Change		Sodium)	C		
		Psychotic Disorder		Provigil (Modafinil)	C		
		Schizoaffective Disorder		Topamax (Topiramate)	C		
		Suicide Attempt		Naltrexone			
				(Naltrexone)	C		
				Depo-Provera 150			
				(Medroxyprogesterone			
				Acetate)	C		
				Albuterol			
				(Salbutamol)	C		
				All Other			
				Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 600 MG (300 Disability MG, 2 IN 1 D) Other	Alcohol Use Bipolar Disorder Condition Aggravated Gun Shot Wound Injury Intentional Misuse Mobility Decreased Pain Scar Suicide Attempt Swelling Face	Consumer	Neurontin (Gabapentin)	PS		

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Date:05/31/05ISR Number: 4678873-7Report Type:Expedited (15-DaCompany Report #2005040889

Age:55 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Pressure Increased	Health	Celebrex (Celecoxib)	PS		ORAL
200 MG (200		Cardiac Murmur	Professional				
MG, 1 IN 1		Cardiac Valve Disease					
D), ORAL		Convulsion		Neurontin			
		Dysstasia		(Gabapentin)	SS		ORAL
900 MG (300		Fluid Retention					
MG, 3 IN 1		Keratorhexis					
D), ORAL		Muscle Spasms		Gabaentin			
		Musculoskeletal		(Gabapentin)	SS		ORAL
(300 MG),		Discomfort					
ORAL		Oedema Peripheral		Bextra (Valdecoxib)	SS		ORAL
ORAL		Pharmaceutical Product		Nexium			
		Complaint		(Esomeprazole)	C		
		Psoriatic Arthropathy		Elavil			
		Rheumatoid Arthritis		(Amitriptyline			
		Stool Analysis Abnormal		Hydrochloride)	C		
		Tremor		Tylox (Oxycodone			
		Urinary Incontinence		Hydrochloride,			
		Weight Decreased		Paracetamol)	C		
				Lasix (Furosemide0	C		
				Klonopin			
				(Clonazepam)	C		
				Voltaren (D9clofenac			
				Sodium)	C		

Date:05/31/05ISR Number: 4678931-7Report Type:Expedited (15-DaCompany Report #2005016042

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Depression	Consumer	Neurontin			

Initial or Prolonged Gun Shot Wound (Gabapentin) PS
 Other Headache
 Paraparesis
 Subdural Haematoma
 Suicide Attempt

Date:05/31/05ISR Number: 4678932-9Report Type:Expedited (15-DaCompany Report #2004106809
 Age:24 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Amnesia	Consumer	Neurontin			
		Atherosclerosis		(Gabapentin)	PS		
		Brain Oedema		Morphine (Morphine)	SS		
3 IN 1 D		Completed Suicide		Duragesic (Fentanyl)	SS		
TRANSDERMAL	175 MG	(1 IN					
		Euphoric Mood					
1 D),		Injury					
TRANSDERMAL		Intentional Misuse		Heroin (Diamorphine)	SS		
		Laryngeal Disorder		Citalopram			
		Loss Of Consciousness		(Citalopram)	SS		
		Mouth Haemorrhage		Propoxyphene			
		Pulmonary Congestion		(Dextropropoxyphene)	SS		
		Pulmonary Oedema					
		Respiratory Arrest					
		Ventricular Hypertrophy					
		Vomiting					

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Date:05/31/05ISR Number: 4678954-8Report Type:Expedited (15-DaCompany Report #2005003387
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged Other		Cervical Vertebral Fracture Coma Conjunctivitis Corneal Ulcer Difficult To Wean From Ventilator Drug Ineffective Dysphagia Gun Shot Wound Pneumonia Haemophilus Self Injurious Behaviour Subdural Haematoma Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:05/31/05ISR Number: 4678959-7Report Type:Expedited (15-DaCompany Report #2005076965
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Gastric Bypass	Health Professional	Neurontin (Gabapentin)	PS		

Date:05/31/05ISR Number: 4679256-6Report Type:Expedited (15-DaCompany Report #FRWYE689924MAY05
Age:75 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Bone Marrow Toxicity Condition Aggravated Marrow Hyperplasia	Health Professional Other	Effexor (Venlafaxine Hydrochloride, Unspec, 0)	PS		ORAL
150 MG 1X PER		Metamyelocyte Count					
1 DAY ORAL	8 DAY	Increased Neutropenia		Calciparin (Heparin Calcium, ,0)	SS		
SUBCUTANEOUS	SC			Ciflox			

500 MG 2X PER

(Ciprofloxacin, ,0) SS

ORAL

1 DAY ORAL 15 DAY

Gentamicin
(Gentamicin Sulfate,
,0) SS

INTRAVENOUS 200 MG 1X PER

1 DAY IV 15 DAY

Imovane (Zopiclone,
, 0) SS

ORAL

ORAL

Neurontin
(Gabapentin, ,0) SS

ORAL

ORAL

Xatral (Alfuzosin) C

Date:05/31/05ISR Number: 4679269-4Report Type:Direct
Age:59 YR Gender:Male I/FU:I

Company Report #CTU 249930

Outcome PT
Death Collapse Of Lung
Hepatic Cirrhosis
Hepatic Function Abnormal
Hip Fracture
Liver Disorder
Starvation

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Swelling

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
2 PATCHES	LASTS 48 HRS		Fentanyl Patches 100	PS		OTHER
OTHER			Neurontin	SS		ORAL
2 TAKEN BY						
MOUTH	ORAL					

Date:05/31/05ISR Number: 4679687-4Report Type:Expedited (15-DaCompany Report #2004070397
 Age:42 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Disability 1200 MG (300 MG, 4 IN 1 Other D), ORAL		Agitation Alcoholism Anger Anxiety Bipolar Disorder Depression Disability Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		ORAL
2 MG 90.5 MG, 4 IN 1 D), ORAL		Drug Level Decreased Drug Tolerance Feeling Abnormal Intentional Misuse Pain Sexual Dysfunction Somnolence Stress At Work Suicidal Ideation Suicide Attempt		Clonazepam (Clonazepam) Prozac (Fluoxetine Hydrochloride) Eskalith (Lithium Carbonate)	SS C C		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Acute Respiratory	Consumer	Neurontin			
Other		Distress Syndrome		(Gabapentin)	PS		
		Anoxic Encephalopathy		Effexor-Xr			
		Anxiety		(Venlafaxine			
		Apnoea		Hydrochloride)	C		
		Brain Oedema		Risperdal			
		Cardiac Arrest		(Risperidone)	C		
		Completed Suicide					
		Depression					
		Injury Asphyxiation					
		Pain					
		Pneumomediastinum					
		Pneumonia Aspiration					
		Pneumonitis					
		Respiratory Arrest					
		Respiratory Failure					
		Soft Tissue Injury					
		Suicidal Ideation					
		Victim Of Crime					
		Vomiting					

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Date:05/31/05ISR Number: 4679690-4Report Type:Expedited (15-DaCompany Report #2005060540
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 20 MG (10 MG, Initial or Prolonged BID INTERVAL: EVERY DAY),ORAL		Abasia Cerebrovascular Accident Confusional State Drug Ineffective	Consumer	Bextra (Valdecoxib)	PS		ORAL
900 MG (300 MG, TID INTERVAL: EVERY DAY), ORAL		Palpitations Tubal Ligation		Neurontin (Gabapentin)	SS		ORAL
				Xanax (Alprazolam) Celexa (Citalopram Hydrobromide) Prevacid (Lansoprazole) Flexeril (Cyclobenzaprine Hydrochloride)	C C C C		

Date:05/31/05ISR Number: 4679709-0Report Type:Expedited (15-DaCompany Report #2004093246
Age:70 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other OPHTHALMIC	ORAL	Completed Suicide Gastrointestinal Disorder	Consumer	Neurontin (Gabapentin)	PS		
		Gun Shot Wound Insomnia Polytraumatism		Levaquin (Levaquin) Metronidazole (Metronidazole) Zestril (Lisinopril)	C C C		

Casodex	
(Bicalutamide)	C
Roxicodone	
(Oxycodone	
Hydrochloride)	C
Protonix	
(Pantoprazole)	C
Aciphex (Rabeprazole	
Sodium)	C
Oxycontin (Oxycodone	
Hydrochloride)	C
K-Dur (Potassium	
Chloride)	C
Antibiotics	C
All Other	
Therapeutic Products	C

Date:05/31/05ISR Number: 4679710-7Report Type:Expedited (15-DaCompany Report #2004106702

Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Consumer	Neurontin			
Other		Drug Ineffective		(Gabapentin)	PS		
		Poisoning Deliberate					

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Date:05/31/05ISR Number: 4679923-4Report Type:Expedited (15-DaCompany Report #2005042717

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Disease Recurrence	Health	Neurontin			
		Lung Neoplasm Malignant	Professional	(Gabapentin)	PS		
		Neoplasm Malignant					

Date:05/31/05ISR Number: 4679924-6Report Type:Expedited (15-DaCompany Report #2004117393

Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Anger	Consumer	Neurontin			
Initial or Prolonged		Anxiety		(Gabapentin)	PS		
900 MG (300							
Disability		Deformity					
MG, 3 IN 1							
Other		Drug Ineffective					
D),							
		Eye Injury		Trazodone			
		Face Injury		(Trazodone)	C		
		Facial Bones Fracture		Mirtazapine			
		Gun Shot Wound		(Mirtazapine)	C		
		Head Injury		Naltrexone			
		Ill-Defined Disorder		(Naltrexone)	C		
		Insomnia					
		Intentional Misuse					
		Intentional Self-Injury					
		Irritability					
		Migraine					
		Overdose					
		Pain					
		Soft Tissue Injury					
		Stress					
		Suicidal Ideation					
		Suicide Attempt					
		Unevaluabe Event					

Date:05/31/05ISR Number: 4679926-XReport Type:Expedited (15-DaCompany Report #2004058034

Age:50 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Adjustment Disorder
Initial or Prolonged	Anger
Disability	Anxiety
Other	Back Pain
	Bipolar Disorder
	Burning Sensation
	Chest Pain
	Confusional State
	Drug Ineffective
	Grandiosity
	Hallucination, Auditory
	Hyperhidrosis
	Intentional Misuse
	Intentional Self-Injury
	Limb Injury
	Nephritis Interstitial
	Off Label Use
	Overdose
	Pain
	Pressure Of Speech
	Psychomotor Hyperactivity

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Dose	Duration	Restlessness Stress Suicide Attempt	Report Source	Product	Role	Manufacturer	Route
1800 MG,		Tinea Versicolour Urethritis	Consumer	Neurontin (Gabapentin)	PS		
2400 MG (600 MG, 4 IN 1 D), ORAL				Ibuprofen (Ibuprofen)	SS		ORAL
				Olanzapine (Olanzapine)	C		
				Temazepam (Temazepam)	C		
				Clonazepam (Clonazepam)	C		
				Bupropion (Bupropion)	C		
				Depakote (Valproate Semisodium)	C		
				Effexor-Xr (Venlafaxine Hydrochloride)	C		
				Zoloft (Sertraline)	C		
				Serzone (Nefazodone Hydrochloride)	C		
				Ambien (Zolpidem Tartrate)	C		
				Gabitril (Tiagabine Hydrochloride)	C		
				Risperidone (Risperidone)	C		
				Griseofulvin (Griseofulvin)	C		
				Clotrimazole (Clotrimazole)	C		
				Vytone (Diiodohydroxyquinol ine, Hydrocortisone)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Condition Aggravated	Consumer	Neurontin			
Other		Depression	Health	(Gabapentin)	PS		ORAL
ORAL		Diarrhoea	Professional	Prozac (Fluoxetine			
		Disease Progression		Hydrochloride)	C		
		Hyperreflexia		Baclofen (Baclofen)	C		
		Hypertension		Prilosec			
		Multiple Sclerosis		(Omeprazole)	C		
		Nausea		Nitrostat			
		Panic Attack		(Nitroglycerin)	C		
		Photophobia		Carbamazepine			
		Weight Decreased		(Carbamazepine)	C		
				Adderall (Amfetamine			
				Aspartate,			
				Amfetamine Sulfate,			
				Dexamfetamine			
				Saccharate,	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:05/31/05ISR Number: 4680868-4Report Type:Direct
 Age:21 YR Gender:Female I/FU:I

Company Report #CTU 250028

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Neurontin	PS		
300 MG 2 TID		Therapeutic Response Unexpected With Drug Substitution					

Date:05/31/05ISR Number: 4680902-1Report Type:Direct
 Age:83 YR Gender:Female I/FU:I

Company Report #CTU 250022

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Dysarthria Somnolence		Gabapentin Caps G5027; 300mg Greenstone	PS	Greenstone	
TAKE 1							
CAPSULE EVERY							
8 HOURS							

Date:06/01/05ISR Number: 4680899-4Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 250119

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Neurontin (Generic)	PS		ORAL
600 MG ORAL		Therapeutic Response Unexpected With Drug Substitution					
BID							

Date:06/01/05ISR Number: 4680906-9Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 250046

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Condition Aggravated		Gabapentin 300mg	PS	Greenstone	
300MG EVERY 6 HOURS		Neuropathic Pain					
		Pharmaceutical Product Complaint					

Date:06/01/05ISR Number: 4680916-1Report Type:Direct Company Report #CTU 250053
 Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Condition Aggravated		Gabapentin 600mg	PS	Greenstone	
TAKE 1 TABLET FOUR TIMES A DAY		Neuropathy Peripheral					
		Pharmaceutical Product Complaint					

Date:06/01/05ISR Number: 4681095-7Report Type:Expedited (15-DaCompany Report #2005044232
 Age:56 YR Gender:Female I/FU:F

Outcome	PT
Other	Alopecia
	Antipsychotic Drug Level Above Therapeutic
	Bladder Prolapse
	Breath Odour
	Drug Ineffective
	Dysgeusia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Complaint	Report Source	Product	Role	Manufacturer	Route
800 MG (1 D), ORAL		Gangrene Hypoaesthesia Inadequate Analgesia Nausea Panic Attack Pharmaceutical Product	Consumer	Neurontin (Gabapentin)	PS		ORAL
		Complaint Tooth Discolouration Tooth Loss		Lithium (Lithium) Cyclobenzaprine (Cyclobenzaprine) Thyroid (Thyroid) All Other Therapeutic Product (All Other Therapeutic Products) Clonazepam (Clonazepam) Avandia (Rosiglitazone Maleate) Naproxen (Naproxen) Zocor (Simvastatin)	SS C C C C C C C C C		

Date:06/02/05ISR Number: 4679968-4Report Type:Expedited (15-DaCompany Report #PHFR2005GB01965

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 40 mg, QD		Blood Creatinine Increased		Baclofen	PS	Novartis Sector: Pharma	ORAL
Initial or Prolonged 36mg/day		Drug Level Increased		Tizanidine	SS		ORAL
Other UNKNOWN	400 mg, BID	Epilepsy		Gabapentin	SS		
2.5mg/day		Glasgow Coma Scale		Bendrofluazide	C		ORAL
		Abnormal Hallucination Loss Of Consciousness Overdose Renal Failure Acute					

Urinary Tract Infection

Date:06/02/05ISR Number: 4681565-1Report Type:Expedited (15-DaCompany Report #2005077695

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Back Pain	Consumer	Neurontin			
Initial or Prolonged	Body Height Decreased		(Gabapentin)	PS		
100 MG (100						
Other	Constipation					
MG 1 IN 1D)						
	Delayed Recovery From		Bextra(Valdecoxib)	SS		
	Anaesthesia		Motrin (Ibuprofen)	SS		
	Dizziness		Fentanyl (Fentanyl)	SS		
TRANSDERMAL	(25 MCG)					
	Drug Ineffective					
TRANSDERMAL						
	Emotional Disorder		Vioxx (Rofecoxib)	SS		
	Feeling Abnormal		Advil (Ibuprofen)	SS		
	Gastritis		Aleve (Naproxen			
	Impaired Driving Ability		Sodium)	C		
	Muscle Spasms		Levoxyl			
	Nausea		(Levothyroxine			
	Paralysis		Sodium)	C		
	Pruritus					
	Somnolence					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/02/05ISR Number: 4681585-7Report Type:Expedited (15-DaCompany Report #2004100388
 Age:50 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG 3 IN 1 D)		Arteriovenous Malformation	Consumer Health	Neurontin (Gabapentin)	PS		
		Aseptic Necrosis Bone	Professional				
		Cerebrovascular Spasm Osteoporosis		Prednisone Tablet (Prednisone)	SS		

Date:06/02/05ISR Number: 4681614-0Report Type:Expedited (15-DaCompany Report #2005042034
 Age:69 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - 200 MG (100 Initial or Prolonged MG, 2 IN 1 Other D), ORAL		Abdominal Tenderness Anuria	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Blood Creatinine Increased	Professional				
		Blood Potassium Increased		Rapamune (Sirolimus)	C		
		Brain Death		Minoxidil			
		Cardiac Arrest		(Minoxidil)	C		
		Clostridial Infection		Humalog (Insulin			
		Dehydration		Lispro)	C		
		Failure To Thrive		Lantus (Insulin			
		Gastrointestinal		Glargine)	C		
		Haemorrhage		Neoral (Ciclosporin)	C		
		Haematemesis		Prednisone			
		Haematochezia		(Prednisone)	C		
		Haemoglobin Decreased		Epoetin Alfa			
		Hepatic Failure		(Epoetin Alfa)	C		
		International Normalised Ratio Increased		Metoprolol			
		Metabolic Acidosis		(Metoprolol)	C		
		Multi-Organ Failure		Percocet (Oxycodone			
		Pneumonia		Hydrochloride, Paracetamol)	C		
		Renal Failure		Pantoprazole			
		Respiratory Arrest		(Pantoprazole)	C		
		Sepsis		Quinine (Quinine)	C		

Shock Haemorrhagic
Urine Output Decreased
Vascular Resistance
Systemic Decreased

Dynacirc
(Isradipine) C
Aldactone
(Spironolactone) C
Furosemide
(Furosemide) C
Nystatin (Nystatin) C
Prevacid
(Lansoprazole) C
Maalox (Aluminium
Hydroxide Gel,
Magnesium Hydroxide) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C

Date:06/02/05ISR Number: 4682467-7Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 250148

Outcome PT
Other Condition Aggravated
Vaginal Candidiasis

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Freedom Of Information (FOI) Report

Vulvovaginal Discomfort

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG CAP			Gabapentin	PS		

Date:06/03/05ISR Number: 4685187-8Report Type:Expedited (15-DaCompany Report #2005055610
 Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	900 MG (1D), ORAL	Drug Ineffective Epilepsy	Foreign Health Professional Company Representative	Gabapentin (Gabapentin) Valproic Acid (Valproic Acid) Venlafaxine (Venlafaxine) Omeprazol (Omeprazole) Olanzapine (Olanzapine)	PS C C C C		ORAL

Date:06/06/05ISR Number: 4682446-XReport Type:Expedited (15-DaCompany Report #US-ROCHE-380104
 Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged INTRAVENOUS	1 DAY	Anaemia Convulsion		Granisetron Hydrochloride	PS	Roche	
SUBCUTANEOUS	87 DAY	Disease Progression		Darbepoetin Alfa	SS		
UNKNOWN		Gastrointestinal		Dronabinol	SS		
UNKNOWN	31 DAY	Haemorrhage		Oxycodone	SS		
UNKNOWN		Haemolytic Anaemia		Gabapentin	SS		
		Liver Transplant		Donepezil			

UNKNOWN	Rejection	Hydrochloride	SS
UNKNOWN	Mental Status Changes	Zolpidem Tartrate	SS
55 DAY	Metabolic Encephalopathy	Morphine Sulfate	C
	Nausea	Chlorpromazine	C
	Pancytopenia	Oxyir	C
	Vomiting	Tacrolimus	C
		Pantoprazole	C
		Prandin	C
		Sertraline	
		Hydrochloride	C
		Glipizide	C
		Nifedipine	C
		Magnesium	C
		Levofloxacin	C
8 DAY		Pentamidine	C
		Pericolace	C
		Bisacodyl	C
		Acetylsalicylic Acid	C
		Frusemide	C
INTRAVENOUS	1 DAY	Sargamostim	C
SUBCUTANEOUS	6 DAY	Dexamethasone	C
INTRAVENOUS	1 DAY	Doxorubicin	C
		Cisplatin	C
		Senna Fruit	C
		Propranolol	C
		Multivitamins	C
		Insulin	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Spironolactone C
 Megestrol Acetate C

Date:06/06/05ISR Number: 4683283-2Report Type:Direct
 Age:36 YR Gender:Male I/FU:I

Company Report #CTU 250375

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Abuser		Gabapentin 400mg			
		Drug Dependence		Teva	PS	Teva	ORAL
400 - 600 MG		Drug Effect Decreased					
4-5 TIMES/WK		Euphoric Mood					
ORAL							

Date:06/06/05ISR Number: 4684909-XReport Type:Direct
 Age:58 YR Gender:Female I/FU:I

Company Report #CTU 250484

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
		Rash		Neurontin 300 Mg Tid	PS		ORAL
300 MG PO TID		Therapeutic Response Unexpected With Drug Substitution					

Date:06/06/05ISR Number: 4684917-9Report Type:Direct
 Age:63 YR Gender:Female I/FU:I

Company Report #CTU 250476

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Amnesia		Neurontin 400 Mg Cap	PS		
1 A DAY		Suicide Attempt					
Hospitalization - Initial or Prolonged							

Date:06/06/05ISR Number: 4685797-8Report Type:Direct
 Age:49 YR Gender:Female I/FU:I

Company Report #CTU 250448

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Neurontin	PS		
600 MG 2 TID		Pharmaceutical Product Complaint					

Date:06/06/05ISR Number: 4685959-XReport Type:Expedited (15-DaCompany Report #050526-0000650
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Fatigue	Consumer	Desoxyn	PS		ORAL
30 MG; Q4H;		Loss Of Consciousness					
PO	18 MON	Murder		Xanax	SS		
2 MG		Road Traffic Accident		Oxycontin	SS		
80 MG; Q4H				Duragesic Patches	SS		
100 MCG				Prilosec	SS		
40 MG; QD				Neurontin	SS		
300 MG; Q4H							

Date:06/06/05ISR Number: 4686297-1Report Type:Expedited (15-DaCompany Report #2002054135
Age: Gender: I/FU:F

Outcome	PT
Other	Drug Exposure During Pregnancy

Freedom Of Information (FOI) Report

Solitary Kidney

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Literature Health Professional	Neurontin (Gabapentin) Phenobarbital (Penobarbital)	PS C		

Date:06/06/05ISR Number: 4686302-2Report Type:Expedited (15-DaCompany Report #2005081403
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Amniotic Fluid Volume Decreased Drug Exposure During Pregnancy	Literature Health Professional	Neurontin (Gabapentin)	PS		

Date:06/06/05ISR Number: 4686304-6Report Type:Expedited (15-DaCompany Report #2005081416
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Caesarean Section Drug Exposure During Pregnancy	Literature Health Professional	Neurontin (Gabapentin)	PS		

Date:06/06/05ISR Number: 4686306-XReport Type:Expedited (15-DaCompany Report #2005081434
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Caesarean Section Drug Exposure During Pregnancy	Literature Health Professional	Neurontin (Gabapentin)	PS		

Date:06/06/05ISR Number: 4686314-9Report Type:Expedited (15-DaCompany Report #2005081451
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Caesarean Section Drug Exposure During Pregnancy	Literature Health Professional	Neurontin (Gabapentin)	PS		

Date:06/06/05ISR Number: 4686328-9Report Type:Expedited (15-DaCompany Report #2005078616
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Back Pain Drug Withdrawal Syndrome	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Intervertebral Disc Protrusion Muscle Spasms Pain Paraesthesia Rotator Cuff Syndrome Treatment Noncompliance Weight Increased		Skelaxin (Metaxalone) Oxycontin (Oxycodone)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/06/05ISR Number: 4686385-XReport Type:Expedited (15-DaCompany Report #2005079866

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

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Duration Anaemia Bacteraemia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other SUBCUTANEOUS	Neutropenia Renal Failure SUBCUTANEOUS	Professional	Calciparine "Difrex" (Heparin Calcium)	SS		
1000 MG (500 MG, 2 IN 1 D) ORAL	Spinal Myelogram Abnormal Urinary Tract Infection White Blood Cell Count Decreased		Ciflox (Ciprofloxacin)	SS		ORAL
INTRAVENOUS 1 D)	200 MG (1 IN		Gentamicin (Gentamicin)	SS		
INTRAVENOUS ORAL			Imovane (Zopiclone)	SS		ORAL
150 MG (1 IN 1 D) ORAL			Effexor (Venlafaxine Hydrochloride)	SS		ORAL
			Xatral (Alfuzosin) Stablon (Tianeptine) Tavanic (Levofloxacin)	C C C		

Date:06/06/05ISR Number: 4686973-0Report Type:Expedited (15-DaCompany Report #2005063347

Age:62 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Death Hospitalization - (DAILY), ORAL	Duration Circulatory Collapse Feeling Abnormal	Foreign Health	Neurontin (Gabapentin)	PS		ORAL

Initial or Prolonged	Hypokinesia	Professional	Zoloft (Sertraline)	C
	Pulmonary Oedema	Company	All Other	
		Representative	Therapeutic Products	
			(All Other	
			Therapeutic	
			Products)	C
			Cartia	
			(Acetylsalicylic	
			Acid)	C
			Endep (Amitriptyline	
			Hydrochloride)	C
			Furosemide	
			(Furosemide)	C
			Gliclazide	
			9gliclazide)	C
			Karvea (Irbesartan)	C
			Lipitor	
			(Atorvastatin)	C
			Metformin(Metformin)	C
			Norvasc (Amlodipine)	C

Date:06/06/05ISR Number: 4687183-3Report Type:Expedited (15-DaCompany Report #2005016042
Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Aggression
Initial or Prolonged	Convulsion
Other	Depression
	Drug Ineffective

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Gun Shot Wound Headache Hemiparesis	Report Source	Product	Role	Manufacturer	Route
		Paraparesis Simple Partial Seizures Subdural Haematoma Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:06/07/05ISR Number: 4684194-9Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0382647A
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 20MG Per day		Chills		Deroxat	PS	Glaxosmithkline	ORAL
Initial or Prolonged 300MG Per day		Hypercalcaemia		Zyloric	SS	Glaxosmithkline	ORAL
100MG Per day		Leukocyturia		Neoral	SS		ORAL
INTRAVENOUS dose	60MG Single 2 HR	Pyelonephritis Pyrexia		Aredia	SS		
INTRAVENOUS day	4000IU per day			Neorecormon	SS		
40MG Per day				Mopral	SS	Glaxosmithkline	ORAL
125MCG Per day				Levothyrox	SS	Glaxosmithkline	ORAL
300MG Twice per day				Neurontin	SS		ORAL
30DROP per day				Rivotril	SS		ORAL
10MG Per day				Stilnox	SS		ORAL

SUBCUTANEOUS	Calciparine	SS	
250MG Per day	Cellcept	SS	ORAL
13 DAY	Phosphoneuros	SS	ORAL
5MG Per day	Amlor	SS	ORAL
4MG Unknown	Kenzen	SS	ORAL

Date:06/07/05ISR Number: 4685784-XReport Type:Direct Company Report #CTU 250630
 Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State		Gabapentin 300 Mg	PS		
2 CAP TID		Incoherent Nausea Pharmaceutical Product Complaint Tinnitus					

Date:06/07/05ISR Number: 4685835-2Report Type:Direct Company Report #CTU 250581
 Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Rash Pruritic		Gabapentin	PS		ORAL
300 MG PO QHS							

Date:06/07/05ISR Number: 4686199-0Report Type:Expedited (15-DaCompany Report #2005078612
 Age: Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Anxiety
Initial or Prolonged	Crying
	Drug Effect Decreased

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1-2 MG (1 MG, ONCE TO TWICE A DAY), ORAL		Drug Ineffective Drug Interaction Limb Operation Phantom Pain Shoulder Operation Suicidal Ideation Weight Decreased	Consumer	Alprazolam (Alprazolam)	PS		ORAL
ORAL				Neurontin (Gabapentin)	SS		ORAL
				Synthroid (Levothyroxine Sodium)	C		
				Multivitamins (Multivitamins)	C		

Date:06/07/05ISR Number: 4686461-1Report Type:Expedited (15-DaCompany Report #2005074293
Age:45 YR Gender:Female I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
1800 MG (600 MG, 3 IN 1 D), ORAL (800 MG), ORAL		Brain Damage Condition Aggravated Drug Effect Decreased Drug Interaction Headache Insomnia	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
ORAL				Motrin (Ibuprofen)	SS		ORAL
		Intervertebral Disc Protrusion Nerve Compression		Hydrocodone (Hydrocodone)	SS		
				Mobic (Meloxicam)	SS		ORAL
		Neuropathy Pain Pain In Extremity Shoulder Pain Treatment Noncompliance		Ultram (Tramadol Hydrochloride)	C		

Date:06/07/05ISR Number: 4686543-4Report Type:Expedited (15-DaCompany Report #2005078325
Age:79 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin			
Other		Drug Ineffective Suicidal Ideation		(Gabapentin)	PS		

Date:06/07/05ISR Number: 4686593-8Report Type:Expedited (15-DaCompany Report #2005081014
Age:60 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abdominal Discomfort Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		
(200 MG,)				Zantac 75 (Ranitidine Hydrochloride)	SS		
75 MG (75 MG,				Tylenol (Paracetamol)	C		
1 IN 1 D)				Colace (Docusate Sodium)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/07/05ISR Number: 4686602-6Report Type:Expedited (15-DaCompany Report #2004117310
 Age:44 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	800 MG (1 D)	Completed Suicide Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		
		Loss Of Consciousness Multiple Drug Overdose Accidental		Ibuprofen (Ibuprofen)	SS		
				Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		
				Nortriptyline Hydrochloride (Nortriptyline Hydrochloride)	SS		
				Venlafaxine (Venlafaxine)	SS		
				Oxazepam (Oxazepam)	C		
				Benzodiazepine Derivatives (Benzodiazepine Derivatives)	C		
				Opioids (Opioids)	C		

Date:06/08/05ISR Number: 4685398-1Report Type:Expedited (15-DaCompany Report #FR-GLAXOSMITHKLINE-B0382567A
 Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	10MG Per day	Abortion Induced		Deroxat	PS	Glaxosmithkline	ORAL
	6UNIT per day	Drug Exposure During Pregnancy		Diantalvic	SS		ORAL
	.5MG per day			Rivotril	SS		ORAL
	900MG per day			Xanax	SS		ORAL
				Neurontin	SS		ORAL

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Death	Abortion Induced		Deroxat	PS	Glaxosmithkline	
10MG Per day						
Congenital Anomaly	Congenital Nose		Diantalvic	SS		
6UNIT per day						
	Malformation		Rivotril	SS		
	Drug Exposure During		Xanax	SS		
.5MG per day						
900MG per day	Pregnancy		Neurontin	SS		
	Facial Dysmorphism					
	Micrognathia					
	Multiple Congenital					
	Abnormalities					
	Oesophageal Atresia					
	Retrognathia					

Date:06/08/05ISR Number: 4685751-6Report Type:Expedited (15-DaCompany Report #KII-2005-0016764

Age:47 YR Gender:Female I/FU:I

Outcome	PT
Hospitalization -	Blood Pressure Systolic
Initial or Prolonged	Increased
Other	Bowel Sounds Abnormal
	Confusional State

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Depressed Level Of Consciousness Dysarthria				
ORAL		Electrocardiogram Qrs Complex Prolonged Electrocardiogram Qt Corrected Interval	Study Health Professional Other			ORAL
ORAL		Prolonged Multiple Drug Overdose Nystagmus Perseveration	Acetaminophen W/Oxycodone (Paracetamol, Oxycodone Hydrochloride)	SS		ORAL
ORAL		Respiratory Rate Decreased	Gabapentin (Gabapentin)	SS		ORAL
ORAL			Amfetamine (Amfetamine)	SS		ORAL
ORAL			Selective Estrogen Receptor Modulator	SS		ORAL
			Remeron (Mirtazapine) Doxepin (Doxepin)	C C		

Date:06/08/05ISR Number: 4686754-8Report Type:Expedited (15-DaCompany Report #2005080588
Age:68 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 MG, 3 IN 1 D), ORAL		Cholestasis Cytolytic Hepatitis Pain	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
				Triflucan (Fluconazole) Paracetamol (Paracetamol)	SS C		

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Life-Threatening	Arthritis	Foreign	Neurontin			
Hospitalization - 800 MG (400	Asthenia	Consumer	(Gabapentin)	PS		ORAL
Initial or Prolonged MG, 2 IN 1	Conjunctival Hyperaemia					
Other D), ORAL	Corneal Disorder					
	Cough		Atorvastatin			
	Drug Hypersensitivity		(Atorvastatin)	C		
	Hyposideraemia		Gliclazide			
	Impetigo		(Gliclazide)	C		
	Leukopenia		Irbesartan			
	Oedema Peripheral		(Irbesartan)	C		
	Platelet Count Decreased		Metformin			
	Poor Venous Access		(Metformin)	C		
	Rash Erythematous					
	Renal Failure					
	Rhabdomyolysis					
	Toxic Epidermal Necrolysis					
	Uveitis					
	Visual Disturbance					
	Vitreous Disorder					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/08/05ISR Number: 4687056-6Report Type:Expedited (15-DaCompany Report #2005081724
 Age:63 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (3 IN 1 D), ORAL	Abdominal Pain Upper Anorexia Back Pain	Foreign Health	Gabapentin (Tablets) (Gabapentin)	PS		ORAL
	Constipation Dysphonia Haemoglobin Increased Headache Muscular Weakness Osteochondrosis Productive Cough Protein Urine Present Pyrexia Red Blood Cells Urine Positive White Blood Cells Urine Positive	Professional Company Representative	Doxazosin (Doxazosin) Antihypertensives (Antihypertensives)	C C		

Date:06/08/05ISR Number: 4687063-3Report Type:Expedited (15-DaCompany Report #2005077695
 Age: Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 100 MG (100 Other MG, 1 IN 1 D)	Body Height Decreased Constipation Dizziness	Consumer	Neurontin (Gabapentin)	PS		
TRANSDERMAL	BACK PAIN Impaired Driving Ability Muscle Spasms Nausea Paralysis Pruritus Somnolence		Bextra (Valdecoxib) Motrin (Ibuprofen) Fentanyl (Fentanyl) Vioxx (Rofecoxib) Advil (Ibuprofen) Aleve (Naproxen Sodium) Levoxyl (Levothyroxine Sodium)	SS SS SS SS SS C C		

All Other
Therapeutic Products
(All Other
Therapeutic
Products) C

Date:06/08/05ISR Number: 4687087-6Report Type:Expedited (15-DaCompany Report #2002054129
Age: Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anomaly Of External Ear	Literature	Neurontin			
Other		Congenital	Health	(Gabapentin)	PS		
		Congenital Acrochordon	Professional	Lamotrigine			
		Drug Exposure During		(Lamotrigine)	SS		
		Pregnancy					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/08/05ISR Number: 4687090-6Report Type:Expedited (15-DaCompany Report #2002054132

Age: Gender:Unknown I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Exposure During Pregnancy Hypospadias	Literature Health Professional	Neurontin (Gabapentin) Valproate (Valproic Acid)	PS SS		

Date:06/08/05ISR Number: 4687094-3Report Type:Expedited (15-DaCompany Report #2005042717

Age: Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Lung Neoplasm Malignant Recurrent Cancer	Consumer Health Professional	Neurontin (Gabapentin)	PS		

Date:06/08/05ISR Number: 4687095-5Report Type:Expedited (15-DaCompany Report #2005079867

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 300 MG(300 Other MG, 1 IN 1D), ORAL		Cerebrovascular Disorder Chest Pain Drug Ineffective Dysphemia Fatigue Headache Hypoaesthesia Memory Impairment Muscle Twitching Respiratory Disorder Speech Disorder Weight Decreased	Consumer	Neurontin (Gabapentin) Zoloft (Setraline) All Other Therapeutic Products(All Other Therapeutic Products)	PS C C		ORAL

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Arthralgia Drug Ineffective	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
2400 MG (800 MG, 3 IN 1 D), ORAL ORAL				Celebrex (Celecoxib)	SS		ORAL
				Lantus (Insulin Glargine)	C		
				Humalog (Insulin Lispro)	C		
				Humulin R (Insulin Human)	C		
				Norvasc (Amlodipine)	C		
				Sinequan (Doxepin)	C		

Outcome PT
Other Drug Exposure During
Pregnancy

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Eclampsia
Twin Pregnancy

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Literature Health Professional	Neurontin (Gabapentin)	PS		

Date:06/08/05ISR Number: 4687285-1Report Type:Expedited (15-DaCompany Report #2005080957
Age:44 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 400 MG (200 MG, 2 IN 1 D), ORAL		Blood Cholesterol Increased Gastric Ulcer Hypertension Intervertebral Disc Protrusion	Consumer	Celebrex (Celecoxib)	PS		ORAL
				Neurontin (Gabapentin)	SS		
				Accupril (Quinapril Hydrochloride)	SS		
				Ambien (Zolpidem Tartrate)	C		
				Effexor (Venlafaxine Hydrochloride)	C		
				Oxycontin (Oxycodone Hydrochloride)	C		
				Maxzide (Hydrochlorothiazide , Triamterene)	C		
				Potassium (Potassium)	C		

Date:06/08/05ISR Number: 4687350-9Report Type:Expedited (15-DaCompany Report #2004058042
Age:34 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 300 MG QHS,		Anxiety Completed Suicide	Consumer Health	Neurontin (Gabapentin)	PS		

MAY	Depression	Professional	
INCREASETO	Drug Ineffective		
Q8H	Feeling Abnormal		
	Gun Shot Wound	Valdecoxib	C
	Hypoaesthesia	Medrol	
	Injury	(Methylprednisolone)	C
	Insomnia	Hydrocodone	C
	Mental Disorder	Ultracet	
	Nervousness	(Paracetamol,	
	Neuralgia	Tramadol	
	Pain	Hydrochloride)	C
	Paraesthesia	Cyclobenzaprine	C
	Stress		
	Suicidal Ideation		
	Weight Decreased		

Date:06/08/05ISR Number: 4687352-2Report Type:Expedited (15-DaCompany Report #2004065552
Age:49 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
Disability
Other

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1 D)		Coagulopathy Colon Injury	Consumer	Neurontin (Gabapentin)	PS		
		Drug Ineffective					
1 MG (0.5 MG, 2 IN 1 D)		Gun Shot Wound Haemothorax		Klonopin (Clonazepam)	SS		
		Hypotension					
3 TABLETS (3 IN 1 D)		Injury Mediastinal Haemorrhage Pain		Percocet (Oxycodone Hydrochloride, Paracetamol)	SS		
		Somnolence					
60 MG (10 MG, AS NEEDED)		Splenic Injury Staphylococcal Abscess Subdiaphragmatic Abscess		Lorcet (Hydrocodone Bitartrate, Paracetamol)	SS		
		Suicide Attempt					
				Protonix (Pantoprazole)	C		
				Xalatan (Latanoprost)	C		
				Nexium (Esomeprazole)	C		
				Gemfibrozil	C		
				Pravachol (Pravastatin Sodium)	C		
				Flonase (Fluticasone Propionate)	C		
				Gaviscon (Sodium Alginate, Sodium Bicarbonate)	C		
				Alternagel (Aluminium Hydroxide Gel, Dried)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abnormal Behaviour	Consumer	Neurontin			
Disability		Anhedonia		(Gabapentin)	PS		
Other		Anxiety		Geodon (Ziprasidone)	SS		
80 MG (40 MG, 2 IN 1 D)		Completed Suicide					
		Drug Toxicity		Acetaminophen			
		Emotional Disorder		(Paracetamol)	SS		
		Incorrect Dose		Paxil (Paroxetine			
60 MG (30 MG, 2 IN 1 D)		Administered		Hydrochloride)	SS		
		Injury					
		Mental Disorder					
		Pain					
		Pharmaceutical Product					
		Complaint					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/08/05ISR Number: 4687354-6Report Type:Expedited (15-DaCompany Report #2005004215
Age:34 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin			
Other		Completed Suicide		(Gabapentin)	PS		
300 MG		Drug Ineffective		Zyprexa (Olanzapine)	C		
		Suicidal Ideation					

Date:06/08/05ISR Number: 4687355-8Report Type:Expedited (15-DaCompany Report #2004074993
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Accidental Exposure	Consumer	Neurontin			
Hospitalization -		Completed Suicide		(Gabapentin)	PS		
3200 MG (800		Drug Ineffective					
Initial or Prolonged		Drug Toxicity		Celebrex (Celecoxib)	SS		
MG, 4 IN 1 D)		Injury		Paxil (Paroxetine			
Other		Intentional Misuse		Hydrochloride)	SS		
		Mental Status Changes		Amitriptyline With			
		Metabolic Acidosis		Perphenazine			
		Overdose		(Amitriptyline			
		Respiratory Distress		Hydrochloride)	SS		
		Respiratory Failure		Oxycontin (Oxycodone			
		Suicide Attempt		Hydrochloride)	SS		
				Paracetamol	SS		
				Nexium			
				(Esomeprazole)	C		
				Baclofen	C		
				Insulin	C		
				Duragesic (Fentanyl)	C		
				Glucophage			
				(Metformin			
				Hydrochloride)	C		

Date:06/08/05ISR Number: 4687356-XReport Type:Expedited (15-DaCompany Report #2005083479
Age:62 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Neurontin (Gabapentin)	PS		
Date:06/08/05ISR Number: 4687361-3Report Type:Expedited (15-DaCompany Report #2005048413 Age:68 YR Gender:Female I/FU:F							
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1800 MG (600	Abdominal Pain Upper Amnesia	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other MG, 3 IN 1		Convulsion	Professional				
D), ORAL		Deafness					
ORAL		Dizziness Tinnitus		Sinemet (Carbidopa, Levodopa)	SS		ORAL
		Transient Ischaemic Attack		Antihypertensives (Anthihypertensives) Anti-Diabetics (Anti-Diabetics)	C C		
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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/08/05ISR Number: 4687362-5Report Type:Expedited (15-DaCompany Report #2005046321
Age:57 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Drug Ineffective	Consumer	Neurontin			
Other		Medication Error	Health	(Gabapentin)	PS		ORAL
1800 MG (600		Nerve Injury	Professional				
MG, 3 IN 1		Pain					
D), ORAL		Pharmaceutical Product Complaint		Vitamins (Vitamins)	C		

Date:06/08/05ISR Number: 4687369-8Report Type:Expedited (15-DaCompany Report #2005072525
Age:67 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Drug Ineffective	Consumer	Celebrex (Celecoxib)	PS		
200 MG (200		Feeling Abnormal	Health				
Initial or Prolonged		Infection	Professional	Neurontin			
MG, 1 IN 1 D)		Pulmonary Thrombosis		(Gabapentin)	SS		
Other		Spinal Fusion Surgery		Estrogens			
		Thrombosis		(Estrogens)	SS		
		Weight Increased		Synthroid			
				(Levothyroxine Sodium)	C		

Date:06/08/05ISR Number: 4687373-XReport Type:Expedited (15-DaCompany Report #2005003387
Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Cervical Vertebral Fracture	Consumer	Neurontin			
Hospitalization -		Conjunctivitis		(Gabapentin)	PS		
Initial or Prolonged		Corneal Ulcer					
Other		Drug Ineffective					
		Dysphagia					

Fracture
 Gun Shot Wound
 Haematoma
 Haemophilus Infection
 Mastoid Disorder
 Pneumonia
 Self Injurious Behaviour
 Subdural Haematoma
 Subdural Hygroma
 Suicide Attempt

Date:06/08/05ISR Number: 4687374-1Report Type:Expedited (15-DaCompany Report #2004106702
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin			
Other		Drug Toxicity		(Gabapentin)	PS		
		Oedema		Tramadol (Tramadol)	SS		
		Polysubstance Abuse		Ethanol (Ethanol)	SS		
		Pulmonary Congestion		Acetaminophen			
				(Acetaminophen)	SS		
				Carisoprodol			
				(Carisoprodol)	SS		
				Meprobamate			

Freedom Of Information (FOI) Report

(Meprobamate)	SS
Hydrocodone	
(Hydrocodone)	SS
Oxycodone	
(Oxycodone)	SS
Promethazine	
(Promethazine)	SS

Date:06/08/05ISR Number: 4687394-7Report Type:Expedited (15-DaCompany Report #2005069198
 Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Memory Impairment Speech Disorder Thinking Abnormal Vision Blurred	Consumer	Neurontin (Gabapentin)	PS		

Date:06/08/05ISR Number: 4687403-5Report Type:Expedited (15-DaCompany Report #2004058034
 Age:50 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization - Initial or Prolonged Disability	Affect Lability Aggression Amnesia
Other	Anger Anxiety Back Pain Benign Prostatic Hyperplasia Chest Pain Concussion Delusion Of Grandeur Dental Discomfort Depressed Mood Drug Ineffective Early Morning Awakening Hallucination, Auditory Headache Hyperhidrosis Hypoaesthesia Hypomania Illusion Initial Insomnia

Intentional Misuse
Limb Injury
Lung Neoplasm
Memory Impairment
Nephritis Interstitial
Nervousness
Off Label Use
Overdose
Pain
Pain In Extremity
Polytraumatism
Pressure Of Speech
Restlessness
Road Traffic Accident
Self-Injurious Ideation
Sleep Disorder
Speech Disorder

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
			Spinal Osteoarthritis Spousal Abuse Stress			
1800 MG,		Consumer	Neurontin (Gabapentin)	PS		
2400 MG (600			Ibuprofen (Ibuprofen)	SS		ORAL
MG, 4 IN 1						
D), ORAL			Soma (Carisoprodol)	SS		
			Olanzapine (Olanzapine)	C		
			Temazepam (Temazepam)	C		
			Clonazepam (Clonazepam)	C		
			Bupropion (Bupropion)	C		
			Depakote (Valproate Semisodium)	C		
			Effexor-Xr (Venlafaxine Hydrochloride)	C		
			Zoloft (Sertraline)	C		
			Serzone (Nefazodone Hydrochloride)	C		
			Ambien (Zolpidem Tartrate)	C		
			Gabitril (Tiagabine Hydrochloride)	C		
			Risperidone (Risperidone)	C		
			Griseofulvin (Griseofulvin)	C		
			Clotrimazole (Clotrimazole)	C		
			Vytone (Diodohydroxyquinol ine, Hydrocortisone)	C		
			Ativan (Lorazepam)	C		

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Adjustment Disorder	Consumer	Neurontin			
1800 MG		Back Pain		(Gabapentin)	PS		ORAL
(DAILY), ORAL		Convulsion					
		Drug Ineffective					
		Dyspnoea					
		Fear					
		Headache					
		Hyperhidrosis					
		Marital Problem					
		Near Drowning					
		Pain In Extremity					
		Palpitations					
		Panic Disorder					
		Suicide Attempt					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/08/05ISR Number: 4687405-9Report Type:Expedited (15-DaCompany Report #2004106748
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (300 Other MG, 3 IN 1 D),		Abnormal Behaviour Aggression Agitation Delusion Depression Drug Ineffective Homicidal Ideation	Consumer	Neurontin (Tablets) (Gabapentin)	PS		
6 MG (3 MG, 2 IN 1 D),		Intentional Misuse Lethargy Multiple Drug Overdose Psychotic Disorder Somnolence Suicidal Ideation Suicide Attempt		Haldol (Haloperidol) Risperidone (Risperidone) Seroquel (Quetiapine Fumarate) Metformin (Metformin)	SS SS C C		

Date:06/08/05ISR Number: 4687712-XReport Type:Direct Company Report #CTU 250739
 Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other ONE PO QID		Abdominal Discomfort Pharmaceutical Product Complaint		Gabapentin 300	PS		ORAL

Date:06/08/05ISR Number: 4687873-2Report Type:Expedited (15-DaCompany Report #2005081577
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abortion Spontaneous Drug Exposure During Pregnancy	Literature Health Professional	Neurontin (Gabapentin)	PS		

Date:06/08/05ISR Number: 4687874-4Report Type:Expedited (15-DaCompany Report #2005081558

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Literature	Neurontin			
		Drug Exposure During	Health	(Gabapentin)	PS		
		Pregnancy	Professional				

Date:06/08/05ISR Number: 4687876-8Report Type:Expedited (15-DaCompany Report #2005081547

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Literature	Neurontin			
		Drug Exposure During	Health	(Gabapentin)	PS		
		Pregnancy	Professional				

Date:06/08/05ISR Number: 4687970-1Report Type:Expedited (15-DaCompany Report #2005081396

Age: Gender:Female I/FU:I

Outcome	Duration	PT
Dose		
Other		Drug Exposure During
		Pregnancy

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Date:06/08/05ISR Number: 4688005-7Report Type:Expedited (15-DaCompany Report #2005081525

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Literature	Neurontin			
		Drug Exposure During	Health	(Gabapentin)	PS		
		Pregnancy	Professional				

Date:06/08/05ISR Number: 4688006-9Report Type:Expedited (15-DaCompany Report #2005081507

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous	Literature	Neurontin			
		Drug Exposure During	Health	(Gabapentin)	PS		
		Pregnancy	Professional				

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/08/05ISR Number: 4688007-0Report Type:Expedited (15-DaCompany Report #2005081495

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous Drug Exposure During Pregnancy	Literature Health Professional	Neurontin (Gabapentin)	PS		

Date:06/08/05ISR Number: 4688008-2Report Type:Expedited (15-DaCompany Report #2005081481

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Caesarean Section Drug Exposure During Pregnancy	Literature Health Professional	Neurontin (Gabapentin)	PS		

Date:06/08/05ISR Number: 4688012-4Report Type:Expedited (15-DaCompany Report #2005081465

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Caesarean Section Drug Exposure During Pregnancy Twin Pregnancy	Literature Health Professional	Neurontin (Gabapentin)	PS		

Date:06/08/05ISR Number: 4688025-2Report Type:Expedited (15-DaCompany Report #2004117393

Age:33 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged 900 MG (300 Disability MG, 3 IN 1 Other D),		Alcohol Use Anxiety Body Dysmorphic Disorder Depression Drug Ineffective	Consumer	Neurontin (Gabapentin) Trazodone	PS		

Gun Shot Wound
Head Injury
Insomnia
Irritability
Migraine
Overdose
Pain
Poisoning
Stress
Suicidal Ideation
Suicide Attempt

(Trazodone) C
Remeron
(Mirtazapine) C
Revia (Naltrexone) C

Date:06/09/05ISR Number: 4687631-9Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 250780

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300MG BID		Drug Ineffective		Gabapentin 300mg	PS		
		Therapeutic Response Unexpected With Drug Substitution					

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/09/05ISR Number: 4688231-7Report Type:Expedited (15-DaCompany Report #KII-2005-0016867
Age:46 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Agitation Belligerence Coma Drug Abuser	Study Health Professional Other	Morphine Sulfate (Similar To Nda 19-516) (Morphine Sulfate) Unknown			
SEE TEXT		Heart Rate Increased Hypertension		Carisoprodol (Carisoprodol)			PS SS
SEE TEXT		Intentional Misuse Respiratory Rate		Gabapentin (Gabapentin)			SS
800 MG, SEE TEXT		Increased					
SEE TEXT		White Blood Cell Count Increased		Other Analgesics And Antipyretics			SS
SEE TEXT				Heroin (Diamorphine)			SS
				Barbiturates Cocaine (Cocaine)			SS SS

Date:06/09/05ISR Number: 4688245-7Report Type:Expedited (15-DaCompany Report #2005044792
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability Other 600 MG (200 MG, 3 IN 1 D), ORAL		Accident Balance Disorder Chest Pain Disorientation	Consumer Health Professional	Neurontin (Gabapentin)			PS ORAL
60 MG, ORAL		Drug Ineffective Fatigue		Morphine Sulfate (Morphine Sulfate)			SS ORAL
		Febrile Neutropenia Feeling Abnormal Fibromyalgia		Oxycontin (Oxycodone Hydrochloride) Gabapentin			SS

ORAL	Heart Rate Increased	(Gabapentin)	SS	ORAL
	Hypersensitivity	Zanaflex (Tizanidine		
	Intervertebral Disc	Hydrochloride)	C	
	Protrusion	Lortab (Hydrocodone		
	Loss Of Consciousness	Bitartrate,		
	Mental Status Changes	Paracetamol)	C	
	Pharmaceutical Product	Zoloft (Sertraline)	C	
	Complaint	Atenolol (Atenolol)	C	
	Respiratory Distress	Flonase (Fluticasone		
		Propionate)	C	
		Allegra-D		
		(Fexofenadine,		
		Pseudoephedrine		
		Hydrochloride)	C	
		Mobic (Meloxicam)	C	
		Soma (Carisoprodol)	C	

Date:06/09/05ISR Number: 4688440-7Report Type:Expedited (15-DaCompany Report #2005083144
Age: Gender:Female I/FU:I

Outcome	PT
Death	Abortion Induced
Congenital Anomaly	Congenital Nose
Other	Malformation
	Drug Exposure During
	Pregnancy
	Facial Dysmorphism
	Micrognathia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Multiple Congenital Abnormalities Oesophageal Atresia	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1 D), ORAL		Retrognathia Tooth Hypoplasia	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
0.5 MG (0.25 MG, 2 IN 1 D), ORAL				Xanax Tablet (Alprazolam)	SS		ORAL
ORAL				Rivotril (Clonazepam)	SS		ORAL
ORAL				Deroxat (Paroxetine Hydrochloride)	SS		ORAL
ORAL				Di-Antalvic (Dextropropoxyphene Hydrochloride, Paracetamol)	SS		ORAL

Date:06/09/05ISR Number: 4688441-9Report Type:Expedited (15-DaCompany Report #2005083143
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 900 MG (300 MG, 3 IN 1 D), ORAL		Abortion Induced Drug Exposure During Pregnancy	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
0.5 MG (0.25				Xanax Tablet (Alprazolam)	SS		ORAL

D), ORAL

ORAL

ORAL

ORAL

Rivotril (Clonazepam)	SS	ORAL
Deroxat (Paroxetine Hydrochloride)	SS	ORAL
Di-Antalvic (Dextropropoxyphene Hydrochloride, Paracetamol)	SS	ORAL

Date:06/10/05ISR Number: 4687446-1Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20050507147

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide		Ibuprofen	PS		
		Drug Ineffective		Nortriptyline			
		Loss Of Consciousness		Hydrochloride	SS		
		Multiple Drug Overdose		Amitriptyline			
		Accidental		Hydrochloride	SS		
				Venlafaxine	SS		
				Neurontin	SS		
				Neurontin	SS		
				Neurontin	SS		
				Oxazepam	C		
				Benzodiazepine			
				Derivatives	C		
				Opioids	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/10/05ISR Number: 4687896-3Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLIN-A0554054A

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blindness		Avandia	PS	Glaxosmithkline	ORAL
4MG Per day	YR			Neurontin	SS		ORAL
300MG At							
night	YR			Glucagon	C		
				Iron	C		
				Lasix	C	Glaxosmithkline	
				Protonix	C		
				Percocet	C		
				Novolog	C		
				Humulin R	C		
				Lipitor	C		
				Rocaltrol	C		
				Alprazolam	C		
				Oscal	C	Glaxosmithkline	
				Aranesp	C		

Date:06/10/05ISR Number: 4688080-XReport Type:Expedited (15-DaCompany Report #FR-ROCHE-406439

Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Induced		Rivotril	PS	Roche	ORAL
UNKNOWN		Drug Exposure During		Neurontin	SS		
		Pregnancy		Xanax	SS		ORAL
		Facial Dysmorphism		Deroxat	SS		ORAL
DOSING		Multiple Congenital					
REGIMEN		Abnormalities					
REPORTED AS		Oesophageal Atresia					
0.5 DOSES A							
DAY.							
DOSING				Di Antalvic	SS		ORAL

REGIMEN

REPORTED AS 6

DOSES A DAY.

Date:06/10/05ISR Number: 4691028-5Report Type:Expedited (15-DaCompany Report #2004030610

Age:75 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1500 MG (3 IN Disability 1 D), ORAL	Arthralgia Muscular Weakness Myalgia	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
Other 200 MG (200 MG, 1 IN 1 D), ORAL	Myopathy Nuclear Magnetic Resonance Imaging Abnormal Oedema Oedema Peripheral Pain In Extremity	Health Professional	Celebrex (Celecoxib) Voltaren (Diclofenac Sodium) Venostasin (Horse Chestnut Extract, Thiamine Hydrochloride) All Other Therapeutic Products (All Other Therapeutic Products) Biomagnesin (Citric Acid, Magnesium	SS C C C		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Citrate, Magnesium
Phosphate Dibasic) C

Date:06/13/05ISR Number: 4689807-3Report Type:Direct
Age:48 YR Gender:Female I/FU:I

Company Report #CTU 251046

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Disability		Incorrect Dose Administered Suicide Attempt		Neurontin Pfizer Seroquel Senteca	PS SS	Pfizer Astra Astra Senteca	

Date:06/13/05ISR Number: 4690208-2Report Type:Expedited (15-DaCompany Report #2005046003
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Convulsion Diarrhoea Drug Ineffective	Consumer Health Professional	Phenytoin Suspension (Phenytoin Sodium)	PS		
UNKNOWN	300 MG,	Nausea					
UNKNOWN		Transient Ischaemic Attack		Gabapentin (Gabapentin)	SS		
UNKNOWN	UNKNOWN			Lipitor (Atorvastatin)	SS		
UNKNOWN	UNKNOWN						

Date:06/13/05ISR Number: 4690377-4Report Type:Expedited (15-DaCompany Report #2005083192
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Cerebrovascular Accident Loss Of Consciousness Nervous System Disorder Therapeutic Response Unexpected With Drug Substitution	Consumer	Neurontin (Gabapentin) Gabapentin(Gabapenti n)	PS SS		

Date:06/13/05ISR Number: 4690681-XReport Type:Expedited (15-DaCompany Report #2005082064
Age:54 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Condition Aggravated	Consumer	Neurontin			
Other		Overdose		(Gabapentin)	PS		ORAL
900 MG (300							
MG, 3 IN 1		Pain In Extremity					
D), ORAL							
				Diuretics			
				(Diuretics)	C		
				All Other			
				Therapeutic Products			
				(All Other			
				Therapeutic			
				Products)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/13/05ISR Number: 4690789-9Report Type:Expedited (15-DaCompany Report #US015151
Age:71 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - 4 MG BID ORAL	Blood Glucose Decreased	Health	Gabitril	PS		ORAL
Initial or Prolonged 300 MG ORAL	Cardiovascular Disorder Cerebral Disorder	Professional	Gabitril Neurontin	SS SS		ORAL
	Convulsion Drug Interaction Dyskinesia Loss Of Consciousness Muscle Twitching Speech Disorder Syncope Tremor					

Date:06/14/05ISR Number: 4690611-0Report Type:Direct Company Report #CTU 251062
Age:70 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged	Convulsion Hypotension Mental Status Changes		Gabapentin Morphine Zosyn	PS C C		

Date:06/14/05ISR Number: 4692005-0Report Type:Expedited (15-DaCompany Report #2005042430
Age:63 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Other 400 MG (200 MG, BID INTERVAL: BID)ORAL	Aphasia Arthralgia Dizziness Dysgraphia Ear Pain Fear	Consumer Health Professional	Neurontin (Gabapentin)	PS		ORAL

Headache
 Memory Impairment
 Meningioma
 Pharmaceutical Product
 Complaint
 Speech Disorder

Date:06/14/05ISR Number: 4692239-5Report Type:Expedited (15-DaCompany Report #CA-2005-007813

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Condition Aggravated Cystitis Dyspnoea	Foreign Consumer Other	Betaseron (Interferon Beta-1b) Injection, 250ug	PS		
SUBCUTANEOUS	8 MIU, EVERY 2D,	Multiple Sclerosis Oedema Peripheral		Gabapentin (Gabapentin)	SS		
SUBCUTANEOUS				Baclofen	C		
				Zanaflex (Tizanidine Hydrochloride)	C		
				Amitriptyline (Amitriptyline)	C		
				Diovan (Valsartan)	C		
				Nexium			

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Freedom Of Information (FOI) Report

(Esomeprazole) C
 Tylenol
 Extra-Strenght C

Date:06/15/05ISR Number: 4692881-1Report Type:Expedited (15-DaCompany Report #2005056759
 Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (400 MG),		Irritable Bowel Syndrome Serotonin Syndrome	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:06/15/05ISR Number: 4693022-7Report Type:Expedited (15-DaCompany Report #2005044232
 Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 800 MG (1 D),		Alopecia Bladder Prolapse Breath Odour	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Drug Ineffective Drug Level Increased Dysgeusia Gangrene Hair Disorder Hypoaesthesia Malaise Panic Attack Pharmaceutical Product Complaint Tooth Discolouration Tooth Loss		Lithium (Lithium) Cyclobenzaprine (Cyclobenzaprine) Thyroid (Thyroid) Clonazepam (Clonazepam) Avandia (Rosiglitazone Maleate) Naproxen (Naproxen) Zocor (Simvastatin)	SS C C C C C C		

Date:06/15/05ISR Number: 4693699-6Report Type:Expedited (15-DaCompany Report #2005085537
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Anorexia	Consumer	Neurontin			
Other		Arthralgia		(Gabapentin)	PS		
		Blindness Unilateral					
		Cerebrovascular Accident					
		Cyst					
		Discomfort					
		Emotional Distress					
		Gait Disturbance					
		Loss Of Consciousness					
		Malaise					
		Pain In Jaw					
		Speech Disorder					
		Tinnitus					
		Wrong Drug Administered					

Date:06/15/05ISR Number: 4693858-2Report Type:Expedited (15-DaCompany Report #2005085463
Age:76 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization -	Aortic Valve Incompetence
Initial or Prolonged	Intestinal Obstruction
Other	Lung 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
200 MG (200 MG, 1 IN 1 D), ORAL		Pneumonia Pulmonary Hypertension Respiratory Failure Ventricular Dysfunction	Consumer	Celebrex (Celecoxib)	PS		ORAL
ORAL				Neurontin (Gabapentin)	SS		ORAL
				Prilosec (Omeprazole)	C		
				Trazodone (Trazodone)	C		
				Antibiotics (Antibiotics)	C		
				Nitrofurantoin (Nitrofurantoin)	C		
				Amitriptyline (Amitriptyline)	C		
				Clonazepam (Clonazepam)	C		
				Folic Acid (Folic Acid)	C		

Date:06/16/05ISR Number: 4693791-6Report Type:Expedited (15-DaCompany Report #2005069871
Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Pressure Increased Body Temperature	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
600 MG (300 MG, 2 IN 1 D), ORAL		Increased Convulsion	Professional Other	Senna (Senna)	C		
				Glycerin (Glycerin)	C		
				Thiamine (Thiamine)	C		

Dihydrocodeine C
 (Dihydrocodeine)
 Meloxicam C
 (Meloxicam)
 Vitamin B (Vitamin B) C

Date:06/16/05ISR Number: 4693880-6Report Type:Expedited (15-DaCompany Report #2005087005
 Age:79 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (600 MG, 2 IN 1 D) 15 DROP	Epilepsy Sepsis Skin Infection Skin Ulcer Somnolence	Foreign Health Professional Company Representative	Neurontin (Gabapentin) Rivotril (Clonazepam)	PS SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/16/05ISR Number: 4693990-3Report Type:Expedited (15-DaCompany Report #2005086493

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Alanine Aminotransferase Increased	Consumer	Neurontin (Gabapentin)	PS		
		Blood Hiv Rna Increased		Crixivan (Indinavir Sulfate)	SS		
		Blood Pressure Increased					
		Convulsion					
		Depression					
		Gastrointestinal Disorder					
		Hypoaesthesia					
		Immune System Disorder					
		Lipoma					
		Nerve Injury					
		Nervousness					
		Overdose					
		Pain					
		Sleep Disorder					
		Suicidal Ideation					

Date:06/16/05ISR Number: 4694074-0Report Type:Expedited (15-DaCompany Report #2004106719

Age:56 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death							
Life-Threatening		Completed Suicide	Consumer	Neurontin (Gabapentin)	PS		ORAL
1500 MG (300		Depression					
Hospitalization -		Drug Ineffective					
MG, 5 IN 1							
Initial or Prolonged		Drug Toxicity					
D), ORAL							
Other		Intentional Misuse		Diphenhydramine (Diphenhydramine)	SS		ORAL
ORAL		Polysubstance Abuse					
		Suicidal Ideation		Toprol (Metoprolol)	SS		
20 MG (1 IN 1							
D),							
(350 MG, 1 OR				Soma (Carisoprodol)	SS		ORAL
2 EVERY							

4-6HOURS AS

NECESSARY),

ORAL

(4 MG, 1-2

EVERY

4-6HOURS AS

NECESSARY),

ORAL

Meprobamate
(Meprobamate) SS
Dilaudid
(Hydromorphone
Hydrochloride) SS

ORAL

Prednisone
(Prednisone) C
Duragesic (Fentanyl) C
Plendil (Felodipine) C
Catapres (Clonidine) C
Altace (Ramipril) C
Potassium
(Potassium) C
Prevacid
(Lansoprazole) C
Clonazepam
(Clonazepam) C
Ursodiol
(Ursodeoxycholic

Freedom Of Information (FOI) Report

Acid)	C
Starlix	
(Nateglinide)	C
Lorazepam	
(Lorazepam)	C
Albuterol	
(Salbutamol)	C
Lasix (Furosemide)	C

Date:06/16/05ISR Number: 4694075-2Report Type:Expedited (15-DaCompany Report #2005003387
 Age:59 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Basal Cell Carcinoma	Consumer	Neurontin			
Hospitalization -	Cervical Vertebral		(Gabapentin)	PS		
Initial or Prolonged	Fracture					
Other	Conjunctivitis					
	Drug Ineffective					
	Dysphagia					
	Gun Shot Wound					
	Pneumonia Haemophilus					
	Self Injurious Behaviour					
	Skull Fracture					
	Subarachnoid Haemorrhage					
	Subdural Haematoma					
	Subdural Hygroma					
	Suicide Attempt					
	Unresponsive To Verbal					
	Stimuli					

Date:06/16/05ISR Number: 4694084-3Report Type:Expedited (15-DaCompany Report #2004058034
 Age:50 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Abnormal Behaviour
Initial or Prolonged	Affect Lability
Disability	Aggression
Other	Amnesia
	Anger
	Anxiety
	Arthralgia
	Back Pain
	Benign Prostatic

Hyperplasia
Chest Pain
Concussion
Confusional State
Deformity
Delusion
Delusional Perception
Dental Discomfort
Depressed Mood
Drug Ineffective
Drug Screen Positive
Early Morning Awakening
Feeling Hot
Grandiosity
Hallucination
Hallucination, Auditory
Headache

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Age:35 YR Gender:Male I/FU:F

Outcome	PT
Death	Activities Of Daily
Hospitalization -	Living Impaired
Initial or Prolonged	Atrophy
Other	Back Pain
	Completed Suicide
	Depression
	Drug Ineffective
	Drug Toxicity
	Intervertebral Disc
	Disorder
	Intervertebral Disc
	Protrusion
	Mass
	Muscle Spasms
	Neck Pain

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pain In Extremity Pulmonary Congestion Pulmonary Oedema	Report Source	Product	Role	Manufacturer	Route
		Spinal Cord Injury Spinal Disorder	Consumer	Neurontin (Gabapentin)	PS		
				Tramadol (Tramadol)	SS		
				Ethanol (Ethanol)	SS		
				Acetaminophen (Paracetamol)	SS		
				Carisoprodol (Carisoprodol)	SS		
				Meprobamate (Meprobamate)	SS		
				Hydrocodone (Hydrocodone)	SS		
				Oxycodone (Oxycodone)	SS		
				Promethazine (Promethazine)	SS		
				Skelaxin (Metaxalone)	C		
				Cataflam (Diclofenac Sodium)	C		
				Darvocet (Dextropropoxyphene Napsilate, Paracetamol)	C		
				Vioxx (Rofecoxib)	C		

Date:06/16/05ISR Number: 4694086-7Report Type:Expedited (15-DaCompany Report #2005049905
Age:32 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Complex Partial Seizures Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		
800 MG (400 MG, 2 IN 1 D),		Potentiation Electroencephalogram Abnormal Loss Of Consciousness Multiple Allergies		Astelín (Azelastine Hydrochloride) Tegretol	SS		

1000 MG (500
 MG, 2 IN 1 D)
 Muscle Rigidity (Carbamazepine) SS
 Petit Mal Epilepsy
 Pharmaceutical Product Ativan (Lorazepam) C
 Complaint

Date:06/17/05ISR Number: 4693282-2Report Type:Expedited (15-DaCompany Report #US-GLAXOSMITHKLINE-A0562784A
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 25MG Single Initial or Prolonged dose	1 DAY	Blood Pressure Increased		Coreg	PS	Glaxosmithkline	ORAL
1 DAY		Dizziness					
		Headache		Aldactone	SS		
1 DAY		Hypotension		Vasotec	SS		
1 DAY		Medication Error		Neurontin	SS		
		Swelling		Toprol Xl	C		
		Vision Blurred		Insulin	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/17/05ISR Number: 4694242-8Report Type:Direct
 Age: Gender:Male I/FU:I

Company Report #CTU 251428

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 MG PO		Loss Of Consciousness		Neurontin	PS		ORAL
		Therapeutic Response Unexpected With Drug Substitution					

Date:06/17/05ISR Number: 4694871-1Report Type:Expedited (15-DaCompany Report #2004106748
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged UNKNOWN Other MG, 3 IN 1 D), UNKNOWN	900 MG, (300	Abnormal Behaviour Aggression Agitation Delusion	Consumer	Neurontin (Tablets) (Gabapentin)	PS		
UNKNOWN UNKNOWN UNKNOWN IN 1 D), UNKNOWN	UNKNOWN 6 MG (3 MG, 2	Depression Drug Ineffective Homicidal Ideation Intentional Misuse Lethargy		Haldol (Haloperidol) Risperidone (Risperidone)	SS SS		
		Multiple Drug Overdose Psychotic Disorder Somnolence Suicidal Ideation Suicide Attempt		Seroquel (Quetiapine Fumarate) Metformin (Metformin)	C C		

Date:06/17/05ISR Number: 4694875-9Report Type:Expedited (15-DaCompany Report #2004012957
 Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization -	Alcohol Use	Consumer	Neurontin	
Initial or Prolonged	Bipolar Disorder		(Gabapentin)	PS
UNKNOWN	600 MG (300			
Disability	Condition Aggravated			
MG, 2 IN 1				
Other	Drug Dependence			
D), UNKNOWN				
	Facial Palsy			
	Gun Shot Wound			
	Head Injury			
	Injury			
	Movement Disorder			
	Nicotine Dependence			
	Pain			
	Palpitations			
	Post Procedural			
	Complication			
	Suicidal Ideation			
	Suicide Attempt			
	Swelling			
	Tenderness			

Date:06/17/05ISR Number: 4694876-0Report Type:Expedited (15-DaCompany Report #2005042128
Age:74 YR Gender:Male I/FU:F

Outcome	PT
Other	Blood Disorder
	Chest Wall Pain
	Drug Effect Decreased
	Metastases To Lung

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pharmaceutical Product Complaint Sarcoma	Report Source	Product	Role	Manufacturer	Route
200 MG (100 MG), ORAL			Consumer Health	Neurontin (Gabapentin)	PS		ORAL
200 MG (100 MG, 2 IN 1 D), ORAL			Professional	Celebrex (Celecoxib)	SS		ORAL
				Cozarr (Losartan Potassium)	C		
				Synthroid (Levothyroxine Sodium)	C		
				Norvasc (Amlodipine)	C		
				Zantac (Ranitidine Hydrochloride)	C		
				Atenolol (Atenolol)	C		
				Furosemide (Furosemide)	C		
				Pravachol (Pravastatin Sodium)	C		
				Ultracet (Paracetamol, Tramadol Hydrochloride)	C		
				Tranxene (Clorazepate Dipotassium)	C		
				Bayer Children'S Aspirin (Acetylsalicylic Acid)	C		

Date:06/17/05ISR Number: 4694882-6Report Type:Expedited (15-DaCompany Report #2005087093

Age: Gender:Female I/FU:I

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
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Other	Cerebrovascular Accident	Consumer	Celebrex (Celecoxib)	PS
	Muscle Spasms		Neurontin	
500 MG, 3 IN	Road Traffic Accident		(Gabapentin)	SS
1 D			Naproxen (Naproxen)	C

Date:06/17/05ISR Number: 4695068-1Report Type:Expedited (15-DaCompany Report #2005086982
Age:15 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization -	Confusional State	Foreign	Neurontin			
Initial or Prolonged	Convulsion	Health	(Gabapentin)	PS		
900 MG (300						
MG,)	Drug Interaction	Professional				
		Company	Tramadol (Tramadol)	SS		
		Representative	Amitriptyline			
			(Amitriptyline)	C		
			Clonidine			
			(Clonidine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/17/05ISR Number: 4695438-1Report Type:Direct
Age:55 YR Gender:Male I/FU:I

Company Report #CTU 251462

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coordination Abnormal		Gabapentin	PS		
				Fosinopril	C		
				Furosemide	C		
				Insulin Nph (Human)	C		
				Morphine	C		
				Paroxetine	C		
				Paroxetine	C		
				Insulin Reg Human			
				Inj Novolin R	C		
				Fentanyl	C		
				Docusate/Sennosides	C		
				Omeprazole	C		
				Epoetin Alpha	C		
				Aspirin	C		
				Lorazepam	C		
				Felodipine	C		

Date:06/17/05ISR Number: 4695642-2Report Type:Direct
Age: Gender:Male I/FU:I

Company Report #CTU 251455

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Diarrhoea		Neurontin (Generic)	PS		
300 MG -6 TAB							
		Muscle Spasms					
DAILY							
		Pharmaceutical Product Complaint					

Date:06/17/05ISR Number: 4695662-8Report Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 251456

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Abortion Induced		Neurontin	PS		ORAL
200 MG P.O.							
Congenital Anomaly		Alpha 1 Foetoprotein					
BID							

(DURATION:
YEARS)
Increased
Amniotic Band Syndrome
Congenital Anomaly
Congenital Foot
Malformation
Gastroschisis
Lower Limb Deformity
Maternal Drugs Affecting
Foetus
Prenatal
Multivitamins C

Date:06/20/05ISR Number: 4694266-0Report Type:Expedited (15-DaCompany Report #US-BOEHRINGER INGELHEIM PHARMACEUTICALS, INC.-2005-
BAge:66 YR Gender:Female I/FU:F

Outcome PT
Hospitalization - Alanine Aminotransferase
Initial or Prolonged Increased
Aspartate
Aminotransferase
Increased
Asthenia
Back Pain
Blood Creatine
Phosphokinase Increased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
		Blood Glucose Increased Cellulitis Clostridium Colitis				
		Condition Aggravated Decreased Appetite Diarrhoea Difficulty In Walking Dyspnoea Erythema Fatigue Feeling Abnormal	Micardis Tablets	PS	B.I. Pharmaceuticals, Inc. /Ridgefield	ORAL
RESPIRATORY (INHALATION)	QID	Gastrooesophageal Reflux Disease	Albuterol	C		
PRN RESPIRATORY (INHALATION)	QID	Myalgia Nausea Oedema Oedema Peripheral Pain In Extremity Rhabdomyolysis Sinus Tachycardia	Atrovent	C		
			Protonix Amaryl Calan Sr K-Dur Mtv Calcium Msm Lasix Synthroid Zetia	C C C C C C C C C C		
every other month			Zyrtec Zitrix	C C		

Date:06/20/05ISR Number: 4695423-XReport Type:Direct
Age:43 YR Gender: I/FU:I

Company Report #CTU 251512

Outcome Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG ONE				Gabapentin	PS		ORAL

BID PO

Pharmaceutical Product

Complaint

Date:06/20/05ISR Number: 4695810-XReport Type:Direct
Age: Gender:Female I/FU:I

Company Report #CTU 251549

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Nausea		Gabapentin (Generic)	PS		ORAL
400 MG PO TID		Pharmaceutical Product Complaint Vomiting					

Date:06/20/05ISR Number: 4696169-4Report Type:Expedited (15-DaCompany Report #2004055469
Age:60 YR Gender:Female I/FU:F

Outcome	PT
Death	Cardiac Arrest
Hospitalization -	Epilepsy
Initial or Prolonged	Hepatic Failure
	Ketoacidosis
	Lactic Acidosis
	Prothrombin Time
	Prolonged
	Respiratory Distress

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Septic Shock

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
(2 IN 1 D), ORAL		Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
3 GRAM (1 IN 1 D), ORAL			Metformin (Metformin Hydrochloride) (Metformin)	SS		ORAL
(6 IN 1 D), ORAL			Rifadine (Rifampicin)	SS		ORAL
(1 D), ORAL			Macrochantin (Nitrofurantoin)	SS		ORAL
400 MG (400 MEQ, 1 IN 1 D), ORAL			Oflocet (Ofloxacin)	SS		ORAL
			Aprovel (Irbesartan) Ferrograd C (Ferrous Sulfate Exsiccated, Sodium Ascorbate)	C C		

Date:06/20/05ISR Number: 4696287-0Report Type:Expedited (15-DaCompany Report #2005086719

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other (2 MG,)		Fatigue Imprisonment Loss Of Consciousness	Health Professional	Xanax (Alprazolam)	PS		
		Murder		Neurontin			

1800 MG (300
MG, 6 IN 1
D),

Desoxyn

(Metamfetamine
Hydrochloride) SS

ORAL

180 MG (30
MG, 6 IN 1
D), ORAL 18 MON

Duragesic
(Fentanyl) SS

(100 MCG, 2
EVERY 72
HOURS),

Prilosec
(Omeprazole) SS

ORAL

40 MG (40 MG,
1 IN 1 D),

Oxycontin
(Oxycodone
Hydrochloride) SS

480 MG (80
MG, 6 IN 1
D),

Freedom Of Information (FOI) Report

Date:06/20/05ISR Number: 4696371-1Report Type:Expedited (15-DaCompany Report #KII-2005-0017035
 Age:63 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged Other	Depressed Level Of Consciousness Hemiparesis Pneumonia	Study Health Professional Other	Oxycodone Hydrochloride (Oxycodone Hydrochloride)	PS		ORAL
ORAL						
	Respiratory Rate Decreased		Morphine Sulfate (Morphine Sulfate)	SS		ORAL
ORAL						
	Septic Shock		Verapamil (Verapamil) Metformin (Metformin) Ssri (Gabapentin (Gabapentin) Benztropeine (Benztropeine) Colchicine (Colchicine) Proton Pump Inhibitor (Furosemide (Furosemide) Hydrochlorothiazide (Hydrochlorothiazide) Acetylsalicylic Acid (Acetylsalicylic Acid) Folic Acid (Folic Acid) Docusate (Docusate)	SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS SS		

Date:06/20/05ISR Number: 4696388-7Report Type:Expedited (15-DaCompany Report #USA-2005-0020549
 Age: Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Fatigue Loss Of Consciousness	Health Professional	Oxycontin Tablets (Oxycodone			

	Murder	Other	Hydrochloride) Cr		
	Road Traffic Accident		Tablet	PS	
80 MG, Q4H,					
			Desoxyn		
			(Metamfetamine		
			Hydrochloride)	SS	ORAL
30 MG, Q4H,					
ORAL					
			Xanax (Alprazolam)	SS	
2 MG, UNK					
			Duragesic (Fentanyl)	SS	
100 MCG,					
			Prilosec		
			(Omeprazole)	SS	
40 MG, SEE					
TEXT,					
			Neurontin		
			(Gabapentin)	SS	
300 MG, Q4H					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/20/05ISR Number: 4696452-2Report Type:Expedited (15-DaCompany Report #2005087274
Age:25 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Anxiety	Health	Neurontin			
ORAL		Priapism	Professional	(Gabapentin)	PS		ORAL
		Retching		Trazodone			
		Vomiting		(Trazodone)	C		

Date:06/20/05ISR Number: 4696462-5Report Type:Expedited (15-DaCompany Report #2004083654
Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Anxiety	Consumer	Neurontin			
Other		Completed Suicide		(Gabapentin)	PS		ORAL
ORAL		Depression					
		Injury					
		Mental Disorder					
		Pain					
		Suicidal Ideation					

Date:06/20/05ISR Number: 4696464-9Report Type:Expedited (15-DaCompany Report #2005004121
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Adverse Drug Reaction	Consumer	Lipitor			
Initial or Prolonged		Anxiety	Health	(Atorvastatin)	PS		ORAL
(20 MG),ORAL							
Other		Asthenia	Professional	Neurontin			
		Blood Creatine		(Gabapentin)	SS		
ORAL		Phosphokinase Increased		Bextra (Valdecoxib)	SS		ORAL
		Blood Testosterone		Synthroid			
		Decreased		(Levothyroxine			
		Chest Pain		Sodium)	SS		ORAL
(0.5 MG),ORAL							
		Demyelination		Aceon (Perindopril			
		Depression		Erbumine)	SS		

Disturbance In Attention	Hydrochlorothiazide	
Feeling Abnormal	(Hydrochlorothiazide	
Hypertension)	SS
Hypoaesthesia	Topamax (Topiramate)	C
Impaired Work Ability	Aleve (Naproxen	
Lethargy	Sodium)	C
Malaise	Asa (Acetylsalicylic	
Mental Disorder	Acid)	C
Muscle Disorder	Levoxyl	
Muscle Tightness	(Levothyroxine	
Muscle Twitching	Sodium)	C
Muscular Weakness	Ambien (Zolpidem	
Myalgia	Tartrate)	C
Myocardial Infarction		
Nervousness		
Pain		
Pain In Extremity		
Sensory Disturbance		
Sleep Apnoea Syndrome		
Therapy Non-Responder		
Thinking Abnormal		
Thyroid Function Test		
Abnormal		
Viral Infection		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/20/05ISR Number: 4696465-0Report Type:Expedited (15-DaCompany Report #2005016042

Age: Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Aggression	Consumer	Neurontin			
Initial or Prolonged	Bacteraemia		(Gabapentin)	PS		
Other	Brain Oedema					
	Cerebrovascular Accident					
	Computerised Tomogram					
	Abnormal					
	Convulsion					
	Depressed Level Of					
	Consciousness					
	Drug Ineffective					
	Dyskinesia					
	Encephalomalacia					
	Gun Shot Wound					
	Headache					
	Hemiparesis					
	Intracranial Injury					
	Major Depression					
	Memory Impairment					
	Paraparesis					
	Pneumonia					
	Pulmonary Congestion					
	Pulmonary Vascular					
	Disorder					
	Respiratory Failure					
	Sinus Disorder					
	Skin Ulcer					
	Subdural Haematoma					
	Suicide Attempt					
	Thrombosis					
	Wound Dehiscence					
	Wound Infection					

Date:06/20/05ISR Number: 4696542-4Report Type:Expedited (15-DaCompany Report #2005071190

Age:55 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -	Chest Pain	Health	Neurontin			
Initial or Prolonged	Dyspnoea	Professional	(Gabapentin)	PS		
1600 MG (800						

Tenormin (Atenolol)	C
Avandamet (Metformin Hydrochloride, Rosiglitazone)	C
Crestor (Rosuvastatin)	C
Nph Insulin (Insulin Injection, Isophane)	C
Insulin, Regular (Insulin)	C
Ultracet (Paracetamol, Tramadol Hydrohchloride)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/20/05ISR Number: 4696555-2Report Type:Expedited (15-DaCompany Report #2005054277

Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
2400 MG (600			Professional				
MG, 4 IN 1							
D), ORAL							

Klonopin (Clonazepam) C

Date:06/20/05ISR Number: 4697329-9Report Type:Expedited (15-DaCompany Report #2005087203

Age:88 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged (300 MG),		Drug Interaction Post Procedural	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Complication	Professional				
		Somnolence		Df118 (Dihydrocodeine Bitartrate)	SS		ORAL
(40 MG), ORAL							

Diazepam (Diazepam) C
Ketoprofen (Ketoprofen) C
Gentamicin (Gentamicin) C

Date:06/21/05ISR Number: 4697158-6Report Type:Direct Company Report #CTU 251600

Age:55 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Confusional State		Neurontin 300 Mg Tid	PS		
TID ORAL X 3							

DAY

Dysarthria

Vertigo

Valproic Acid	C
Mood Stabilizer	C
Ranitidine	C
Felodipine	C
Prilosec	C
Metoprolol	C

Date:06/21/05ISR Number: 4697596-1Report Type:Expedited (15-DaCompany Report #US_0408105111

Age:79 YR Gender:Female I/FU:F

Outcome	PT
Death	Abdominal Pain
Hospitalization -	Abdominal Pain Lower
Initial or Prolonged	Affective Disorder
Other	Agitation
	Anaemia
	Anxiety
	Aortic Aneurysm
	Arthritis
	Atherosclerosis
	Back Pain
	Bipolar Disorder
	Blood Calcium Decreased
	Bowel Sounds Abnormal
	Bronchial Infection
	Bronchitis

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Freedom Of Information (FOI) Report

Carbon Dioxide Increased
Cardiomegaly
Chronic Obstructive
Airways Disease
Exacerbated
Coma
Compression Fracture
Condition Aggravated
Confusional State
Cystitis Interstitial
Decreased Appetite
Delirium
Delusional Disorder,
Persecutory Type
Depressed Level Of
Consciousness
Diarrhoea
Dissociation
Dizziness
Drug Ineffective
Drug Interaction
Drug Toxicity
Dyspnoea Exertional
Electrocardiogram St
Segment Abnormal
Encephalopathy
Eosinophil Count
Increased
Fluid Intake Reduced
Grunting
Haemorrhage
Hallucination
Heart Rate Increased
Hunger
Hyperglycaemia
Hyperhidrosis
Hypertension
Hypotension
Inappropriate Schedule Of
Drug Administration
Increased Appetite
Interstitial Lung Disease
Major Depression
Medication Error
Micturition Urgency
Nausea
Nightmare

Oral Intake Reduced
Pain
Pancreatitis Haemorrhagic
Pollakiuria
Protein Total Decreased
Respiration Abnormal
Restlessness
Shoulder Pain
Suicidal Ideation
Thoracic Vertebral
Fracture
Tremor
Upper Respiratory Tract
Infection

Peri-Colace	C
Methylprednisolone	C
Proventil	
(Salbutamol Sulfate)	C
Combivent	C
Duragesic (Fentanyl)	C
Celexa (Citalopram	
Hydrobromide)	C
Wellbutrin	
(Bupropion	
Hydrochloride)	C
Depakote (Valproate	
Semisodium)	C
Senokot (Senna	
Alexandrina)	C
Elmiron (Pentosan	

Freedom Of Information (FOI) Report

Polysulfate Sodium)	C	
Macrobid		
(Nitrofurantoin)	C	
Ultram (Tramadol		
Hydrochloride)	C	
Caltrate + D	C	
Advair	C	
Flovent (Fluticasone		
Propionate)	C	
Oxycontin (Oxycodone		
Hydrochloride)	C	
Mellaril		
(Thioridazine		
Hydrochloride)	C	
Simethicone	C	
Darvocet-N	C	
Nardil (Phenelzine)	C	
Prilosec (Omeprazole		
Ratiopharm)	C	Ratiopharm
Ativan (Lorazepam)	C	
Entric Aspirin		
(Acetylsalicylic		
Acid)	C	
Mavik (Trandolapril)	C	
Os-Cal	C	
Humabid (Guaifenesin		
L.A.)	C	

Date:06/21/05ISR Number: 4697984-3Report Type:Expedited (15-DaCompany Report #2005088879
 Age:81 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Chest Pain	Foreign	Gabapentin			
Other		Drug Withdrawal Syndrome	Literature	(Gabapentin)	PS		
		Hypertension	Health				
		Influenza Like Illness	Professional				
		Mental Status Changes					
		Upper Respiratory Tract					
		Infection					

Date:06/21/05ISR Number: 4698308-8Report Type:Expedited (15-DaCompany Report #2005083143
 Age:35 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abortion Induced Congenital Anomaly	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 Other MG, 3 IN 1 D), ORAL		Congenital Jaw Malformation	Professional				
0.5 MG (0.25 MG, 2 IN 1 D), ORAL		Congenital Nose Malformation		Xanax Tablet (Alprazolam)	SS		ORAL
		Drug Exposure During Pregnancy					
ORAL		Facial Dysmorphism Intra-Uterine Death		Rivotril (Clonazepam)	SS		ORAL
ORAL		Micrognathia Oesophageal Atresia		Deroxat (Paroxetine Hydrochloride)	SS		ORAL
ORAL		Retrognathia		Di-Antalvic (Dextropropoxyphene Hydrochloride, Paracetamol)	SS		ORAL

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/21/05ISR Number: 4698332-5Report Type:Expedited (15-DaCompany Report #2005083144

Age:0 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Abortion Induced	Foreign	Neurontin			
Congenital Anomaly		Congenital Jaw	Health	(Gabapentin)	PS		ORAL
900 MG		Malformation	Professional				
Other		Congenital Nose					
(300MG, 3 IN		Malformation		Xanax Tablet			
1 D), ORAL		Drug Exposure During		(Alprazolam)	SS		ORAL
0.5 MG (0.25		Pregnancy					
MG, 2 IN 1		Facial Dysmorphism					
D), ORAL		Foetal Malformation		Rivotril			
ORAL		Intra-Uterine Death		(Clonazepam)	SS		ORAL
		Micrognathia		Deroxat (Paroxetine			
ORAL		Oesophageal Atresia		Hydrochloride)	SS		ORAL
		Retrognathia		Di-Antalvic			
ORAL				(Dextropropoxyphene			
				Hydrochloride,			
				Paracetamol)	SS		ORAL

Date:06/22/05ISR Number: 4696897-0Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20050603536

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Fatigue		Duragesic	PS		
TRANSDERMAL		Loss Of Consciousness		Desoxyn	SS		
OROPHARINGEAL		18 MON		Desoxyn	SS		
OROPHARINGEAL		Road Traffic Accident					
		18 MON		Xanax	SS		
				Oxycontin	SS		
				Neurontin	SS		

Date:06/22/05ISR Number: 4697302-0Report Type:Expedited (15-DaCompany Report #2005UW08507

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose				Prilosec	PS		ORAL
Other		Fatigue		Desoxyn	SS		ORAL
18	MON	Imprisonment					
		Loss Of Consciousness		Desoxyn	SS		ORAL
18	MON	Road Traffic Accident		Xanax	SS		
				Oxycontin	SS		
				Duragesic Patches	SS		
				Neurontin	SS		

Date:06/22/05ISR Number: 4697698-XReport Type:Expedited (15-DaCompany Report #2005088508

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin			
		Leg Amputation		(Gabapentin)	PS		

Date:06/22/05ISR Number: 4698109-0Report Type:Direct Company Report #CTU 251670

Age:86 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Mental Status Changes		Gabapentin	PS		ORAL
200MG BID				200mg			
Initial or Prolonged		Myoclonus					
ORAL							

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Cardizem Cd	C
Lanoxin	C
Novolog	C
Lasix	C
Vitamin B12	C
Calcium/Vitamin D	C
Aricept	C
Miacalcin Ns	C
Flonase	C
Prevacid	C
Effexor	C
Lopi	C

Date:06/22/05ISR Number: 4698488-4Report Type:Expedited (15-DaCompany Report #2005088816
 Age:48 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Cerebral Disorder Depression	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other	Drug Ineffective Nasopharyngitis		Vitamins (Vitamins) Keppra (Levetiracetam) All Other Therapeutic Products (All Other Therapeutic Products)	SS C C		
			Ativan (Lorazepam) Tegretol (Carbamazepine)	C C		

Date:06/22/05ISR Number: 4698588-9Report Type:Expedited (15-DaCompany Report #2005088825
 Age:72 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Condition Aggravated Convulsion	Consumer	Neurontin (Gabapentin)	PS		ORAL
Other	Head Injury		Gabapentin			

ORAL	Loss Of Consciousness	(Gabapentin)	SS	ORAL
	Pharmaceutical Product Complaint	Dilantin (Phenytoin Sodium)	C	
		All Other Therapeutic Products (All Other Therapeutic Products)	C	

Date:06/22/05ISR Number: 4698590-7Report Type:Expedited (15-DaCompany Report #2005016042

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Aggression
Initial or Prolonged	Bacteraemia
Other	Brain Oedema
	Cerebral Disorder
	Cerebrovascular Accident
	Convulsion
	Depressed Level Of

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Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 900 MG (600 MG, 1/2 TABLET 3 TIMES DAILY), ORAL		Deep Vein Thrombosis	Foreign Health Professional Company Representative	Neurontin (Gabapentin)	PS		ORAL

Domperidone (Domperidone)	C
Mist Carminative (Camphor, Cardamom Oil, Rhizoma Zingiberæ, Tinctura Capsici)	C
Diclofenac	

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Diclofenac Sodium) C

Date:06/22/05ISR Number: 4699316-3Report Type:Expedited (15-DaCompany Report #2005086715
Age:72 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged ORAL	Blood Alkaline Phosphatase Abnormal	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
Other ORAL	Blood Bilirubin Increased C-Reactive Protein	Professional	Tegretol (Carbamazepine)	SS		ORAL
	Increased Eosinophilia Gamma-Glutamyltransferase Abnormal Haemoglobin Decreased Jaundice Pruritus		Glimperide (Glimperide) Insulin Glargine (Insulin Glargine) Indapamide (Indapamide)	C C C		

Date:06/22/05ISR Number: 4699427-2Report Type:Expedited (15-DaCompany Report #ACO_0169_2005
Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 36 MG PO Initial or Prolonged 800 MG PO	Epilepsy Glasgow Coma Scale	Foreign Health	Zanaflex Gabapentin	PS SS		ORAL ORAL
40 MG PO	Abnormal Hallucination Intentional Misuse Loss Of Consciousness Overdose Renal Failure Acute Urinary Tract Infection	Professional Other	Baclofen Ciprofloxacin Bendrofluazide	SS C C		ORAL

Date:06/23/05ISR Number: 4698858-4Report Type:Expedited (15-DaCompany Report #2005080588
Age:68 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cholestasis	Foreign	Neurontin			
Hospitalization - 900 MG (300		Cytolytic Hepatitis	Health	(Gabepentin)	PS		ORAL
Initial or Prolonged MG, 3 IN 1		Disease Recurrence	Professional				
D), ORAL		Pain					
				Triflucan (Flunconazole)	SS		
				Paracetamol (Paracetamol)	C		
				Duragesic (Fentanyl)	C		
				Solu-Medrol (Methylprednisolone			
				Sodium Succinate)	C		
				Mopral (Omeprazole)	C		
				Kardegic (Acetylsalicylate			
				Lysine)	C		
				Lasilix (Furosemide)	C		
				Lopressor (Metoprolol			
				Tartrate)	C		
				Cordarone (Amiodarone			
				Hydrochloride)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Zocor (Simvastatin) C
 Rocephin
 (Ceftriaxone Sodium) C
 Flagyl "Searle"
 (Metronidazole) C
 Actiskenan (Morphine
 Sulfate) C
 Zyloric "Glaxo
 Wellcome"
 (Allopurinol) C
 Rivotril
 (Clonazepam) C
 Oxycontin (Oxycodone
 Hydrochloride) C

Date:06/23/05ISR Number: 4698957-7Report Type:Direct
 Age:70 YR Gender:Female I/FU:I

Company Report #CTU 251837

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 300MG TID		Drug Ineffective		Gabapentene	PS		ORAL
PO				Gabapentene Different Manufacture	SS		ORAL
300MG TID							
PO							

Date:06/23/05ISR Number: 4699381-3Report Type:Direct
 Age:43 YR Gender: I/FU:I

Company Report #CTU 251870

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose 300 MG 1 BID		Muscle Spasms		Gabapentin	PS		ORAL
PO		Pharmaceutical Product Complaint					

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Duration						
Life-Threatening	Cervical Vertebral	Consumer	Neurontin			
Hospitalization -	Fracture		(Gabapentin)	PS		
Initial or Prolonged	Conjunctivitis					
Other	Corneal Ulcer					
	Difficult To Wean From					
	Ventilator					
	Drug Ineffective					
	Dysphagia					
	Ear Canal Injury					
	Fracture					
	Glasgow Coma Scale					
	Abnormal					
	Gun Shot Wound					
	Mastoid Disorder					
	Pneumonia Haemophilus					
	Self Injurious Behaviour					
	Subdural Haematoma					
	Subdural Hygroma					
	Suicide Attempt					

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/23/05ISR Number: 4699820-8Report Type:Expedited (15-DaCompany Report #2005089492

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Attention Deficit/Hyperactivity	Consumer	Lipitor (Atorvastatin)	PS		ORAL
Other 1200 MG (600 MG, 2 IN 1 D), ORAL		Disorder Back Disorder Back Pain		Neurontin (Gabapentin)	SS		ORAL
		Blood Cholesterol Increased		Zoloft (Sertraline)	SS		ORAL
100 MG (100 MG, 1 IN 1 D), ORAL		Fall Heart Rate Increased					
ORAL		Hypertension Migraine		Seroquel (Quetiapine Fumarate)	SS		ORAL
		Post Procedural Complication Surgery Weight Increased		Ambien (Zolpidem Tartrate) Nexium (Esomeprazole)	C C		

Date:06/24/05ISR Number: 4700267-6Report Type:Direct

Company Report #CTU 251930

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 800 MG TID PO		Convulsion Pharmaceutical Product Complaint		Gabapentin-Ndc# 00228263711	PS		ORAL

Date:06/24/05ISR Number: 4700270-6Report Type:Direct

Company Report #CTU 251931

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Convulsion		Gabapentin Ndc#			
		Pharmaceutical Product		00228263711	PS		ORAL
800 MG TID PO		Complaint					

Date:06/24/05ISR Number: 4700284-6Report Type:Direct Company Report #CTU 251934
 Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective		Gabapentin 300 Mg			
				Ivax #4382	PS	Ivax	ORAL
300 MG /DAY							
ORAL							

Date:06/24/05ISR Number: 4701231-3Report Type:Expedited (15-DaCompany Report #2005088786
 Age: Gender:Female I/FU:I

Outcome	PT
Other	Amnesia
	Breast Mass
	Convulsion
	Coordination Abnormal
	Crying
	Depression
	Difficulty In Walking
	Drug Ineffective
	Dysarthria
	Hypoaesthesia

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
ORAL		Mood Swings Pain Pain In Extremity Suicidal Ideation Thinking Abnormal	Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:06/24/05ISR Number: 4701242-8Report Type:Expedited (15-DaCompany Report #2004012957
Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 600 MG (300 Disability MG, 2 IN 1 D) Other		Affective Disorder - Bipolar Disorder Condition Aggravated Dependence Drug Abuser Facial Palsy Gun Shot Wound Injury Overdose Pain Palpitations Scar Skin Injury Suicidal Ideation Suicide Attempt Swelling Tenderness	Consumer	Neurontin (Gabapentin)	PS		

Date:06/27/05ISR Number: 4700560-7Report Type:Direct Company Report #CTU 251964
Age:46 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
300 MG TID PO		Rash Generalised Rash Pruritic		Gabapentin 300 Mg Tid Effexor Xr Lipitor	PS C C		ORAL

Lisinopril

C

Date:06/27/05ISR Number: 4700597-8Report Type:Expedited (15-DaCompany Report #2005090328
Age:55 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Distension Ascites	Foreign Health	Gabapentin (Gabapentin)	PS		ORAL
UNKNOWN (1200 MG), ORAL		Joint Swelling	Professional				
		Oedema Peripheral Rash Swelling Face		Omeprazole (Omeprazole) Diclofenac (Diclofenac Sodium) Tolbutamide (Tolbutamide) Metformin (Metformin) Lisinopril (Lisinopril) Simvastatin (Simvastatin) Cyclizine	C C C C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

(Cyclizine) C
 Stemetil
 (Prochlorperazine) C
 Ibuprofen
 (Ibuprofen) C
 Paracetamol
 (Paracetamol) C
 Co-Codamol (Codeine
 Phosphate,
 Paracetamol) C

Date:06/27/05ISR Number: 4701719-5Report Type:Expedited (15-DaCompany Report #2005077350
 Age:76 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Cardiac Failure	Foreign	Gabapentin			
Other		Condition Aggravated	Consumer				
	900 MG (300 MG, 3 IN 1 D), ORAL		Other	(Gabapentin)	PS		ORAL
				Dextropropoxifen			
				(Dextropropoxyphene)	C		
				Acenocoumarol			
				(Acenocoumarol)	C		ORAL

Date:06/27/05ISR Number: 4701891-7Report Type:Expedited (15-DaCompany Report #2004106748
 Age:29 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abnormal Behaviour Aggression	Consumer	Neurontin (Tablets) (Gabapentin)	PS		
Other	900 MG (300 MG,3 IN 1 D)	Agitation					
		Condition Aggravated		Neurontin			

900 MG (300	Depression	(Gabapentin)	SS
MG,3 IN 1 D)	Drug Ineffective		
	Hallucination, Auditory	Haldol (Haloperidol)	SS
	Homicidal Ideation	Risperidone	
6 MG (3 MG,2	Intentional Misuse	(Risperidone)	SS
IN 1 D)	Lethargy		
	Multiple Drug Overdose	Seroquel (Quetiapine	
	Psychotic Disorder	Fumarate)	C
	Somnolence	Metformin	
	Stress	(Metformin)	C
	Suicidal Ideation		
	Suicide Attempt		

Date:06/27/05ISR Number: 4701968-6Report Type:Expedited (15-DaCompany Report #2004117310
Age:37 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin			
SEE IMAGE		Drug Ineffective		(Gabapentin)	PS		
		Loss Of Consciousness		Ibuprofen			
		Multiple Drug Overdose		(Ibuprofen)	SS		
		Accidental		Amitriptyline			
				Hydrochloride			

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Freedom Of Information (FOI) Report

(Amitriptyline Hydrochloride)	SS
Nortriptyline Hydrochloride (Nortriptyline Hydrochloride)	SS
Venlafaxine (Venlafaxine0)	SS
Oxazepam (Oxazepam)	C
Benzodiazepine Derivatives (Benzodiazepine Derivatives)	C
Opioids (Opioids)	C

Date:06/27/05ISR Number: 4701971-6Report Type:Expedited (15-DaCompany Report #2005090303
Age:65 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Disturbance In Attention	Consumer	Neurontin			
Other		Drug Dose Omission		(Gabapentin)	PS		ORAL
900 MG (300		Fatigue					
MG, 3 IN 1		Herpes Zoster					
D), ORAL		Hostility		Amaryl (Glimepiride)	C		
		Memory Impairment		Warfarin 9warfarin0	C		
		Mobility Decreased		Glucophage (Metformin Hydrochloride0	C		
				Prinzide (Hydrochlorothiazide , Lisinopril)	C		
				Atenolo (Atenolol)	C		
				Acetylsalicylic Acid 9acetylsalicylic Acid)	C		

Date:06/27/05ISR Number: 4701994-7Report Type:Expedited (15-DaCompany Report #2004077855
Age:14 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abnormal Behaviour Aggression	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG		Coma					
(UNKNOWN),		Concussion					
ORAL		Developmental Delay		Levothroid			
		Drug Ineffective		(Levothyroxine Sodium)	C		
		Dysarthria		Clindamycin			
		Gait Disturbance		(Clindamycin)	C		
		Hallucination, Auditory		Seroquel (Quetiapine Fumarate)	C		
		Loss Of Consciousness		Lithium (Lithium)	C		
		Nightmare		Multivitamins			
		Pain		(Multivitamins)	C		
		Personality Change		Paxil (Paroxetine Hydrochloride)	C		
		Polytraumatism		Clozaril (Clozapine)	C		
		Suicide Attempt		Ativan (Lorazepam)	C		
				Colace (Docusate Sodium)	C		

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Freedom Of Information (FOI) Report

Provigil (Modafinil) C
 Topamax (Topiramate) C
 Naltrexone
 (Naltrexone) C
 Depo-Provera 150
 (Medroxyprogesterone
 Acetate) C
 Albuterol
 (Salbutamol) C
 All Other
 Therapeutic Products
 (All Other
 Therapeutic
 Products) C

Date:06/27/05ISR Number: 4702038-3Report Type:Expedited (15-DaCompany Report #2005079867
 Age:51 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 300 MG (300 Other MG, 1 IN 1 D), ORAL		Chest Pain Drug Ineffective Dysphemia Dyspnoea Fatigue Headache Hypoaesthesia Memory Impairment Muscle Twitching Speech Disorder Weight Decreased	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Zoloft (Sertraline0 All Other Therapeutic Products (All Othertherapeutic Products)	C		C

Date:06/27/05ISR Number: 4702040-1Report Type:Expedited (15-DaCompany Report #2005063311
 Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 900 MG (900		Asthenia Drug Ineffective	Consumer Health	Neurontin (Gabapentin)	PS		ORAL

MG, 1 IN 1	Fatigue	Professional		
D), ORAL	Pharmaceutical Product			
900 MG (900	Complaint		Gabapentin	
MG, 1 IN 1	Sleep Apnoea Syndrome		(Gabapentin)	SS
D), ORAL	Somnolence			ORAL
	Weight Decreased			
			Glucotrol	
			(Glipizide)	C
			Glucophage	
			(Metformin	
			Hydrochloride)	C
			Synthroid	
			(Levothyroxine	
			Sodium)	C
			Provera	
			(Medroxyprogesterone	
			Acetate)	C
			Estrace (Estradiol)	C
			Folic Acid (Folic	
			Acid)	C
			Methotrexate	
			(Methotrexate)	C

Freedom Of Information (FOI) Report

Multivitamins (Multivitamins)	C
Xalatan (Latanprost)	C
Alprazolam (Alprazolam)	C
Lomotil (Atropine Sulfate, Diphenoxylate Hydrochloride)	C
Dilaudid (Hydromorhone Hydrochloride)	C
Acetaminophen W/Codeine (Codeine, Paracetamol)	C
Propoxyphene Hcl And Acetaminophen (Dextropropoxyphene Hydrochloride, Paracetamol)	C
Entex Pse (Guaifenesin, Pseudoephedrine Hydrochloride)	C
Vicodin Es (Hydrocodone Bitartrate, Paracetamol)	C

Date:06/27/05ISR Number: 4702191-1Report Type:Expedited (15-DaCompany Report #2005083192
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebral Disorder	Consumer	Neurontin			
		Cerebrovascular Accident		(Gabapentin)	PS		
		Loss Of Consciousness		Gabapentin			
		Pharmaceutical Product		(Gabapentin)	SS		
		Complaint					

Date:06/28/05ISR Number: 4701367-7Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20050507147
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	800 mg, twice	Completed Suicide		Motrin 800 Mg	PS		
	to three	Loss Of Consciousness					
	times, as	Multiple Drug Overdose					
	needed	Accidental					
				Nortriptyline			
				Hydrochloride	SS		
				Amitriptyline			
				Hydrochloride	SS		
				Venlafaxine	SS		
				Neurontin	SS		
				Neurontin	SS		
				Neurontin	SS		
				Oxazepam	C		
				Benzodiazepine			
				Derivatives	C		
				Opioids	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/28/05ISR Number: 4703009-3Report Type:Direct
 Age:75 YR Gender:Female I/FU:I

Company Report #CTU 252090

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Unevaluable Event		Neurontin	PS		

Date:06/28/05ISR Number: 4703725-3Report Type:Expedited (15-DaCompany Report #2005072171
 Age:50 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Angioneurotic Oedema	Consumer	Celebrex (Celecoxib)	PS		ORAL
200 MG (200		Diarrhoea	Health				
MG, DAILY),		Disease Progression	Professional				
ORAL		Drug Ineffective		Neurontin			
		Fibromyalgia		(Gabapentin)	SS		
10 MG (10 MG,		Gastrointestinal Pain		Bextra (Valdecoxib)	SS		ORAL
1 IN 1 D),		Hip Arthroplasty					
ORAL		Inflammation					
		Musculoskeletal Disorder		Ibuprofen			
2400 MG (800		Osteoarthritis		(Ibuprofen)	SS		
MG, 3 IN 1 D)		Pain					
				Oxycodone			
				Hydrochloride			
				(Oxycodone			
				Hydrochloride)	C		
				Oxycocet (Oxycodone			
				Hydrochloride,			
				Paracetamol)	C		
				Sertraline			
				Hydrochloride			
				(Sertraline			
				Hydrochloride)	C		

Freedom Of Information (FOI) Report

Wellbutrin
 (Bupropion
 Hydrochloride) C
 Seroquel (Quetiapine
 Fumarate) C
 Aricept (Donepezil) C

Date:06/29/05ISR Number: 4703775-7Report Type:Direct
 Age: Gender:Female I/FU:I

Company Report #CTU 252242

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Effect Decreased		Gabapentin 300 Mg	PS		ORAL
300 MG 3 CAPS		Drug Intolerance					
BID PO		Pharmaceutical Product Complaint					

Date:06/29/05ISR Number: 4704585-7Report Type:Expedited (15-DaCompany Report #2005081724
 Age:63 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Abdominal Pain Upper	Foreign	Gabapentin	PS		ORAL
1200 MG (3 IN		Anorexia	Health				
Initial or Prolonged		Back Pain	Professional	Doxazosin			
1 D), ORAL		Condition Aggravated	Company	(Doxazosin)	C		
		Constipation	Representative	Antihypertensives			
		Dysphonia	Other				
		Headache					
		Muscular Weakness					
		Osteochondrosis		(Antihypertensives)	C		
		Productive Cough					
		Protein Urine Present					
		Pyrexia					
		Red Blood Cells Urine					
		Positive					
		Urine Analysis Abnormal					

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Fatigue Imprisonment	Health Professional	Neurontin (Gabapentin)	PS		
1800 MG (300 MG, 6 IN 1 D) 2 MG		Legal Problem					
		Loss Of Consciousness Road Traffic Accident		Xanax Tablet (Alprazolam)	SS		
				Desoxyn (Metamfetamine Hydrochloride)	SS		ORAL
180 MG (30 MG, 6 IN 1 D), ORAL							
40 MG 1 D				Prilosec (Omeprazole)	SS		
480 MG (80 MG, 6 IN 1 D)				Oxycontin (Oxycodone Hydrochloride)	SS		
200 MG, 1 IN 3 D				Duragesic (Fentanyl)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:06/29/05ISR Number: 4705089-8Report Type:Expedited (15-DaCompany Report #2005090547

Age:79 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Cholesterol Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Cardiac Disorder		Digoxin (Digoxin)	C		
		Cardiac Pacemaker Replacement		Prinivil (Lisinopril)	C		
		Chest Pain		Synthroid (Levothyroxine Sodium)	C		
		Cough		Toprol (Metoprolol)	C		
		Drug Dose Omission		Tamsulosin (Tamsulosin)	C		
		Implantable Defibrillator Insertion		Lasix (Furosemide)	C		
		Medical Device Complication		Plavix (Clopidogrel Sulfate)	C		
		Pneumonia		Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
		Therapy Non-Responder		Nitroglycerin (Nitroglycerin)	C		
		Treatment Noncompliance					
		Vascular Occlusion					

Date:06/29/05ISR Number: 4705247-2Report Type:Expedited (15-DaCompany Report #2005AL002272

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Asterixis Coordination Abnormal	Literature Health	Gabapentin Capsules, 300 Mg (Purepac)	PS		ORAL
300 MG ; BID;		Fall	Professional				
PO		Gait Disturbance		Glyburide	C		
		Myoclonus		Metoprolol	C		
		Off Label Use		Hydrochlorothiazide	C		
				W/ Losartan	C		
				Fentanyl	C		
				Omeprazole	C		
				Docusate Sodum	C		
				Bisacodyl	C		
				Epoetin Alfa	C		

Sucralfate

C

Date:06/29/05ISR Number: 4705697-4Report Type:Expedited (15-DaCompany Report #2005092127
Age:52 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Blood Creatine Phosphokinase Increased	Foreign Literature	Gabapentin (Gabapentin)	PS		
100 MG (100 MG, 1 IN 1 D)		Blood Creatinine Increased Hypoaesthesia Myopathy	Health Professional	Ranitidine (Ranitidine Hydrochloride) Clonazepam (Clonazepam) Sertraline (Sertraline) Erythropoietin (Erythropoietin) Vitamin B6 (Vitamin B6)	C C C C C		

Freedom Of Information (FOI) Report

Multivitamins
(Multivitamins) C

Date:06/29/05ISR Number: 4705892-4Report Type:Expedited (15-DaCompany Report #2005092140
Age:43 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Blood Creatine Phosphokinase Increased	Foreign Literature	Gabapentin (Gabapentin)	PS		
100 MG (100 MG, 1 IN 1 D)		Blood Creatinine Increased Eye Disorder Hypoaesthesia Oral Myopathy	Health Professional	Amlodipine (Amlodipine) Atorvastatin (Atorvastatin) Rocaltrol (Calcitriol) Multivitamins (Multivitamins) Zopiclone (Zopiclone) Erythropoietin (Erythropoietin) Iron Sucrose (Saccharated Iron Oxide)	C C C C C C		

Date:06/29/05ISR Number: 4705936-XReport Type:Expedited (15-DaCompany Report #2005092295
Age:35 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Anxiety Disorder Drug Ineffective Intentional Misuse Overdose Physical Assault Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:06/29/05ISR Number: 4706105-XReport Type:Expedited (15-DaCompany Report #2005092341
Age:31 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Carbon Monoxide Poisoning Drug Ineffective Suicide Attempt	Consumer	Neurontin (Gabapentin) Carbon Monoxide (Carbon Monoxide)	PS SS		

RESPIRATORY

(INHALATION) INHALATION

Date:06/30/05ISR Number: 4704599-7Report Type:Expedited (15-DaCompany Report #2005091715
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Drug Tolerance Increased	Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL		Gastric Bypass Protein Urine Present Renal Disorder		Micardis (Telmisartan) Klonopin (Clonazepam)	C C		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Prozac (Fluoxetine
Hydrochloride) C
Proscar
(Finasteride) C
Flomax "Boehringer
Ingelheim"
(Tamsulosin
Hydrochloride) C

Date:06/30/05ISR Number: 4704657-7Report Type:Expedited (15-DaCompany Report #2005083192
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Cerebrovascular Accident	Consumer	Neurontin (Gabapentin)	PS		
		Loss Of Consciousness		Gabapentin			
		Nervous System Disorder		(Gabapentin)	SS		
		Pharmaceutical Product					
		Complaint					

Date:06/30/05ISR Number: 4704755-8Report Type:Expedited (15-DaCompany Report #KII-2005-0017140
Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Alanine Aminotransferase Increased	Study Health	Morphine Sulfate (Similar To 19-516)	PS		ORAL
Other		Aspartate	Professional	(Morphine Sulfate)			
ORAL		Aminotransferase Increased	Other	Amitriptyline (Amitriptyline)	SS		
		Blood Creatinine Increased		Acetaminophen W/Hydrocodone			
		Blood Ph Decreased		Bitartrate			
		Chromaturia		(Paracetamol, Hydrocodone)	SS		
ORAL		Coma					
		Hypotension		Carisoprodol			
		Multiple Drug Overdose		(Carisoprodol)	SS		ORAL
ORAL		Oxygen Saturation		Gabapentin			

ORAL	Decreased	(Gabapentin)	SS	ORAL
	Pneumonia Aspiration	Antiinflammatory/Ant		
	Respiratory Depression	irheumatic Products	SS	
	Sinus Tachycardia	Acetylsalicylic Acid		
	Urine Analysis Abnormal	(Acetylsalicylic		
	Vomiting	Acid)	SS	

Date:07/01/05ISR Number: 4706952-4Report Type:Expedited (15-DaCompany Report #2005090778
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Abdominal Pain Upper Aphasia	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (200 MG, 3 IN 1 D), ORAL		Cerebrovascular Accident					
		Foaming At Mouth					
800 MG (200 MG, 3 IN 1 D), ORAL		Hepatic Pain Pharmaceutical Product		Gabapentin (Gabapentin)	SS		ORAL
		Complaint					
		Pruritus					
		Tremor		Oxycontin (Oxycodone Hydrochloride) Clinoril (Sulindac)	C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Lortab (Hydrocodone Bitartrate, Paracetamol) C

Date:07/01/05ISR Number: 4707106-8Report Type:Expedited (15-DaCompany Report #2005-DE-02336GD
Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Drug Toxicity Polysubstance Abuse	Literature	Paracetamol (Paracetamol)	PS		
				Oxazepam(Oxazepam)	SS		
				Alprazolam	SS		
				Oxycodone (Oxycodone)	SS		
				Propoxyphene (Dextropropoxyphene)	SS		
				Diphenhydramine (Diphenhydramine)	SS		
				Temazepam (Temazepam)	SS		
				Gabapentin	SS		
				Metoprolol (Metoprolol)	SS		

Date:07/01/05ISR Number: 4707107-XReport Type:Expedited (15-DaCompany Report #2005056759
Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged (400 MG),		Drug Ineffective Irritable Bowel Syndrome Serotonin Syndrome	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL
ORAL							

Date:07/05/05ISR Number: 4706787-2Report Type:Expedited (15-DaCompany Report #L05-USA-02220-01
Age:54 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Death	Completed Suicide	Literature	Citalopram	PS
	Drug Abuser	Health	Metaxalone	SS
	Drug Toxicity	Professional	Gabapentin	SS
TABLET				
	Intentional Misuse		Acetaminophen	SS
	Overdose		Flurazepam	SS

Date:07/05/05ISR Number: 4706803-8Report Type:Expedited (15-DaCompany Report #2005092344

Age:38 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		Carbon Monoxide Poisoning	Consumer	Neurontin			
Other		Drug Ineffective		(Gabapentin)	PS		
		Injury		Carbon Monoxide			
		Suicide Attempt		(Carbon Monoxide)	SS		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/05/05ISR Number: 4706806-3Report Type:Expedited (15-DaCompany Report #2005092334
Age:49 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		

Date:07/05/05ISR Number: 4706807-5Report Type:Expedited (15-DaCompany Report #2005092310
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Ineffective Injury Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:07/05/05ISR Number: 4706809-9Report Type:Expedited (15-DaCompany Report #2005091971
Age:79 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Blood Triglycerides Increased	Consumer	Lipitor (Atorvastatin)	PS		ORAL
(1 IN 1 D), ORAL		Carotid Artery Occlusion					
300 MG (100 MG, 3 IN 1 D), ORAL		Coronary Artery Occlusion Suicidal Ideation		Neurontin (Gabapentin)	SS		ORAL

Asa (Acetylsalicylic Acid)	C
Furosemide	C
Alprazolam	C
Protonix (Pantoprazole)	C
Klor-Con (Potassium Chloride)	C
Lotensin (Benazepril	

Hydrochloride) C
Zantac (Ranitidine
Hydrochloride) C
Propranolol C
Folic Acid C
Multivitamins C
Calcium C
Famotidine C
Detrol (Tolterodine
L-Tartrate) C
Tamoxifen C
Tiazac (Diltiazem
Hydrochloride) C

Date:07/05/05ISR Number: 4706812-9Report Type:Expedited (15-DaCompany Report #2005070434
Age:51 YR Gender:Female I/FU:F

Outcome PT
Hospitalization - Agoraphobia
Initial or Prolonged Blood Cholesterol
Disability Increased
Other Body Height Decreased
Cerebrovascular Accident
Convulsion

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route	
300 MG (100 MG, 3 IN 1 D)		Disturbance In Attention	Consumer Health	Neurontin (Gabapentin)	PS			
		Economic Problem						
		Insomnia	Professional					
		Intervertebral Disc Degeneration						
		Marital Problem						
		Memory Impairment			Premarin (Estrogens Conjugated)	C		
		Mental Disorder			Prozac (Fluoxetine Hydrochloride)	C		
		Nervous System Disorder			Seroquel	C		
		Nervousness						
		Osteoarthritis						
Panic Attack								
Treatment Noncompliance								
Tremor								

Date:07/05/05ISR Number: 4706821-XReport Type:Expedited (15-DaCompany Report #2002065565

Age:55 YR Gender: I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death	ORAL	Completed Suicide	Literature Consumer	Gabapentin (Gabapentin)	PS		
OPHTHALMIC		Intentional Misuse					
ORAL		Overdose	Health Professional	Diltiazem (Diltiazem)	SS		ORAL
ORAL				Atenolol (Atenolol)	SS		ORAL

Date:07/05/05ISR Number: 4707673-4Report Type:Expedited (15-DaCompany Report #2005083952

Age:47 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability	800 MG (400 MG 2 IN 1	Amnesia	Foreign Health	Neurontin (Tablets) (Gabapentin)	PS		
Other		Apathy					
OPHTHALMIC		Asthenia	Professional				

D)ORAL

Balance Disorder

Fatigue
Somnolence

Neurobion
(Cyancobalamin,
Pyridoxine
Hydrochloride,
Thiamine

C

Date:07/05/05ISR Number: 4707679-5Report Type:Expedited (15-DaCompany Report #2005091119

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Neutropenia	Foreign Health Professional	Neurontin (Gabapentin) Rivotril (Clonazepam) Orocal (Calcium Carbonate)	PS C C		ORAL

Date:07/05/05ISR Number: 4707949-0Report Type:Direct

Company Report #CTU 252450

Age:43 YR Gender: I/FU:I

Outcome

PT

Muscle Spasms

Therapeutic Response

Unexpected With Drug

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Substitution

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
300 MG TID PO			Gabapentin	PS		ORAL

Date:07/05/05ISR Number: 4708174-XReport Type:Direct
 Age: Gender:Female I/FU:I Company Report #CTU 252466

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		No Adverse Effect		Neurontin 100 Mg	PS		
100 MG				Prevacid 30 Mg	SS		
30 MG							

Date:07/05/05ISR Number: 4708194-5Report Type:Direct
 Age: Gender: I/FU:I Company Report #CTU 252511

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose		No Adverse Effect		Gabapentin (Generic Of Neurontin)	PS		
100 MG 3X./D							

Date:07/05/05ISR Number: 4708426-3Report Type:Expedited (15-DaCompany Report #2005068211
 Age:86 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Flatulence	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
900 MG (300 Other MG, 3 IN 1 D)		Malaise	Professional				
		Oral Intake Reduced					
ORAL		Pharmaceutical Product					
		Complaint		Gabapentin (Gabapentin)	SS		ORAL
1800 MG (600		Weight Decreased					

ORAL

Bextra (Valdecoxib)	SS
Methadone Hydrochloride (Methadone Hydrochloride)	SS
Isosorbide	C
Lisinopril	C
Furosemide	C
Potassium	C
Warfarin	C
Terazosin	C
Nifedipine	C
All Other Therapeutic Products	C
Simvastatin	C

Date:07/05/05ISR Number: 4708973-4Report Type:Expedited (15-DaCompany Report #2005091349
 Age:75 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration						
Hospitalization - Initial or Prolonged	Atrial Fibrillation	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL	Biliary Cirrhosis					
Other	Cardiac Failure	Professional	Nifedipine	SS		ORAL
ORAL	Lung Disorder		Kardegic (Acetylsalicylate Lysine)	C		
	Lymphadenopathy					

Freedom Of Information (FOI) Report

Hyperium
(Rilmenidine) C

Date:07/05/05ISR Number: 4709104-7Report Type:Expedited (15-DaCompany Report #2005019428
Age:66 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Atrial Fibrillation	Consumer	Lipitor	PS		ORAL
Other		Headache		(Atorvastatin)			
10 MG (10 MG,		Malaise					
1 IN 1 D),		Rash Generalised					
ORAL		Therapy Non-Responder		Neurontin	SS		ORAL
600 MG (1 IN		Treatment Noncompliance		(Gabapentin)			
1 D), ORAL		Trigeminal Neuralgia					
50 MG (1 IN 1				Atenolol (Atenolol)	SS		ORAL
D), ORAL							
200 MG (1 IN				Tegretol	SS		ORAL
1 D), ORAL				(Carbamazepine)			
				Lisinopril	C		
				(Lisinopril)			
				Hydrochlorothiazide	C		
				(Hydrochlorothiazide			
)			

Date:07/05/05ISR Number: 4709106-0Report Type:Expedited (15-DaCompany Report #2005048449
Age:46 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Aggression	Consumer	Neurontin	PS		
Other		Cervical Vertebral		(Gabapentin)			
900 MG (300							

MG, 3 IN 1	Fracture		
D),	Completed Suicide		
100 MG (100	Drug Ineffective	Diclofenac	
MG, 1 IN 1	Haemorrhage	(Diclofenac Sodium)	SS
D),	Intentional Self-Injury		
	Petechiae		
		Diflucan	
		(Fluconazole)	C
		Diphenoxylate	
		(Diphenoxylate)	C
		Propoxyphene	
		Napsylate	
		(Dextropropoxyphene	
		Napsylate)	C
		Levothyroxine	
		(Levothyroxine)	C
		All Other	
		Therapeutic Products	
		(All Other	
		Therapeutic	
		Products)	C
		Glucosamine	
		(Glucosamine)	C
		Chondroitin	
		(Chondroitin	
		Sulfate)	C
		Relafen (Nabumetone)	C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/05/05ISR Number: 4709108-4Report Type:Expedited (15-DaCompany Report #2005071563
 Age:6 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 400 MG, ORAL	Arthralgia Bipolar Disorder	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
Other	Convulsion	Professional	Lamictal			
	Coordination Abnormal		(Lamotrigine)	SS		
	Difficulty In Walking		Zyprexa (Olanzapine)	SS		
	Disease Recurrence		Seroquel (Quetiapine			
	Dizziness		Fumarate)	SS		
	Drug Ineffective		Depakote (Valproate			
	Increased Appetite		Semisodium)	SS		
	Insomnia		Depakene (Valproate			
	Mania		Sodium)	C		
	Metabolic Disorder		Claritin			
	Migraine		(Loratadine)	C		
	Speech Disorder					
	Suicidal Ideation					
	Weight Increased					

Date:07/05/05ISR Number: 4709162-XReport Type:Expedited (15-DaCompany Report #2005092301
 Age:45 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	Drug Ineffective Intentional Misuse Overdose Suicide Attempt	Consumer	Neurontin (Gabapentin) Antidepressants (Antidepressants)	PS SS		

Date:07/05/05ISR Number: 4709188-6Report Type:Expedited (15-DaCompany Report #2005092327
 Age:63 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Death Other	Completed Suicide Depression Drug Ineffective Insomnia	Consumer	Neurontin (Gabapentin)	PS		

Date:07/05/05ISR Number: 4709191-6Report Type:Expedited (15-DaCompany Report #2005071384
Age: Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Condition Aggravated Diabetic Neuropathy	Consumer	Neurontin (Gabapentin)	PS		
UNKNOWN	300 MG	(100 Hip Arthroplasty					
MG, 3 IN 1							
D); UNKNOWN							
				Procardia (Nifedipine)	C		
				Lipitor (Atorvastatin)	C		

Date:07/06/05ISR Number: 4708557-8Report Type:Expedited (15-DaCompany Report #2005080957
Age:44 YR Gender:Female I/FU:F

Outcome	PT
Other	Blood Cholesterol 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
400 MG (200 MG, 2 IN 1 D), ORAL		Gastric Ulcer Hypertension Intervertebral Disc Protrusion	Consumer	Celebrex (Celecoxib)	PS		ORAL
				Neurontin (Gabapentin)	SS		
				Accupril (Quinapril Hydrochloride)	SS		
				Ambien (Zolpidem Tartrate)	C		
				Effexor (Venlafaxine Hydrochloride)	C		
				Oxycontin (Oxycodone Hydrochloride)	C		
				Maxzide (Hydrochlorothiazide, Triamterene)	C		
				Potassium (Potassium)	C		

Date:07/06/05ISR Number: 4708581-5Report Type:Expedited (15-DaCompany Report #2005093793

Age: Gender:Male I/FU:I

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - ORAL	Initial or Prolonged	Abdominal Pain Upper	Consumer	Celebrex (Celecoxib)	PS		ORAL
40 MG (20 MG, Other 1 IN 1 D)		Drug Effect Decreased		Bextra (Valdecoxib)	SS		ORAL
ORAL		Hernia Repair					
		Nephrolithiasis					
3600 MG(600 MG, 1200 MG		Post Procedural Complication		Neurontin (Gabapentin)	SS		ORAL
		Skin Infection					

TID) ORAL

Urinary Hesitation

Urinary Retention

Cyclobenzaprine	C
Lexapro	
(Escitalopram)	C
Amitriptyline	C
Doxazosin	C
Diclofenac	
(Diclofenac Sodium)	C
Oxycontin (Oxycodone	
Hydrochloride)	C

Date:07/07/05ISR Number: 4708944-8Report Type:Expedited (15-DaCompany Report #2005075005

Age: Gender:Female I/FU:F

Outcome	PT
Disability	Amnesia
Other	Back Pain
	Blood Alkaline
	Phosphatase Increased
	Blood Cholesterol
	Increased
	Confusional State
	Difficulty In Walking
	Drug Abuser
	Exercise Capacity
	Decreased

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Eye Disorder Fluid Retention Foot Fracture					
1.5 MG (0.5 MG, 3 IN 1 D), ORAL		High Density Lipoprotein Decreased	Consumer Health	Xanax Tablet (Alproazolam)	PS		ORAL
		Intervertebral Disc Disorder	Professional				
500 MG (100 MG, 3 IN 1 D), ORAL		Memory Impairment Oedema Peripheral Pain In Extremity Vitreous Detachment		Neurontin (Gabapentin)	SS		ORAL
		Weight Increased Weight Loss Poor		Fioricet (Butalbital, Caffeine, Paracetamol) Avapro (Irbesartan) Prevacid (Lansoprazole0	SS C C		

Date:07/07/05ISR Number: 4710173-9Report Type:Expedited (15-DaCompany Report #2005080812
Age:71 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Drug Ineffective Knee Arthroplasty	Consumer Health	Neurontin (Tablets) (Gabapentin)	PS		ORAL
2400 MG (200 MG, 3 IN 1 D) ORAL			Professional				
400 MG (200 MG, 2 IN 1 D) ORAL				Celebrex (Celecoxib)	SS		ORAL
				Lantus (Insulin)			

Glargine)	C
Humalog (Insulin Lispro)	C
Humulin R (Insulin Human)	C
Norvasc (Amlodipine)	C
Sinequan (Doxepin)	C

Date:07/07/05ISR Number: 4710491-4Report Type:Expedited (15-DaCompany Report #2005049905
 Age:32 YR Gender:Female I/FU:F

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
800 MG (400 MG, 2 IN 1 D),		Complex Partial Seizures Drug Interaction	Consumer	Neurontin (Gabapentin)	PS		
		Petit Mal Epilepsy					
		Pharmaceutical Product					
		Complaint		Astelina (Azelastine Hydrochloride)	SS		
1000 MG (500 MG, 2 IN 1 D),				Tegretol (Carbamazepine)	SS		
				Ativan (Lorazepam)	C		

Freedom Of Information (FOI) Report

Date:07/07/05ISR Number: 4710509-9Report Type:Expedited (15-DaCompany Report #2005081724
 Age:63 YR Gender:Male I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (3 IN 1 D), ORAL	Abdominal Pain Upper Anorexia Back Pain	Foreign Health Professional	Gabapentin (Tablets) (Gabapentin)	PS		ORAL
	Constipation Dysphonia Haemoglobin Increased Headache Muscular Weakness Osteochondrosis Productive Cough Protein Urine Present Pyrexia Red Blood Cells Urine Positive White Blood Cells Urine Positive	Company Representative	Doxazosin (Doxazosin) Antihypertensives (Antihypertensives)	C C		

Date:07/07/05ISR Number: 4710516-6Report Type:Expedited (15-DaCompany Report #2005086982
 Age:15 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 900 MG (300 Other MG,)	Confusional State Convulsion Drug Interaction	Foreign Health Professional	Neurontin (Gabapentin)	PS		
	Drug Level Above Therapeutic Drug Toxicity	Company Representative	Tramadol (Tramadol) Amitriptyline (Amitriptyline) Clonidine (Clonidine)	SS C C		

Date:07/08/05ISR Number: 4710184-3Report Type:Direct
 Age:52 YR Gender:Male I/FU:I

Company Report #CTU 252789

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening		Back Injury		Neurontin	PS		
600 MG							
Hospitalization -		Convulsion		Bextra	SS		
300 MG							
Initial or Prolonged		Joint Injury		Vioxx	SS		
Disability		Limb Injury		Theraphyin	C		
Other		Road Traffic Accident					

Date:07/08/05ISR Number: 4711040-7Report Type:Expedited (15-DaCompany Report #2005094694
Age:64 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Renal Pain	Foreign	Neurontin			
		Urethral Stricture	Consumer	(Gabapentin)	PS		ORAL
900 MG (300							
MG, 3 IN 1		Urinary Retention					
D), ORAL							
				Ciprofloxacin			
				(Ciprofloxacin)	C		
				Dexibuprofen			
				(Dexibuprofen)	C		

Feeling Cold
Gastrooesophageal Reflux
Disease
Gingival Disorder
Glossodynia
Hostility
Hypersomnia
Hypokinesia
Hyporeflexia
Impaired Healing
Injury
Irritability
Limb Injury
Muscle Spasms
Myalgia
Neck Pain
Oesophageal Spasm
Overdose
Pain In Extremity
Pain In Jaw

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Sedation Sleep Disorder Somnolence					
1800 MG (600 MG, 3 IN 1 D); ORAL		Tenderness Therapy Non-Responder	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Tooth Abscess	Professional				
		Treatment Noncompliance					
20 (20MG, 1 IN 1 D); ORAL		Tremor Vertigo		Lipitor (Atorvastatin)	SS		ORAL
				Lithium (Lithium)	SS		
				Naproxen (Naproxen)	SS		
1000 MG (1 IN 2 D); ORAL				Amoxicillin (Amoxicillin)	SS		ORAL
				Rofecoxib (Rofecoxib)	SS		
				All Other Therapeutic Products	C		
				Levothyroxine Sodium	C		
				Vitamins (Vitamins)	C		
				Diltiazem Hydrochloride	C		

Date:07/08/05ISR Number: 4711359-XReport Type:Expedited (15-DaCompany Report #2005094528
Age:40 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Drug Ineffective Injury Intentional Misuse Multiple Drug Overdose Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		
				All Other Therapeutic Products (All Other Therapeutic Products)	SS		

Date:07/08/05ISR Number: 4711361-8Report Type:Expedited (15-DaCompany Report #2005094556
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin			
		Intentional Misuse		(Gabapentin)	PS		
		Overdose		Xanax Tablet			
		Suicide Attempt		(Alprazolam)	SS		

Date:07/08/05ISR Number: 4711362-XReport Type:Expedited (15-DaCompany Report #2005094661
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Carbon Monoxide Poisoning	Consumer	Neurontin			
		Drug Ineffective		(Gabapentin)	PS		
		Suicide Attempt		Carbon Monoxide			
				(Carbon Monoxide)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/08/05ISR Number: 4711685-4Report Type:Expedited (15-DaCompany Report #2005095535

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other	2700 MG (2700 MG, 1 IN 1 D), UNKNOWN	Retinal Detachment	Health Professional Company Representative	Neurontin (Gabapentin)	PS		

Date:07/11/05ISR Number: 4712013-0Report Type:Direct Company Report #CTU 252966

Age:33 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Life-Threatening Hospitalization - 300MG 3 Initial or Prolonged TIMES A DAY ORAL		Asthenia Convulsion Dyskinesia Fall Head Injury Paraesthesia Speech Disorder		Neurontin 300mg Purepac	PS	Purepac	ORAL

Date:07/11/05ISR Number: 4712324-9Report Type:Expedited (15-DaCompany Report #2004031173

Age:20 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Disability ORAL Other 80 MG (40 MG, 2 IN 1 D), ORAL		Anhedonia Anxiety Completed Suicide Drug Toxicity Emotional Distress	Consumer Health Professional	Neurontin (Gabapentin) Geodon (Ziprasidone)	PS SS		ORAL ORAL

60 MG (30 MG, 2 IN 1 D), ORAL	Incorrect Dose Administered Mental Disorder Pain Pharmaceutical Product Complaint Polytraumatism Sudden Death	Acetaminophen (Paracetamol) Paxil (Paroxetine Hydrochloride)	SS SS	ORAL
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Date:07/11/05ISR Number: 4712329-8Report Type:Expedited (15-DaCompany Report #2005094668
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Ineffective Injury Asphyxiation	Consumer	Neurontin (Gabapentin)	PS		

Date:07/11/05ISR Number: 4712341-9Report Type:Expedited (15-DaCompany Report #2004105644
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged Other		Coma Dependence Post-Traumatic Stress Disorder Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/11/05ISR Number: 4712343-2Report Type:Expedited (15-DaCompany Report #2005094535
 Age:44 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin			
Life-Threatening		Drug Ineffective		(Gabapentin)	PS		
Other		Intentional Misuse					
		Multiple Drug Overdose					

Date:07/11/05ISR Number: 4712457-7Report Type:Expedited (15-DaCompany Report #2005086719
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Fatigue	Health	Xanax Tablet			
		Loss Of Consciousness	Professional	(Alprazolam)	PS		
2 MG		Murder		Neurontin			
		Road Traffic Accident		(Gabapentin)	SS		
1800 MG (300							
MG, 6 IN 1 D)							
				Desoxyn			
				(Metamfetamine			
				Hydrochloride)	SS		ORAL
180 MG (30							
MG, 6 IN 1							
D), ORAL	18	MON					
				Duragesic (Fentanyl)	SS		
200 MCG, 1 IN							
3 D							
				Prilosec			
				(Omeprazole)	SS		
40 MG (40 MG,							
1 IN 1 D)							
				Oxycontin (Oxycodone			
				Hydrochloride)	SS		
480 MG (80							
MG, 6 IN 1 D)							

Date:07/11/05ISR Number: 4712508-XReport Type:Expedited (15-DaCompany Report #2005094545
Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Intentional Self-Injury Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:07/11/05ISR Number: 4712880-0Report Type:Expedited (15-DaCompany Report #2005087113
Age:11 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization - Initial or Prolonged		Crying Fear	Foreign Health	Neurontin (Gabapentin)	PS		
900 MG (300 MG, 3 IN 1 D)		Hallucination, Visual	Professional				
		Hypertonia Middle Insomnia Pallor Pupil Fixed	Company Representative	Epitomax (Topiramate) Urbanyl (Clobazam)	C C		

Date:07/11/05ISR Number: 4713515-3Report Type:Expedited (15-DaCompany Report #2005080588
Age:68 YR Gender:Male I/FU:F

Outcome	PT
Death	Cholestasis
Hospitalization - Initial or Prolonged	Cytolytic Hepatitis Disease Recurrence

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Neuropathic Pain Overdose	Report Source	Product	Role	Manufacturer	Route
900 MG (300	MG,3 IN 1		Foreign Health	Neurontin (Gabapentin)	PS		ORAL
D),ORAL			Professional	Triflucan (Fluconazole)	SS		
				Paracetamol (Paracetamol)	C		
				Duragesic (Fentanyl)	C		
				Solu-Medrol (Methylprednisolone			
				Sodium Succinate)	C		
				Mopral (Omeprazole)	C		
				Kardegic (Acetylsalicylate			
				Lysine)	C		
				Lasilix (Furosemide)	C		
				Lopressor (Metoprolol			
				Tartrate)	C		
				Cordarone (Amiodarone			
				Hydrochloride)	C		
				Zocor (Simvastatin)	C		
				Rocephin (Ceftriaxone Sodium)	C		
				Flagyl "Searle"			
				(Metronidazole)	C		
				Actiskenan (Morphine			
				Sulfate)	C		
				Zyloric "Glaxo Wellcome"			
				(Allopurinol)	C		
				Rivotril			
				(Clonazepam)	C		
				Oxycontin (Oxycodone			
				Hydrochloride)	C		

Date:07/11/05ISR Number: 4713614-6Report Type:Expedited (15-DaCompany Report #2005091119
Age:54 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged ORAL	Neutropenia	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Professional	Rivotril (Clonazepam)	C		
			Orocal (Calcium Carbonate)	C		

Date:07/12/05ISR Number: 4712134-2Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20050601494
Age: Gender:Female I/FU:F

Outcome	PT
Other	Back Pain Brain Damage Drug Interaction

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Headache Intervertebral Disc Protrusion					
		Nerve Compression		Motrin 800 Mg	PS		
OROPHARINGEAL							
		Neuropathy		Ultram	C		
OROPHARINGEAL							
		Pain Pain In Extremity Shoulder Pain		Anti-Inflammatory Medications	C		
OROPHARINGEAL				Neurontin	I		
OROPHARINGEAL				Neurontin	I		
OROPHARINGEAL				Neurontin	I		
OROPHARINGEAL				Hydrocodone Mobic	I I		

Date:07/12/05ISR Number: 4713849-2Report Type:Expedited (15-DaCompany Report #2005094952
Age:62 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged SEE IMAGE		Blood Glucose Decreased Blood Pressure Increased	Consumer	Accupril (Quinapril Hydrochloride)	PS		
Disability Other 600 MG, ORAL		Bone Infection Cataract		Neurontin (Gabapentin)	SS		ORAL
		Cerebrovascular Accident Diabetic Complication Diabetic Retinopathy Drug Effect Decreased Fall Limb Injury Localised Infection Memory Impairment Myopia Nail Dystrophy Post Procedural Complication Speech Disorder Toe Amputation		Insulin (Insulin) Furosemide (Furosemide) Daily Vitamins (Vitamins Nos)	SS C C		

Upper Limb Fracture
Vision Blurred
Wrist Fracture

Date:07/12/05ISR Number: 4713863-7Report Type:Expedited (15-DaCompany Report #2005094675
Age:49 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability		Feeling Abnormal Vomiting	Consumer	Neurontin (Gabapentin)	PS		ORAL

ORAL

Date:07/12/05ISR Number: 4714637-3Report Type:Expedited (15-DaCompany Report #2005074070
Age:66 YR Gender:Male I/FU:I

Outcome	PT
Hospitalization - Initial or Prolonged Disability	Abdominal Pain Upper Anorexia Asthenia Condition Aggravated Constipation Disease Recurrence Fatigue 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
900 MG (300 MG,3 IN 1 D),ORAL		Nausea Pain Skin Burning Sensation Weight Decreased	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
ORAL				Carbimazole (Carbimazole)	SS		ORAL
				Restex (Benserazide Hydrochloride, Levodopa)	C		
				Tramal (Tramadol Hydrochloride)	C		

Date:07/13/05ISR Number: 4714940-7Report Type:Expedited (15-DaCompany Report #2005091349
Age:75 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Alanine Aminotransferase Increased	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
Other		Antibody Test Positive Antimitochondrial	Professional	Nifedipine (Nifedipine)	SS		ORAL
ORAL		Antibody Positive Antinuclear Antibody Positive		Kardegic (Acetylsalicylate Lysine)	C		
		Arrhythmia Ascites Aspartate Aminotransferase Increased		Hyperium (Rilmenidine)	C		
		Atrial Fibrillation Biliary Cirrhosis Primary Blood Alkaline Phosphatase Increased Blood Bilirubin Increased Cardiac Failure					

Hepatic Enzyme Increased
Lung Disorder
Lymphadenopathy

Date:07/13/05ISR Number: 4715451-5Report Type:Expedited (15-DaCompany Report #2004106658
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Ineffective Gun Shot Wound Head Injury Intentional Self-Injury Panic Attack	Consumer	Neurontin (Gabapentin)	PS		

Date:07/14/05ISR Number: 4714595-1Report Type:Direct Company Report #CTU 253264
Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other 600 MG PO		Condition Aggravated Loss Of Consciousness		Neurontin 600 Mg -Generic	PS		ORAL

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/14/05ISR Number: 4714804-9Report Type:Direct
Age:50 YR Gender:Male I/FU:I

Company Report #CTU 253212

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 600 TID ORAL		Drug Ineffective Therapeutic Response Unexpected With Drug Substitution		Gabapentin 600 Greenstone	PS	Greenstone	ORAL

Date:07/14/05ISR Number: 4716488-2Report Type:Expedited (15-DaCompany Report #2005039647
Age:20 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Disability 1200 MG (600 MG, 2 IN 1 D), ORAL		Abasia Cardiovascular Disorder Condition Aggravated Pain In Extremity Peripheral Coldness	Foreign Consumer	Neurontin (Tablets) (Gabapentin) All Other Therapeutic Producuts (All Other Therapeutic Products) Cortisone (Cortisone)	PS C C		ORAL

Date:07/14/05ISR Number: 4716562-0Report Type:Expedited (15-DaCompany Report #2005094694
Age:64 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 900 MG (300 MG, 3 IN 1 D), ORAL		Anaesthetic Complication Pain Post Procedural Complication	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL

Post Procedural Pain
Renal Pain
Urethral Stricture
Urinary Incontinence
Urinary Retention

Ciprofloxacin
(Ciprofloxacin) C
Dexibuprofen
(Dexibuprofen) C

Date:07/14/05ISR Number: 4716648-0Report Type:Expedited (15-DaCompany Report #2004106748
Age:29 YR Gender:Male I/FU:F

Outcome PT
Hospitalization - Abnormal Behaviour
Initial or Prolonged Aggression
Other Agitation
Condition Aggravated
Delusion
Depression
Drug Ineffective
Hallucination, Auditory
Hallucination, Visual
Homicidal Ideation
Intentional Misuse
Lethargy
Multiple Drug Overdose
Psychotic Disorder
Refusal Of Treatment By
Patient

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Somnolence Stress Suicidal Ideation	Consumer	Neurontin (Tablets) (Gabapentin)	PS		
900 MG (300 MG, 3 IN 1 D)		Suicide Attempt Treatment Noncompliance		Neurontin (Gabapentin)	SS		
900 MG (300 MG, 3 IN 1 D)				Haldol (Haloperidol) Risperidone (Risperidone)	SS SS		
6 MG (3 MG, 2 IN 1 D)				Seroquel (Quetiapine Fumarate) Metformin (Metformin)	C C		

Date:07/14/05ISR Number: 4716696-0Report Type:Expedited (15-DaCompany Report #2005074293
Age:45 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Brain Damage Condition Aggravated	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
1800 MG (600 MG, 3 IN 1 D), ORAL		Drug Effect Decreased Drug Interaction		Motrin (Ibuprofen)	SS		ORAL
(800 MG, UNKNOWN), ORAL		Insomnia Intervertebral Disc Protrusion		Hydrocodone (Hydrocodone)	SS		
		Malaise Nerve Compression					

ORAL		Neuropathy		Mobic (Meloxicam)	SS		ORAL
		Pain		Ultram (Tramadol Hydrochloride)	C		
		Pharmaceutical Product Complaint					
Date:07/15/05		ISR Number: 4715428-X		Report Type:Direct		Company Report #CTU 253395	
Age:	Gender:Female	I/FU:I					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Pharmaceutical Product		Gabapentin 100 Mg	PS		
3X		Complaint					
		Tremor					
Date:07/15/05		ISR Number: 4715674-5		Report Type:Direct		Company Report #CTU 253398	
Age:67 YR	Gender:Female	I/FU:I					
Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Dyskinesia		Gabapentin 100 Mg/3			
		Muscle Spasms		Days. 200 Mg 3 Days	PS		ORAL
ONCE A DAY		Tremor					
ORAL				Gabapentin 300 Mg			
				/Day	SS		ORAL
ONCE A DAY							
ORAL				Losec	C		
				Premarin	C		
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Freedom Of Information (FOI) Report

Date:07/15/05ISR Number: 4717378-1Report Type:Expedited (15-DaCompany Report #2005003387
 Age:59 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - Initial or Prolonged		Cerebral Haematoma	Consumer	Neurontin (Gabapentin)	PS		
Other		Fracture					
		Conjunctivitis					
		Corneal Ulcer					
		Drug Ineffective					
		Dysphagia					
		Fracture					
		Gun Shot Wound					
		Mastoid Disorder					
		Pneumonia Haemophilus					
		Self Injurious Behaviour					
		Subdural Haematoma					
		Subdural Hygroma					
		Suicide Attempt					
		Tracheostomy					
		Unresponsive To Verbal Stimuli					

Date:07/15/05ISR Number: 4717449-XReport Type:Expedited (15-DaCompany Report #2005071563
 Age:6 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Abasia	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (300		Arthralgia	Health				
Other		Bedridden	Professional				
MG, 1 IN 1		Bipolar Disorder					
D), ORAL		Condition Aggravated		Lamictal (Lamotrigine)	SS		
		Convulsion		Zyprexa (Olanzapine)	SS		
		Dizziness		Seroquel (Quetiapine Fumarate)	SS		ORAL
ORAL		Drug Ineffective					
		Increased Appetite					
		Insomnia		Depakote (Valproate Semisodium)	SS		ORAL
ORAL		Mania					

Metabolic Disorder
 Migraine
 Speech Disorder
 Suicidal Ideation
 Weight Increased

Depakene (Valproate
 Sodium) C
 Claritin
 (Loratadine) C
 Valproic Acid
 (Valproic Acid) C

Date:07/18/05ISR Number: 4718156-XReport Type:Direct
 Age:57 YR Gender:Male I/FU:I

Company Report #CTU 253427

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - 2 MG PO QHS		Blood Creatine Increased	Terazosin	PS		ORAL
Initial or Prolonged 600 MG PO TID		Hyperkalaemia	Gabapentin	SS		ORAL
			Terazosin Hcl	C		
			Omeprazole	C		
			Albuterol Inh	C		
			Metoprolol Tartrate	C		
			Phenazopyridine Hcl	C		
			Clopidogrel			
			Bisulfate	C		
			Flunisolide Nasal Spray	C		

Freedom Of Information (FOI) Report

Guaifenesin	C
Aspirin	C
Tiotropium Bromide	
Inh	C
Gabapentin	C
Irbesartan	C
Isosorbide Dinitrate	C
Diphenhydramine Hcl	C
Metoclopramide Hcl	C
Simvastatin	C
Advair	C
Carbamazepine	C
Oyst Cal D	C
Simethicone	C
Nitroglycerin	C
Acetaminophen	C
Albuterol So4 Inh	C
Loperamide Hcl	C
Ciprofloxacin 0.3%	
Opth Drops	C

Date:07/19/05ISR Number: 4718691-4Report Type:Expedited (15-DaCompany Report #2004032207
 Age:49 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin			
Other		Drug Ineffective		(Gabapentin)	PS		ORAL
300 MG (300		Injury Asphyxiation					
MG, 1 IN 1		Intervertebral Disc					
D), ORAL		Disorder		Vicodin			
		Neuropathy Peripheral		(Hydrochloride			
		Polytraumatism		Bitartrate,			
		Self Injurious Behaviour		Paracetamol)	C		
		Somnolence		Cardura	C		
		Stress		Celexbra (Celecoxib)	C		
		Treatment Noncompliance		Remeron			
				(Mirtazapine)	C		
				Klonopin			
				(Clonazepam)	C		
				Voltaren (Diclofenac			
				Sodium)	C		

Outcome	PT
Hospitalization -	Blood Glucose Decreased
Initial or Prolonged	Blood Pressure Increased
Disability	Bone Infection
Other	Cataract
	Cerebrovascular Accident
	Diabetic Foot Infection
	Diabetic Retinopathy
	Drug Effect Decreased
	Fall
	Localised Infection
	Memory Impairment
	Nail Disorder
	Oedema Peripheral

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Peripheral Coldness Pharmaceutical Product Complaint	Report Source	Product	Role	Manufacturer	Route
10 MG ORAL		Speech Disorder Toe Amputation	Consumer	Accupril (Quinapril Hydrochloride)	PS		ORAL
600 MG ORAL		Upper Limb Fracture Wrist Fracture		Neurontin (Gabapentin)	SS		ORAL
				Insulin (Insulin) Furosemide Daily Vitamins (Vitamins Nos)	SS C C		

Date:07/19/05ISR Number: 4718807-XReport Type:Expedited (15-DaCompany Report #2005098075
Age:42 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL		Amnesia Drug Interaction	Health Professional	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Disability Other		Drug Screen False Positive Dyskinesia Dyspnoea Dysstasia Fatigue Influenza Loss Of Consciousness Motor Dysfunction Nasopharyngitis Sciatica Somnolence		Morphine (Morphine) Desyrel (Trazodone Hydrochloride) Zoloft (Sertraline)	SS C C		

Date:07/19/05ISR Number: 4719440-6Report Type:Expedited (15-DaCompany Report #2005098035
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Back Pain	Consumer	Zyrtec (Tablets)			

Initial or Prolonged	Diabetes Mellitus	(Cetirizine)	PS	ORAL
1 IN 1 D ORAL				
Other	Hallucination	Celebrex (Celecoxib)	SS	ORAL
200 MG (200				
MG, 1 IN 1 D)	Intervertebral Disc			
ORAL	Disorder			
	Nerve Injury	Neurontin		
3600 MG (3 IN	Rheumatoid Arthritis	(Gabapentin)	SS	ORAL
11D) ORAL	Spinal Fracture			
		Bextra (Valdecoxib)	SS	

Date:07/19/05ISR Number: 4719530-8Report Type:Expedited (15-DaCompany Report #2005098827
Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Accidental Exposure	Consumer	Lipitor			
80 MG (40 MG,		Asthenia		(Atorvastatin)	PS		ORAL
2 IN 1 D)		Dizziness					
ORAL		Gallbladder Operation					
		Hernia		Neurontin			
		Loss Of Consciousness		(Gabapentin)	SS		
		Malaise		Glucotrol			
		Nausea		(Glipizide)	C		
		Wrong Drug Administered		Glucophage Xr			

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Freedom Of Information (FOI) Report

(Metformin
Hydrochloride) C
Metoprolol C
Captopril C

Date:07/19/05ISR Number: 4720376-5Report Type:Expedited (15-DaCompany Report #2005079827
Age:65 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 400 MG (400 MG, 1 IN 1 D), ORAL		Bladder Pain	Consumer	Celebrex (Celecoxib)	PS		ORAL
		Bone Disorder					
		Drug Ineffective					
900 MG (300 MG, 3 IN 1 D)		Haemangioma Hypertonic Bladder		Gabapentin (Gabapentin)	SS		
		Intervertebral Disc Protrusion Neurological Symptom Neuropathy Peripheral Osteoarthritis Pain Scoliosis Somnolence Tension Toothache Treatment Noncompliance Urinary Incontinence		Elvail (Amitriptyline Hydrochloride) Arthrotec(Diclofenac Sodium, Misoprostol) All Other Therapeutic Products (All Other Therapeutic Products) Valsartan (Valsartan)	SS SS SS		

Date:07/19/05ISR Number: 4720742-8Report Type:Expedited (15-DaCompany Report #2005099146
Age:48 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Constipation Diarrhoea	Health Professional	Neurontin (Gabapentin)	PS		ORAL
		Drug Ineffective		Elavil (Amitriptyline			

10 MG (10

Hydrochloride)

SS

ORAL

MG,1 IN 1

D),ORAL

Advil

(Ibuprofen)

C

Dicyclomine

(Dicycloverine)

C

Aygestin

(Norethisterone

Acetate)

C

Date:07/19/05ISR Number: 4720744-1Report Type:Expedited (15-DaCompany Report #2004117310

Age:37 YR Gender:Female I/FU:F

Outcome	PT
Death	Accidental Overdose
Hospitalization -	Back Pain
Initial or Prolonged	Completed Suicide
Other	Consciousness Fluctuating
	Depressed Level Of
	Consciousness
	Depression
	Drug Dependence

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FDA - Adverse Event Reporting System (AERS)

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Dose	Duration	Drug Ineffective Homicidal Ideation Intentional Misuse	Report Source	Product	Role	Manufacturer	Route
300 MG (100 MG,3 IN 1 D); 1800 MG (600 MG,3 IN 1D)		Loss Of Consciousness Multiple Drug Overdose Accidental Neck Pain	Consumer	Neurontin (Gabapentin)	PS		
				Ibuprofen (Ibuprofen)	SS		
				Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		
				Nortriptyline Hydrochloride (Nortriptyline Hydrochloride)	SS		
				Venlafaxine (Venlafaxine)	SS		
				Oxazepam (Oxazepam)	C		
				Benzodiazepine Derivatives (Benzodiazepine Derivatives)	C		
				Opioids (Opioids)	C		

Date:07/20/05ISR Number: 4718726-9Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20050507147
Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death 800 mg, twice to three times, as needed		Completed Suicide Loss Of Consciousness Multiple Drug Overdose Accidental		Motrin 800 Mg	PS		

Nortriptyline	
Hydrochloride	SS
Amitriptyline	
Hydrochloride	SS
Venlafaxine	SS
Neurontin	SS
Neurontin	SS
Neurontin	SS
Neurontin	SS
Oxazepam	C
Benzodiazepine	
Derivatives	C
Opioids	C

Date:07/20/05ISR Number: 4719088-3Report Type:Expedited (15-DaCompany Report #2005099151
Age:21 YR Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 1800 MG (600 MG, 3 IN 1 D),ORAL	Respiratory Distress	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
			Valproate Sodium (Valproate Sodium)	C		

Date:07/20/05ISR Number: 4720982-8Report Type:Expedited (15-DaCompany Report #2005099543

Age: Gender: I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Infection Neutropenia	Foreign Health Professional	Gabapentin (Gabapentin)	PS		

Date:07/20/05ISR Number: 4722349-5Report Type:Expedited (15-DaCompany Report #2005098035

Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Diabetes Mellitus
Initial or Prolonged	Hallucination
Other	Intervertebral Disc Injury

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Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
(1 IN 1 D), ORAL		Nerve Injury Rheumatoid Arthritis Spinal Fracture	Consumer	Zyrtec (Tablets) (Cetirizine)	PS		ORAL
3600 MG (3 IN 1 D), ORAL				Neurontin (Gabapentin)	SS		ORAL
200 MG (200 MG, 1 IN 1 D), ORAL				Bextra (Valdecoxib) Celebrex (celecoxib)	SS C		ORAL

Date:07/20/05ISR Number: 4722441-5Report Type:Expedited (15-DaCompany Report #2005099177

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other 1250 MG (1 D) ORAL		Drug Abuser Drug Withdrawal Syndrome Lethargy Somnolence Tooth Disorder Tooth Loss	Consumer	Neurontin (Gabapentin)	PS		ORAL
				Ditropan (Oxybutynin)	C		
				Senokot (Senna Fruit)	C		
				Zocor (Simvastatin)	C		
				Benadryl	C		
				Acetylsalicylic Acid	C		
				Multivitamins With Minerals	C		
				Vitamin C	C		

Age: Gender:Female I/FU:F

Outcome	PT
Hospitalization -	Abdominal Pain
Initial or Prolonged	Abnormal Behaviour
Other	Agitation
	Anorexia
	Anxiety
	Arthritis
	Balance Disorder
	Condition Aggravated
	Dehydration
	Depression
	Diabetes Mellitus
	Diarrhoea
	Hostility
	Impulsive Behaviour
	Insomnia
	Mania
	Mood Swings
	Muscle Spasms
	Obesity
	Paralysis
	Pneumonia
	Suicidal Ideation
	Suicide Attempt

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Tremor Visual Disturbance	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Neurontin (Gabapentin)	PS		

Date:07/20/05ISR Number: 4722530-5Report Type:Expedited (15-DaCompany Report #2005099567
Age:37 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose			Consumer	Neurontin (Gabapentin)	PS		ORAL
Other		Cerebrovascular Accident Factor V Deficiency					

600 MG (300
MG, 2 IN 1 D)

ORAL

Ceclor (Cefaclor)	C
Theophylline	C
Tambocor (Flecainide Acetate)	C
Lanoxin (Digoxin)	C
Clinidine	C
Lotensin (Benazepril Hydrochloride)	C
Glucophage (Metformin Hydrochloride)	C
Bumex (Bumetanide)	C
Micro-K (Potassium Chloride)	C
Coumadine (Warfarin Sodium)	C
Methotrexate	C
Prilosec	C
Nexium (Esomeprazole)	C
Verapamil	C
Diltiazem	C
Serevent (Salmeterol Xinafoate)	C
Aerobid (Flunisolide)	C

Date:07/20/05ISR Number: 4722612-8Report Type:Expedited (15-DaCompany Report #2005075005
Age: Gender:Female I/FU:F

Outcome	PT
Disability	Amnesia
Other	Anxiety
	Back Pain
	Blood Alkaline
	Phosphatase Increased
	Blood Cholesterol
	Increased
	Confusional State
	Difficulty In Walking
	Drug Dependence
	Eye Disorder
	Fluid Retention

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	High Density Lipoprotein Decreased Intervertebral Disc Disorder Migraine	Report Source	Product	Role	Manufacturer	Route
1.5 MG (0.5 MG, 3 IN 1 D), ORAL			Consumer Health	Xanax Tablet (Alprazolam)	PS		ORAL
		Oedema Peripheral	Professional				
		Retinal Detachment					
500 MG (100 MG, 3 IN 1 D), ORAL		Unevaluable Event Weight Increased		Neurontin (Gabapentin)	SS		ORAL
		Weight Loss Poor					
				Fioricet (Butalbital, Caffeine, Paracetamol)	SS		
				Avapro (Irbesartan)	C		
				Prevacid (Lansoprazole)	C		

Date:07/20/05ISR Number: 4723724-5Report Type:Direct
Age:53 YR Gender:Male I/FU:I

Company Report #CTU 253792

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Confusional State Lethargy		Gabapentin 300 Mg Daily X 1 Day Bid X Daythen Then Ii	PS		

Date:07/20/05ISR Number: 4724703-4Report Type:Direct
Age:32 YR Gender:Female I/FU:I

Company Report #CTU 253740

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Agitation Condition Aggravated Family Stress		Zoloft 150 Pfizer U.S/ Pfizer Us Pharmaceutic	PS	Pfizer U.S/ Pfizer	

150	DAILY	Fatigue			Us Pharmaceutical	ORAL
		Feeling Abnormal				
	ORAL	Headache	Neurontin 300			
		Hyperacusis	Teva Pharm	SS	Teva Pharm	ORAL
300	TWICE	Impaired Work Ability				
	DAILY	Mood Swings				
	ORAL	Somnolence				

Date:07/21/05ISR Number: 4723966-9Report Type:Expedited (15-DaCompany Report #2005042730
Age:42 YR Gender:Male I/FU:F

Outcome	PT
Life-Threatening	Abnormal Behaviour
Hospitalization -	Agitation
Initial or Prolonged	Anorexia
Disability	Atelectasis
Other	Blindness
	Brain Contusion
	Catheter Related
	Complication
	Cerebral Haemorrhage
	Cognitive Disorder
	Delirium
	Diabetes Insipidus

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Disorder Of Globe Drug Ineffective Dyspepsia	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1 D)		Dysphagia Eye Excision Facial Bones Fracture Gun Shot Wound Haemorrhage Hypocalcaemia Hypomagnesaemia Loss Of Consciousness Malnutrition Pleural Effusion Pneumocephalus Pneumonia Bacterial Pseudomonas Infection Pyrexia Respiratory Failure Soft Tissue Injury Suicide Attempt	Consumer	Neurontin (Gabapentin) Paxil (Paroxetine Hydrochloride) Thorazine (Chlorpromazine Hydrochloride) Ambien (Zolpidem Tartrate)	PS C C C		

Date:07/21/05ISR Number: 4723967-0Report Type:Expedited (15-DaCompany Report #2005099291

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other ORAL		Blood Pressure Decreased Body Height Decreased Cystitis Hypoesthesia Kidney Infection Multiple-Drug Resistance Neuralgia Pain Post Procedural Pain Postoperative Adhesion Urinary Tract Infection Uterine Operation	Consumer	Neurontin (Gabapentin) Ciprofloxacin (Ciprofloxacin) Macrobid (Nitrofurantoin) Antihypertensives (Antihypertensives) Oxycontin (Oxycodone Hydrochloride) Percocet (Oxycodone Hydrochloride, Paracetamol) Ativan (Lorazepam) Prevacid	PS SS SS SS C C C		ORAL

(Lansoprazole) C
Voltaren (Diclofenac
Sodium) C
Acetylsalicylic Acid
(Acetylsalicylic
Acid) C
Calcium (Calcium) C
Evista (Raloxifene
Hydrochloride) C
Zetia (Ezetimibe) C

Date:07/21/05ISR Number: 4723969-4Report Type:Expedited (15-DaCompany Report #2005099295
Age:64 YR Gender:Male I/FU:I

Outcome PT
Other Blood Pressure Systolic
Decreased
Crying
Depression

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1 D), ORAL		Dizziness Feeling Abnormal Haemorrhage	Consumer	Neurontin (Gabapentin)	PS		ORAL
75 MG		Impaired Healing Neuralgia Pain Skin Exfoliation Sleep Apnoea Syndrome Somnolence Wound Secretion		Acetylsalicylic Acid (Acetylsalicylic Acid) Plavix (Clopidogrel Sulfate) Warfarin Sodium (Warfarin Sodium) Insulin (Insulin) Coreg (Carvedilol) Klor-Con (Potassium Chloride) Isosorbide Dinitrate (Isosorbide Dinitrate) Zetia (Ezetimibe) Metolazone (Metolazone) Digitex (Digoxin) Zocor (Simvastatin) Zestril (Lisinopril) Lasix (Furosemide)	SS SS SS C C C C C C C C C C C C C		

Date:07/21/05ISR Number: 4724475-3Report Type:Expedited (15-DaCompany Report #2005073479
Age:45 YR Gender:Female I/FU:F

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
900 MG (300 MG, 3 IN 1		Arrhythmia Blood Pressure Fluctuation	Foreign Consumer	Neurontin (Gabapentin)	PS		ORAL

D), ORAL

Blood Pressure Systolic

Increased
Cardiac Failure
Gastric Perforation

Ixel (Milnacipran) C
Ludiomil Tablet
(Maprotiline
Hydrochloride) C
Neozine
(Levomepromazine) C

Date:07/21/05ISR Number: 4724505-9Report Type:Expedited (15-DaCompany Report #S05-CAN-02456-01

Age:47 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coma Suicide Attempt	Foreign Health Professional Other	Citalopram Hydrobromide (Citalopram) Clozapine Gabapentin Seroquel (Quetiapine Fumarate)			PS SS SS SS

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/21/05ISR Number: 4725062-3Report Type:Expedited (15-DaCompany Report #2005081014
 Age:60 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abdominal Discomfort	Consumer	Neurontin			
		Drug Interaction		(Gabapentin)	PS		
		Spinal Operation		Zantac 75			
				(Ranitidine			
				Hydrochloride)	SS		
				Tylenol			
				(Paracetamol)	C		
				Colace (Docusate			
				Sodium)	C		

Date:07/22/05ISR Number: 4722885-1Report Type:Expedited (15-DaCompany Report #PHFR2005GB02481
 Age:65 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Life-Threatening		Haematuria		Methylphenidate	PS	Novartis Sector: Pharma	ORAL
				Gabapentin	SS		ORAL

Date:07/22/05ISR Number: 4723814-7Report Type:Expedited (15-DaCompany Report #2005100287
 Age:29 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin			
		Intentional Misuse		(Gabapentin)	PS		
		Intentional Self-Injury					
		Overdose					
		Suicide Attempt					

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Breast Cancer Female	Consumer	Celebrex (Celecoxib)	PS		ORAL
ORAL		Drug Ineffective		Neurontin			
		Oedema Peripheral		(Gabapentin)	SS		ORAL
900 MG (300		Somnolence					
MG, 3 IN 1							
D), ORAL							
				Glucophage			
				(Metformin			
				Hydrochloride)	C		
				Glipizide			
				(Glipizide)	C		
				Plavix (Clopidogrel			
				Sulfate)	C		
				Monopril (Fosinopril			
				Sodium)	C		
				Clonidine			
				(Clonidine)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/22/05ISR Number: 4725033-7Report Type:Expedited (15-DaCompany Report #2005103174

Age:31 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Ineffective Gun Shot Wound Inadequate Analgesia Intentional Self-Injury	Consumer	Neurontin (Gabapentin)	PS		

Date:07/22/05ISR Number: 4725785-6Report Type:Direct Company Report #CTU 254107

Age:84 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - 800 MG ORALLY Initial or Prolonged TID		Mental Status Changes		Neurontin	PS		ORAL
150 MG, ORALLY,QD				Effexor Xr	SS		ORAL
0.5 MG, ORALLY, QD				Atavan	SS		ORAL
600 MG, ORALLY, QD				Oscal + D	SS		ORAL
10 MG, ORALLY, QD				Ditropan	SS		ORAL
5 MG, ORALLY, QD				Coumadin	SS		ORAL
300 MG, ORALLY, QD				Dilantin	SS		ORAL

10 MG,	Zolpidem - Ambien	SS	ORAL
ORALLY, QHS			
20 MG,	Kadian	SS	ORAL
ORALLY, QD			
15 ML ,	Enulose	SS	ORAL
ORALLY, QD			
5 MG, ORALLY,	Roxicodone	SS	ORAL
QD			
10 MG ,	Zyrtec	SS	ORAL
ORALLY, QD			
	Augmentin	SS	
	Lorazepam	C	
	Zolpidem	C	
	Morphine Sulfate	C	
	Oxycodone	C	
	Cetirizine	C	

Date:07/25/05ISR Number: 4725159-8Report Type:Expedited (15-DaCompany Report #2005071187
Age:84 YR Gender:Male I/FU:F

Outcome PT
Hospitalization - Anticonvulsant Drug Level
Initial or Prolonged Decreased
Other Blood Potassium Decreased
Clostridium Difficile
Toxin Test Positive
Diarrhoea
Drug Ineffective
Grand Mal Convulsion
Incorrect Dose
Administered
Medication Error

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Pharmaceutical Product Complaint Status	Report Source	Product	Role	Manufacturer	Route
		Epilepticus					
		Tremor	Consumer	Neurontin (Gabapentin)	PS		ORAL
400 MG (200 MG, 2 IN 1 D), ORAL				Dilantin (Phenytoin Sodium)	SS		ORAL
360 MG (230MG AM, 130MG PM), ORAL				Zocor (Simvastatin)	C		
				Acetylsalicylic Acid (Acetylsalicylic Acid)	C		
				Potassium Chloride (Potassium Chloride)	C		
				Multivitamins (Multivitamins)	C		
				Aldactone (Spironolactone)	C		
				Plavix (Clopidogrel Sulfate)	C		
				Lasix (Furosemide)	C		
				Prilosec (Omeprazole)	C		
				Tamsulosin (Tamsulosin)	C		
				Toprol (Metoprolol)	C		
				Metronidazole (Metronidazole)	C		
				All Other Therapeutic Products(All Other Therapeutic Products)	C		

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Difficulty In Walking Drug Abuser Fatigue	Consumer Health Professional	Desoxyn (Methamphetamine Hcl)	PS		ORAL
30 MG; Q4H; PO	18	MON					
2MG		Impaired Driving Ability Imprisonment		Xanax (Alprazolam)	SS		
80 MG; Q4H		Intentional Misuse Loss Of Consciousness		Oxycontin (Oxycodone Hydrochloride)	SS		
100 MCG		Murder		Duragesic Patches	SS		
40 MG, QD		Pain Road Traffic Accident		Prilosec (Omeprazole)	SS		
300 MG; Q4H		Speech Disorder		Neurontin (Gabapentin)	SS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/25/05ISR Number: 4725218-XReport Type:Expedited (15-DaCompany Report #2005099668

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged ORAL Other		Back Pain Blood Disorder	Consumer	Neurontin (Gabapentin)	PS		ORAL
1600 MG (400 MG, 4 IN 1 D)		Blood Glucose Increased Carbohydrate Metabolism Disorder		Vicodin (Hydrocodone Bitartrate, Paracetamol)	SS		
		Cardiac Disorder					
		Lung Disorder		Amitriptyline (Amitriptyline0	C		

Date:07/25/05ISR Number: 4725246-4Report Type:Expedited (15-DaCompany Report #2005103178

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		

Date:07/25/05ISR Number: 4725332-9Report Type:Expedited (15-DaCompany Report #2005103308

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

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Depression Drug Ineffective Homicidal Ideation Suicidal Ideation	Consumer	Neurontin (Gabapentin)	PS		

Date:07/25/05ISR Number: 4726177-6Report Type:Direct Company Report #CTU 254244

Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Other Drug Effect Decreased Gabapentin 100 Mg PS ORAL

100 MG TID PO

Pharmaceutical Product
Complaint
Somnolence

Date:07/25/05ISR Number: 4726330-1Report Type:Expedited (15-DaCompany Report #2005101464

Age:27 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Abortion Spontaneous Drug Exposure During Pregnancy	Consumer	Neurontin (Gabapentin) Keppra (Levetiracetam) Topamax (Topiramate) Tylenol (Paracetamol)	PS SS C C		

Date:07/26/05ISR Number: 4727282-0Report Type:Expedited (15-DaCompany Report #2005016042

Age: Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Aggression
Initial or Prolonged	Agitation
Other	Amnesia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Bacteraemia Bipolar Disorder Brain Oedema	Consumer	Neurontin (Gabapentin)	PS		
		Cerebrovascular Accident Coma Computerised Tomogram Abnormal Drug Dependence Drug Ineffective Encephalomalacia Faecal Incontinence General Physical Health Deterioration Gun Shot Wound Head Injury Major Depression Paraparesis Pneumonia Pulmonary Congestion Respiratory Failure Simple Partial Seizures Sinus Disorder Skin Ulcer Subdural Haematoma Suicide Attempt Thrombosis Traumatic Brain Injury Urinary Incontinence Wound Dehiscence Wound Infection					

Date:07/27/05ISR Number: 4727757-4Report Type:Expedited (15-DaCompany Report #2005EU001591
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 1.00 MG, UID/QD, ORAL		Blood Triglycerides Increased	Foreign Health	Prograf (Tacrolimus) Capsule, 0.5mg	PS		ORAL
		Deep Vein Thrombosis	Professional				
2.00 MG,		Epistaxis	Other	Rapamune (Sirolimus)	SS		ORAL

UID/QD, ORAL	Hepatic Steatosis			
1500.00 MG,	Hypoalbuminaemia	Cellcept		
UID/QD, ORAL	Neuropathy Peripheral	(Mycophenolate		
	Pancreatic Disorder	Mofetil) Tablet,		
	Pericardial Effusion	500mg	SS	ORAL
	Pulmonary Embolism			
300.00 MG,		Zyloric "Glaxo		
UID/QD, ORAL		Wellcome"		
		(Allopurinol)		
		Tablet, 300mg	SS	ORAL
2400.00 MG,		Neurontin		
UID/QD, ORAL		(Gabapentin)	SS	ORAL
25.00,		Laroxyl		
UID/QD, ORAL		(Amitriptyline		
		Hydrochloride) 25mg	SS	ORAL
		Rivotril		
		(Clonazepam)	C	
		Orofcac (Calcium		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Carbonate) C
 Lasilix (Furosemide) C
 Levothyrox
 (Levothyroxine
 Sodium) C

Date:07/27/05ISR Number: 4728259-1Report Type:Expedited (15-DaCompany Report #2005106436
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Coma	Consumer	Neurontin (Gabapentin)	PS		

Date:07/27/05ISR Number: 4728265-7Report Type:Expedited (15-DaCompany Report #2005103498
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Injury Multiple Drug Overdose Suicide Attempt	Consumer	Neurontin (Gabapentin) All Other Therapeutics Products (All Other Therapeutics Products)	PS SS		

Date:07/27/05ISR Number: 4728267-0Report Type:Expedited (15-DaCompany Report #2005104601
 Age:34 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Intentional Self-Injury Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:07/27/05ISR Number: 4728324-9Report Type:Expedited (15-DaCompany Report #2004063520
 Age:33 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose Other		Alopecia Anxiety	Consumer	Neurontin (Gabapentin)	PS		ORAL
300 MG (300 MG, 1 IN 1 D), ORAL (1 IN 1 D), ORAL		Body Height Decreased Chest Discomfort Depression Disturbance In Attention		Zoloft (Sertraline)	SS		ORAL
		Divorced Drug Ineffective Drug Interaction Dyspnoea Feeling Abnormal Headache		Xanax Tablet (Alprazolam) Lexapro (Escitalopram)Q Gabapentin (Gabapentin)	SS SS SS		ORAL
ORAL		Heart Rate Increased Mood Altered		Claritin (Loratadine)	SS		ORAL
ORAL		Myocardial Infarction Neck Pain Panic Attack Psychomotor Hyperactivity Road Traffic Accident Stress Suicidal Ideation		Buspar (Buspirone Hydrochloride) Prozac (Fluoxetine Hydrochloride) Zypreka Zyrtec	C C C C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/27/05ISR Number: 4728334-1Report Type:Expedited (15-DaCompany Report #2004083654

Age:61 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Anxiety	Consumer	Neurontin (Tablets)			
Other		Completed Suicide		(Gabapentin)	PS		ORAL
(600 MG),		Coronary Artery					
ORAL		Atherosclerosis		Alcohol (Ethanol)	SS		
		Depression		Diazepam (Diazepam)	SS		
15 MG (5 MG,		Drug Toxicity					
3 IN 1 D)		Emotional Distress		Oxycodone			
		Gait Disturbance		(Oxycodone)	SS		
7.5 MG		Injury		Prandin			
		Intentional Misuse		(Repaglinide)	C		
		Multiple Drug Overdose		Lidoderm (Lidocaine)	C		
		Nervous System Disorder		Verapamil			
		Pain		(Verapamil)	C		
		Suicidal Ideation		Hydrochlorothiazide/			
		Vascular Calcification		Triamtere			
				(Hydrochlorothiazide			
				Triamterene)	C		
				Clonidine			
				(Clonidine)	C		
				Ambien (Zolpidem			
				Tartrate)	C		
				Nexium			
				(Esomeprazole)	C		
				Oxycontin (Oxycodone			
				Hydrochloride)	C		

Date:07/27/05ISR Number: 4728347-XReport Type:Expedited (15-DaCompany Report #2005104607

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide	Consumer	Neurontin			
Other		Drug Ineffective		(Gabapentin)	PS		
		Suicidal Ideation					

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose	Duration					
Hospitalization -	Feeling Guilty	Consumer	Zoloft (Sertraline)	PS		ORAL
ORAL						
Initial or Prolonged	Hunger		Neurontin			
	Increased Appetite		(Gabapentin)	SS		ORAL
ORAL						
40 MG	Nausea		Geodon (Ziprasidone)	SS		
	Neurosis		Lamictal			
	Road Traffic Accident		(Lamotrigine0	C		
	Stress		All Other			
	Suicide Attempt		Therapeutic Products			
	Ulcer		(All Other			
	Victim Of Crime		Therapeutic			
			Products)	C		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/27/05ISR Number: 4728354-7Report Type:Expedited (15-DaCompany Report #2005003617
 Age:28 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 3200 MG (800 Other MG, 4 IN 1 D), ORAL	Affective Disorder Antisocial Personality Disorder Anxiety Disorder Attention Deficit/Hyperactivity Disorder Bipolar Ii Disorder Borderline Personality Disorder Drug Effect Decreased Gastrooesophageal Reflux Disease Gun Shot Wound Haemothorax Homicidal Ideation Intermittent Explosive Disorder Lung Injury Major Depression Personality Disorder Polysubstance Dependence Psychotic Disorder Suicidal Ideation Suicide Attempt Treatment Noncompliance	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Ativan (Lorazepam)	C		
			Celexa (Citalopram Hydrobromide)	C		
			Ambien (Zolpidem Tartrate)	C		

Date:07/27/05ISR Number: 4728464-4Report Type:Expedited (15-DaCompany Report #2005077695
 Age:88 YR Gender:Female I/FU:F

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Hospitalization - Initial or Prolonged 100 MG (100 Other MG 1 IN 1 D)	Back Pain Body Height Decreased Constipation	Consumer	Neurontin (Gabapentin)	PS		

		Crying	Bextra (Valdecoxib)	SS
		Delayed Recovery From	Mortin (Ibuprofen)	SS
		Anaesthesia	Fentanyl (Fentanyl)	SS
TRANSDERMAL	25 MCG,			
		Dizziness		
TRANSDERMAL				
		Drug Ineffective	Vioxx (Rofecoxib)	SS
		Feeling Abnormal	Advil (Ibuprofen)	SS
		Gastritis	Aleve (Naproxen	
		Impaired Driving Ability	Sodium)	C
		Insomnia	Levoxyl	
		Muscle Spasms	(Levothyroxine	
		Nausea	Sodium)	C
		Paralysis	All Other	
		Pruritus	Therapeutic Products	
		Psychomotor Hyperactivity	(All Other	
		Somnolence	Therapeutic	
		Stomach Discomfort	Products)	C
		Unevaluable Event		

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:07/27/05ISR Number: 4728533-9Report Type:Expedited (15-DaCompany Report #2005103484
Age:76 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Completed Suicide Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		

Date:07/27/05ISR Number: 4728553-4Report Type:Expedited (15-DaCompany Report #2005028641
Age:47 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Life-Threatening Hospitalization - 300 MG ORAL		Coma Intentional Misuse	Consumer	Neurontin (Gabapentin)	PS		ORAL
Initial or Prolonged Other ORAL		Multiple Drug Overdose Suicide Attempt		Hydrocodone (Hydrocodone)	SS		ORAL
ORAL				Soma (Carisoprodol)	SS		ORAL

Date:07/27/05ISR Number: 4728574-1Report Type:Expedited (15-DaCompany Report #2005104597
Age:57 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other		Completed Suicide Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		

Date:07/27/05ISR Number: 4728584-4Report Type:Expedited (15-DaCompany Report #2004071970
Age:39 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death Other 1800 MG (600 MG, 3 IN 1 D)		Anxiety Brain Death Brain Herniation	Consumer	Neurontin (Gabapentin)	PS		ORAL

ORAL	Brain Oedema				
	Cerebral Atherosclerosis	Nortriptyline	C		
	Completed Suicide	Nexium			
	Drug Ineffective	(Esomeprazole)	C		
	Gun Shot Wound	Allegra			
	Injury	(Fexofenadine			
	Pain	Hydrochloride)	C		
	Skull Fracture	Amitriptyline	C		
	Subarachnoid Haemorrhage	Multivitamins			
	Subdural Haemorrhage	W/Minerals (Minerals			
		Nos, Vitamins Nos)	C		
		Vitamin C	C		

Date:07/27/05ISR Number: 4728607-2Report Type:Expedited (15-DaCompany Report #2005103258
Age:28 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective Suicide Attempt	Consumer	Neurontin (Gabapentin)	PS		

Date:07/28/05ISR Number: 4729425-1Report Type:Expedited (15-DaCompany Report #2005103489
Age: Gender:Male I/FU:I

Outcome	PT
Other	Depression Drug Withdrawal Syndrome

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Restlessness

Dose	Duration	Report Source	Product	Role	Manufacturer	Route
6400 MG (4 IN 1 D), ORAL		Consumer	Neurontin (Gabapentin)	PS		ORAL

Date:07/28/05ISR Number: 4729698-5Report Type:Expedited (15-DaCompany Report #2004106658
Age:51 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Agoraphobia	Consumer	Neurontin (Gabapentin)	PS		
Other		Anxiety		Vistaril (Hydroxyzine Embonate)	C		
		Completed Suicide		Xanax (Alprazolam)	C		
		Depression		Buspar (Buspirone Hydrochloride)	C		
		Drug Ineffective		Risperdal (Risperidone)	C		
		Gun Shot Wound		Celexa (Citalopram Hydrobromide)	C		
		Head Injury		Prozac (Fluoxetine Hydrochloride)	C		
		Panic Attack		Ativan(Lorazepam)	C		
				Valium (Diazepam)	C		
				Seroquel (Quetiapine Fumarate)	C		
				Trazodone	C		
				Lexapro (Escitalopram)	C		

Date:07/28/05ISR Number: 4729724-3Report Type:Expedited (15-DaCompany Report #2004077855
Age:14 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other		Abnormal Behaviour	Consumer	Neurontin			

Aggression

(Gabapentin)

PS

ORAL

Coma

Levothyroid

Concussion

(Levothyroxine

Drug Abuser

Sodium)

C

Drug Ineffective

Clindamycin

C

Dysarthria

Seroquel (Quetiapine

Feeling Drunk

Fumarate)

C

Gait Disturbance

Lithium

C

Loss Of Consciousness

Multivitamins

C

Pain

Paxil (Paroxetine

Personality Change

Hydrochloride)

C

Polytraumatism

Clozaril (Clozapine)

C

Suicide Attempt

Ativan (Lorazepam)

C

Colace (Docusate

Sodium)

C

Provigil (Modafinil)

C

Topamax (Topiramate)

C

Naltrexone

C

Depo-Provera 150

(Medroxyprogesterone

Acetate)

C

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Albuterol
 (Salbutamol) C
 All Other
 Therapeutic Products C

Date:07/28/05ISR Number: 4729887-XReport Type:Expedited (15-DaCompany Report #2004106702
 Age:35 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Death		Activities Of Daily	Consumer	Neurontin			
Hospitalization -		Living Impaired		(Gabapentin)	PS		
Initial or Prolonged		Central Nervous System		Tramadol	SS		
Other		Mass		Ethanol (Ethanol)	SS		
		Completed Suicide		Acetaminophen			
		Depression		(Paracetamol)	SS		
		Drug Ineffective		Carisoprodol			
		Drug Toxicity		(Carisoprodol)	SS		
		Intentional Misuse		Meprobamate			
		Intervertebral Disc		(Meprobamate)	SS		
		Protrusion		Hydrocodone			
		Lumbar Spinal Stenosis		(Hydrocodone)	SS		
		Pulmonary Congestion		Oxycodone			
		Pulmonary Oedema		(Oxycodone)	SS		
		Spinal Cord Injury		Promethazine			
		Cervical		(Promethazine)	SS		
		Spinal Osteoarthritis		Skelaxin			
				(Metaxalone)	C		
				Cataflam (Diclofenac			
				Sodium)	C		
				Darvocet			
				(Dextropropoxyphene			
				Napsilate,			
				Paracetamol)	C		
				Vioxx (Rofecoxib)	C		

Date:07/28/05ISR Number: 4729916-3Report Type:Expedited (15-DaCompany Report #CA-2005-007813
 Age:54 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization -		Blood Pressure Decreased	Foreign	Betaseron			
Initial or Prolonged		Blood Pressure Increased	Consumer	(Interferon Beta			

Other	Cardiac Arrest	Health	-1b) Injection,	
Required	Catheter Related	Professional	250ug	PS
SUBCUTANEOUS	8 MIU, EVERY			
Intervention to	Complication	Other		
2D,				
Prevent Permanent	Condition Aggravated			
SUBCUTANEOUS				
Impairment/Damage	Cystitis		Gabapentin	
	Dyspnoea		(Gabapentin)	SS
	Hypokinesia		Ciprofloxacin	
	Multiple Sclerosis		(Ciprofloxacin)	C
	Muscle Disorder		Baclofen	C
	Nausea		Zanaflex (Tizanidine	
	Oedema Peripheral		Hydrochloride)	C
	Pyrexia		Amitriptyline	
	Sepsis		(Amitriptyline)	C
	Urinary Tract Infection		Diovan "Novartis"	
			(Valsartan)	C
			Lasix	C
			Vitamin B12	C
			Simvastatin	

D), ORAL

Hypoaesthesia

Pain In Extremity
Peripheral Coldness
Post Procedural
Complication

All Other
Therapeutic Products
(All Other
Therapeutic
Products) C
Cortisone C
(Cortisone)
Prednisone C
(Prednisone)
Cilostazol
(Cilostazol) C

Date:07/29/05ISR Number: 4734152-0Report Type:Expedited (15-DaCompany Report #2005104854
Age:40 YR Gender:Female I/FU:I

Outcome PT
Other Abasia
Adverse Drug Reaction
Alcoholism
Anxiety
Arthritis

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Attention Deficit/Hyperactivity Disorder	Report Source	Product	Role	Manufacturer	Route
120 MG (60 MG, 2 IN 1 D), ORAL		Bipolar Disorder	Consumer	Geodon (Ziprasidone)	PS		ORAL
		Cartilage Injury					
		Condition Aggravated					
ORAL		Disturbance In Attention Drug Dose Omission		Neurontin (Gabapentin)	SS		ORAL
		Gait Disturbance		Risperdal (Risperidone)	SS		
		Heart Rate Increased		Klonopin (Clonazepam)	C		
		Psychomotor Agitation		Flexeril (Cyclobenzaprine Hydrochloride)	C		
		Refusal Of Treatment By Patient		Metoprolol (Metoprolol)	C		
		Tendonitis		Hyoscyamine (Hyoscyamine)	C		
		Thinking Abnormal					
		Weight Increased					

Date:08/01/05ISR Number: 4730634-6Report Type:Expedited (15-DaCompany Report #2005UW10991
 Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged		Overdose		Seroquel	PS	Zeneca Pharmaceutical	ORAL
				Neurontin	SS		

Date:08/01/05ISR Number: 4731590-7Report Type:Direct Company Report #CTU 254829
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Required 900 MG PO TID		Coordination Abnormal		Gabapentin	PS		ORAL
Intervention to Prevent Permanent		Disorientation		Felodipine	C		
		Dysarthria		Nph	C		

Impairment/Damage	Neuropathy	Funasteride	C
	Phantom Pain	Terazosin	C
	Pleural Effusion	Niacin	C
	Tremor	Warfarin	C

Date:08/01/05ISR Number: 4732462-4Report Type:Expedited (15-DaCompany Report #2005105240

Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose Duration Hospitalization - Initial or Prolonged 1200 MG (300 Other MG 2 IN 1 D) ORAL	Blood Cholesterol Increased Blood Glucose Decreased Burning Sensation Chest Pain Fatigue Neuropathy Peripheral Pruritus Rash Macular Thrombosis Tremor	Consumer	Neurontin (Gabapentin)	PS		ORAL
			Lipitor (Atorvastatin)	SS		
			Benadryl (Diphenhydramine)	SS		
			Plavix (Clopidogrel Sulfate)	SS		
			Humulin (Insulin Human, Insulin Isophane, Human Biosynthetic)	SS		

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Freedom Of Information (FOI) Report

Humalog (Insulin Lispro) SS

Date:08/01/05ISR Number: 4732621-0Report Type:Expedited (15-DaCompany Report #2005104514
Age:83 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	2400 MG (800	Aortic Calcification Aortic Valve Stenosis	Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Other	MG, 3 IN 1	Blood Cholesterol					
		Increased					
		Cardiac Disorder Carotid Artery Occlusion Discomfort Drug Effect Decreased Eye Disorder High Density Lipoprotein Increased Knee Arthroplasty Low Density Lipoprotein Abnormal Mitral Valve Calcification Transient Ischaemic Attack		Vitamins (Vitamins)	C		

Date:08/01/05ISR Number: 4732625-8Report Type:Expedited (15-DaCompany Report #2002057211
Age:60 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Other	2400 MG	Back Pain Balance Disorder	Consumer Health	Neurontin (Gabapentin)	PS		ORAL
		Condition Aggravated	Professional				
		Drug Dependence Drug Ineffective Drug Withdrawal Syndrome		Darvocet (Dextropropoxyphene Napsilate,			

(DAILY), ORAL

ORAL	Fluid Retention	Paracetamol)	SS	ORAL
	Hangover	Ultram (Tramadol		
	Mental Disorder	Hydrochloride)	SS	
	Migraine	Clonopin		
	Muscular Weakness	(Clonazepam)	C	
	Neuropathy	Vioxx (Rofecoxib)	C	
	Swelling	Voltaren (Diclofenac		
	Tremor	Sodium)	C	
	Weight Increased			

Date:08/01/05ISR Number: 4732802-6Report Type:Expedited (15-DaCompany Report #2005086715
Age:72 YR Gender:Male I/FU:F

Outcome	PT
Hospitalization -	Appetite Disorder
Initial or Prolonged	Blood Alkaline
Other	Phosphatase Increased
	C-Reactive Protein
	Increased
	Electrocardiogram
	Abnormal
	Eosinophilia

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Dose	Duration	Gamma-Glutamyltransferase Increased Haemoglobin Decreased	Report Source	Product	Role	Manufacturer	Route
ORAL		Jaundice Pruritus	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
ORAL		Transaminases Increased	Professional	Tegretol (Carbamazepine)	SS		ORAL
				Glimepiride	C		
				Insulin Glargine	C		
				Indapamide	C		

Date:08/01/05ISR Number: 4732826-9Report Type:Expedited (15-DaCompany Report #2005105634
Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged 2400 MG (2400 MG, 1 IN 1 D), ORAL		Blood Triglycerides Increased	Foreign Health	Neurontin (Gabapentin)	PS		ORAL
		Circulating Anticoagulant Positive	Professional				
25 MG (25 MG, 1 IN 1 D), ORAL		Deep Vein Thrombosis Epistaxis Hepatic Steatosis Hypoalbuminaemia		Amitriptyline Hydrochloride (Amitriptyline Hydrochloride)	SS		ORAL
		Neuropathy					
0.5 MG (0.5 MG, 1 IN 1 D), ORAL		Oedema Peripheral Pain In Extremity		Prograf (Tacrolimus)	SS		ORAL
		Pancreatic Disorder					
		Pericardial Effusion					
300 MG (300		Pulmonary Embolism		Zyloric "Faes" (Allopurinol)	SS	Faes	ORAL

MCG, 1 IN 1

D), ORAL

2 MG (2 MG, 1

IN 1 D), ORAL

1500 MG (500

MG, 3 IN 1

D), ORAL

Rapamne (Sirolimus) SS ORAL

Cellcept (Mycophenolate Mofetil) SS ORAL

Rivotril (Clonazepam) C
Orocal (Calcium Carbonate) C
Lasilix (Furosemide) C
Levothyrox (Levothyroxine Sodium) C

Date:08/01/05ISR Number: 4732840-3Report Type:Expedited (15-DaCompany Report #2005104931

Age:51 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Hospitalization - Initial or Prolonged	1800 MG (1 IN	Dizziness Nausea Palpitations Sleep Disorder Small Intestinal Obstruction	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL
	1 D), ORAL			All Other Therapeutic Products (All Other Therapeutic			

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Products) C

Date:08/02/05ISR Number: 4736300-5Report Type:Expedited (15-DaCompany Report #2005107357
 Age:33 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Drug Ineffective	Consumer	Neurontin (Gabapentin)	PS		
DEPRESSION		Intentional Self-Injury					
		Suicide Attempt					

Date:08/02/05ISR Number: 4736305-4Report Type:Expedited (15-DaCompany Report #2005107362
 Age: Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Death		Completed Suicide	Consumer	Neurontin (Gabapentin)	PS		
		Drug Ineffective					

Date:08/02/05ISR Number: 4736370-4Report Type:Expedited (15-DaCompany Report #2004121824
 Age:55 YR Gender:Male I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Difficulty In Walking	Consumer	Neurontin (Gabapentin)	PS		ORAL
600 MG (300		Hypokinesia					
MG, 2 IN 1 D)		Vision Blurred					
ORAL		Visual Acuity Reduced		Ditropan (Oxybutynin)	C		

Date:08/02/05ISR Number: 4736517-XReport Type:Expedited (15-DaCompany Report #2005104540
 Age:8 YR Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							

Hospitalization - Initial or Prolonged 900 MG (300 MG, 3 IN 1 D), ORAL	Leukopenia Neutropenia	Foreign Health Professional Company Representative	Neurontin (Tablets) (Gabapentin)	PS	ORAL
--	---------------------------	--	-------------------------------------	----	------

Date:08/02/05ISR Number: 4736557-0Report Type:Expedited (15-DaCompany Report #2005103846
Age: Gender: I/FU:I

Outcome Dose Hospitalization - Initial or Prolonged 900 MG (900 MG), ORAL	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Blood Creatine Increased Hypotension	Foreign Health Professional	Neurontin (Gabapentin)	PS		ORAL

Date:08/03/05ISR Number: 4734849-2Report Type:Direct Company Report #CTU 255233
Age:80 YR Gender:Male I/FU:I

Outcome Dose Other	Duration	PT	Report Source	Product	Role	Manufacturer	Route
		Somnolence		Gabapentin	PS		

FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/03/05ISR Number: 4735275-2Report Type:Expedited (15-DaCompany Report #2005105713

Age: Gender:Female I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Arthritis	Consumer	Neurontin			
		Diabetes Mellitus		(Gabapentin)	PS		
		Intervertebral Disc		Aricept (Donepezil)	SS		
		Protrusion		Xalatan			
		Malaise		(Latanoprost)	SS		
		Parkinson'S Disease		Zoloft (Sertraline)	SS		
		Tremor		Synthroid			
				(Levothyroxine			
				Sodium)	C		

Date:08/04/05ISR Number: 4734442-1Report Type:Expedited (15-DaCompany Report #FR-ROCHE-411366

Age:60 YR Gender:Male I/FU:I

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Hospitalization -		Blood Triglycerides		Cellcept	PS	Roche	ORAL
Initial or Prolonged		Increased		Laroxyl	SS	Roche	ORAL
		Deep Vein Thrombosis		Rapamune	SS		ORAL
		Hepatic Steatosis		Zyloric 300	SS		ORAL
		Hypoalbuminaemia		Prograf	SS		ORAL
		Neuropathy Peripheral		Neurontin	SS		ORAL
		Pancreatic Disorder		Rivotril	C		
UNKNOWN							
		Pericardial Effusion		Orocal	C		
UNKNOWN							
		Pulmonary Embolism		Lasilix	C		
UNKNOWN							
				Levothyroxine	C		
UNKNOWN							

Date:08/04/05ISR Number: 4735175-8Report Type:Expedited (15-DaCompany Report #2005099146

Age:48 YR Gender:Female I/FU:F

Outcome	Duration	PT	Report Source	Product	Role	Manufacturer	Route
Dose							
Other		Constipation	Health	Neurontin			
		Diarrhoea	Professional	(Gabapentin)	PS		ORAL
ORAL							

Drug Ineffective

Elavil
(Amitriptyline
Hydrochloride)

SS

ORAL

10 MG(10 MG,

1 IN 1D),

ORAL

Advil (Ibuprofen)

C

Dicyclomine

(Dicycloverine)

C

Aygestin

(Norethisterone

Acetate)

C

Date:08/04/05ISR Number: 4735193-XReport Type:Expedited (15-DaCompany Report #2005105630

Age:55 YR Gender:Male I/FU:I

Outcome

PT

Hospitalization -

Blood Cholesterol

Initial or Prolonged

Increased

Other

Cardiac Disorder

Carpal Tunnel Syndrome

Drug Ineffective

Dyspnoea

Hyperhidrosis

Hypertension

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

		Hypoaesthesia Tremor	Report Source	Product	Role	Manufacturer	Route
Dose	Duration		Consumer	Celebrex (Celecoxib)	PS		ORAL
400 MG (400 MG, 1 IN 1 D), ORAL							
				Neurontin (Gabapentin)	SS		ORAL
ORAL				Mobic (Meloxicam) Hydrocodone (Hydrocodone)	SS C		
				Ambien (Zolpidem Tartrate)	C		
				Lidoderm Patch (Lidocaine Hydrochloride)	C		

Date:08/04/05ISR Number: 4735670-1Report Type:Expedited (15-DaCompany Report #2005106144
Age:35 YR Gender:Female I/FU:I

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Consumer	Neurontin (Gabapentin)	PS		ORAL
Dose		Homicidal Ideation - Memory Impairment					
Hospitalization - Initial or Prolonged							
1500 MG (2 IN Other		Suicidal Ideation					
1 D), ORAL							

Date:08/04/05ISR Number: 4736302-9Report Type:Expedited (15-DaCompany Report #2005104514
Age:83 YR Gender:Male I/FU:F

		PT	Report Source	Product	Role	Manufacturer	Route
Outcome	Duration		Consumer	Neurontin (Tablets) (Gabapentin)	PS		ORAL
Dose		Aortic Calcification Aortic Valve Stenosis					
Hospitalization - Initial or Prolonged							
2400 MG (800 Other		Blood Cholesterol					
MG, 3 IN 1							

D), ORAL

Increased

Cardiac Disorder
Carotid Artery Occlusion
Drug Effect Decreased
Eye Disorder
Knee Arthroplasty
Mitral Valve
Calcification
Transient Ischaemic
Attack

Vitamins

C

Date:08/04/05ISR Number: 4736518-1Report Type:Direct
Age:41 YR Gender:Male I/FU:I

Company Report #CTU 255430

Outcome
Other

PT
Acne
Condition Aggravated
Coordination Abnormal
Cough
Dandruff
Dry Eye
Dry Skin
Dysarthria
Enuresis
Fatigue
Fungal Skin Infection

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FDA - Adverse Event Reporting System (AERS)

Freedom Of Information (FOI) Report

Date:08/16/05ISR Number: 4745872-6Report Type:Expedited (15-DaCompany Report #FR-BRISTOL-MYERS SQUIBB COMPANY-13064902
 Age:72 YR Gender:Male I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Hospitalization -						
Initial or Prolonged	Acute Generalised		Elisor Tabs	PS	Bristol-Myers Squibb Company	ORAL
	Exanthematous Pustulosis		Zyloric	SS		
	Enterobacter Infection		Rocephine	SS		
	Rash Maculo-Papular					
INTRAMUSCULAR			Laroxyl	SS		
			Neurontin	SS		
			Risperdal	SS		
			Lasilix	C		
			Zestril	C		
			Cordarone	C		
			Sectral	C		

Date:08/16/05ISR Number: 4746039-8Report Type:Expedited (15-DaCompany Report #US-JNJFOC-20050507147
 Age: Gender:Female I/FU:I

Outcome	PT	Report Source	Product	Role	Manufacturer	Route
Dose						
Death						
800 mg, twice	Completed Suicide		Motrin 800 Mg	PS		
	Loss Of Consciousness					
to three						
times, as	Multiple Drug Overdose					
needed						
			Nortriptyline			
			Hydrochloride	SS		
			Amitriptyline			
			Hydrochloride	SS		
			Venlafaxine	SS		
			Neurontin	SS		
			Neurontin	SS		
			Neurontin	SS		
			Neurontin	SS		
			Oxazepam	C		
			Benzodiazepine			
			Derivatives	C		
			Opioids	C		

Summary report for FOI selections:

Selection by inexact search of active ingredient:

GABAPENTIN%

Selection by inexact search of Tradename/Verbatim:

NEURONTIN%

Total number of reports: 8,847

From: 01-NOV-1997

To: Present

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